
O'Donnell A, McParlin C, Robson SC, Beyer F, Moloney E, Bryant A, Bradley J, Muirhead C, Nelson-Piercy C, Newbury-Birch D, Norman J, Simpson E, Swallow B, Yates L, Vale L. *Treatments for hyperemesis gravidarum and nausea and vomiting in pregnancy: a systematic review and economic assessment*. London: NIHR Journals Library, 2016. Health Technology Assessment.

Copyright:

© Queen's Printer and Controller of HMSO 2016. This work was produced by O'Donnell et al. under the terms of a commissioning contract issued by the Secretary of State for Health. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

DOI link to article:

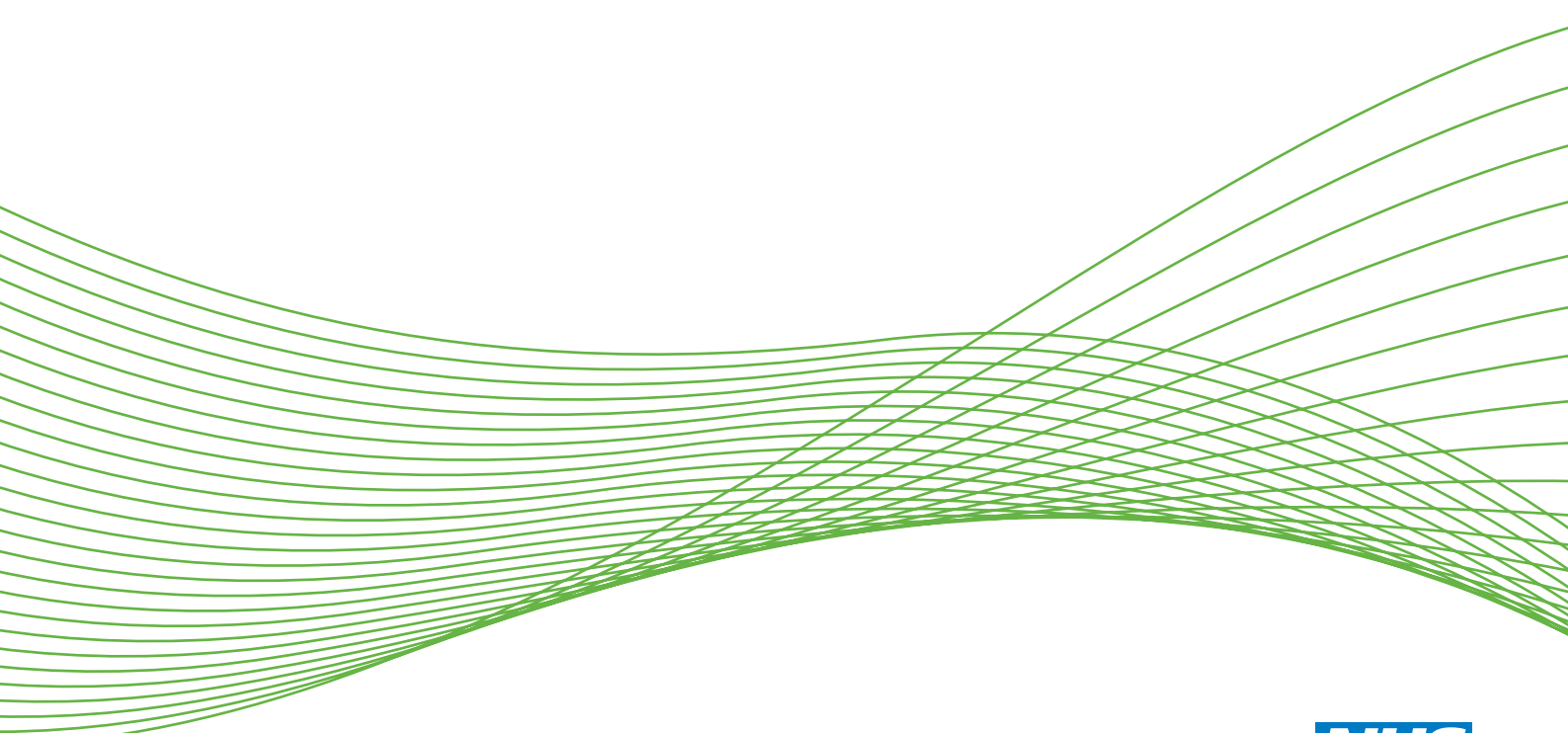
<https://dx.doi.org/10.3310/hta20740>

Date deposited:

12/01/2017

Treatments for hyperemesis gravidarum and nausea and vomiting in pregnancy: a systematic review and economic assessment

Amy O'Donnell, Catherine McParlin, Stephen C Robson, Fiona Beyer, Eoin Moloney, Andrew Bryant, Jennifer Bradley, Colin Muirhead, Catherine Nelson-Piercy, Dorothy Newbury-Birch, Justine Norman, Emma Simpson, Brian Swallow, Laura Yates and Luke Vale



**National Institute for
Health Research**

Treatments for hyperemesis gravidarum and nausea and vomiting in pregnancy: a systematic review and economic assessment

Amy O'Donnell,¹ Catherine McParlin,²
Stephen C Robson,³ Fiona Beyer,¹ Eoin Moloney,⁴
Andrew Bryant,¹ Jennifer Bradley,¹ Colin Muirhead,¹
Catherine Nelson-Piercy,⁵ Dorothy Newbury-Birch,¹
Justine Norman,⁶ Emma Simpson,¹ Brian Swallow,⁷
Laura Yates⁸ and Luke Vale^{4*}

¹Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK

²Newcastle upon Tyne Hospitals NHS Foundation Trust, Newcastle upon Tyne, UK

³Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK

⁴Health Economics Group, Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK

⁵Women's Health Academic Centre, King's Health Partners, King's College London, London, UK

⁶North Tyneside Clinical Commissioning Group, Whitley Bay, UK

⁷Expert Advisor

⁸UK Teratology Information Service (UKTIS) and Institute of Genetic Medicine, Newcastle University, Newcastle upon Tyne, UK

*Corresponding author

Declared competing interests of authors: Luke Vale is a member of the funding panel for the National Institute for Health Research (NIHR) Programme Grants for Applied Research and NIHR Health Technology Assessment. He is also a Director of the NIHR Research Design Service in the North East. Stephen C Robson is a panel member of NIHR Efficacy and Mechanism Evaluation. Catherine Nelson-Piercy is a co-developer of the Royal College of Obstetricians and Gynaecologists green-top guideline on management of nausea vomiting and hyperemesis gravidarum.

Published October 2016

DOI: 10.3310/hta20740

This report should be referenced as follows:

O'Donnell A, McParlin C, Robson SC, Beyer F, Moloney E, Bryant A, *et al.* Treatments for hyperemesis gravidarum and nausea and vomiting in pregnancy: a systematic review and economic assessment. *Health Technol Assess* 2016;**20**(74).

Health Technology Assessment is indexed and abstracted in *Index Medicus/MEDLINE*, *Excerpta Medica/EMBASE*, *Science Citation Index Expanded (SciSearch®)* and *Current Contents®/Clinical Medicine*.

ISSN 1366-5278 (Print)

ISSN 2046-4924 (Online)

Impact factor: 4.058

Health Technology Assessment is indexed in MEDLINE, CINAHL, EMBASE, The Cochrane Library and the ISI Science Citation Index.

This journal is a member of and subscribes to the principles of the Committee on Publication Ethics (COPE) (www.publicationethics.org/).

Editorial contact: nhredit@southampton.ac.uk

The full HTA archive is freely available to view online at www.journalslibrary.nihr.ac.uk/hta. Print-on-demand copies can be purchased from the report pages of the NIHR Journals Library website: www.journalslibrary.nihr.ac.uk

Criteria for inclusion in the *Health Technology Assessment* journal

Reports are published in *Health Technology Assessment* (HTA) if (1) they have resulted from work for the HTA programme, and (2) they are of a sufficiently high scientific quality as assessed by the reviewers and editors.

Reviews in *Health Technology Assessment* are termed 'systematic' when the account of the search appraisal and synthesis methods (to minimise biases and random errors) would, in theory, permit the replication of the review by others.

HTA programme

The HTA programme, part of the National Institute for Health Research (NIHR), was set up in 1993. It produces high-quality research information on the effectiveness, costs and broader impact of health technologies for those who use, manage and provide care in the NHS. 'Health technologies' are broadly defined as all interventions used to promote health, prevent and treat disease, and improve rehabilitation and long-term care.

The journal is indexed in NHS Evidence via its abstracts included in MEDLINE and its Technology Assessment Reports inform National Institute for Health and Care Excellence (NICE) guidance. HTA research is also an important source of evidence for National Screening Committee (NSC) policy decisions.

For more information about the HTA programme please visit the website: <http://www.nets.nihr.ac.uk/programmes/hta>

This report

The research reported in this issue of the journal was funded by the HTA programme as project number 12/152/01. The contractual start date was in September 2013. The draft report began editorial review in December 2014 and was accepted for publication in March 2015. The authors have been wholly responsible for all data collection, analysis and interpretation, and for writing up their work. The HTA editors and publisher have tried to ensure the accuracy of the authors' report and would like to thank the reviewers for their constructive comments on the draft document. However, they do not accept liability for damages or losses arising from material published in this report.

This report presents independent research funded by the National Institute for Health Research (NIHR). The views and opinions expressed by authors in this publication are those of the authors and do not necessarily reflect those of the NHS, the NIHR, NETSCC, the HTA programme or the Department of Health. If there are verbatim quotations included in this publication the views and opinions expressed by the interviewees are those of the interviewees and do not necessarily reflect those of the authors, those of the NHS, the NIHR, NETSCC, the HTA programme or the Department of Health.

© Queen's Printer and Controller of HMSO 2016. This work was produced by O'Donnell *et al.* under the terms of a commissioning contract issued by the Secretary of State for Health. This issue may be freely reproduced for the purposes of private research and study and extracts (or indeed, the full report) may be included in professional journals provided that suitable acknowledgement is made and the reproduction is not associated with any form of advertising. Applications for commercial reproduction should be addressed to: NIHR Journals Library, National Institute for Health Research, Evaluation, Trials and Studies Coordinating Centre, Alpha House, University of Southampton Science Park, Southampton SO16 7NS, UK.

Published by the NIHR Journals Library (www.journalslibrary.nihr.ac.uk), produced by Prepress Projects Ltd, Perth, Scotland (www.prepress-projects.co.uk).

Health Technology Assessment Editor-in-Chief

Professor Hywel Williams Director, HTA Programme, UK and Foundation Professor and Co-Director of the Centre of Evidence-Based Dermatology, University of Nottingham, UK

NIHR Journals Library Editor-in-Chief

Professor Tom Walley Director, NIHR Evaluation, Trials and Studies and Director of the EME Programme, UK

NIHR Journals Library Editors

Professor Ken Stein Chair of HTA Editorial Board and Professor of Public Health, University of Exeter Medical School, UK

Professor Andree Le May Chair of NIHR Journals Library Editorial Group (EME, HS&DR, PGfAR, PHR journals)

Dr Martin Ashton-Key Consultant in Public Health Medicine/Consultant Advisor, NETSCC, UK

Professor Matthias Beck Chair in Public Sector Management and Subject Leader (Management Group), Queen's University Management School, Queen's University Belfast, UK

Professor Aileen Clarke Professor of Public Health and Health Services Research, Warwick Medical School, University of Warwick, UK

Dr Tessa Crilly Director, Crystal Blue Consulting Ltd, UK

Dr Eugenia Cronin Senior Scientific Advisor, Wessex Institute, UK

Ms Tara Lamont Scientific Advisor, NETSCC, UK

Professor William McGuire Professor of Child Health, Hull York Medical School, University of York, UK

Professor Geoffrey Meads Professor of Health Sciences Research, Health and Wellbeing Research and Development Group, University of Winchester, UK

Professor John Norrie Health Services Research Unit, University of Aberdeen, UK

Professor John Powell Consultant Clinical Adviser, National Institute for Health and Care Excellence (NICE), UK

Professor James Raftery Professor of Health Technology Assessment, Wessex Institute, Faculty of Medicine, University of Southampton, UK

Dr Rob Riemsma Reviews Manager, Kleijnen Systematic Reviews Ltd, UK

Professor Helen Roberts Professor of Child Health Research, UCL Institute of Child Health, UK

Professor Jonathan Ross Professor of Sexual Health and HIV, University Hospital Birmingham, UK

Professor Helen Snooks Professor of Health Services Research, Institute of Life Science, College of Medicine, Swansea University, UK

Professor Jim Thornton Professor of Obstetrics and Gynaecology, Faculty of Medicine and Health Sciences, University of Nottingham, UK

Professor Martin Underwood Director, Warwick Clinical Trials Unit, Warwick Medical School, University of Warwick, UK

Please visit the website for a list of members of the NIHR Journals Library Board:
www.journalslibrary.nihr.ac.uk/about/editors

Editorial contact: nihredit@southampton.ac.uk

Abstract

Treatments for hyperemesis gravidarum and nausea and vomiting in pregnancy: a systematic review and economic assessment

Amy O'Donnell,¹ Catherine McParlin,² Stephen C Robson,³ Fiona Beyer,¹ Eoin Moloney,⁴ Andrew Bryant,¹ Jennifer Bradley,¹ Colin Muirhead,¹ Catherine Nelson-Piercy,⁵ Dorothy Newbury-Birch,¹ Justine Norman,⁶ Emma Simpson,¹ Brian Swallow,⁷ Laura Yates⁸ and Luke Vale^{4*}

¹Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK

²Newcastle upon Tyne Hospitals NHS Foundation Trust, Newcastle upon Tyne, UK

³Institute of Cellular Medicine, Newcastle University, Newcastle upon Tyne, UK

⁴Health Economics Group, Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK

⁵Women's Health Academic Centre, King's Health Partners, King's College London, London, UK

⁶North Tyneside Clinical Commissioning Group, Whitley Bay, UK

⁷Expert Advisor

⁸UK Teratology Information Service (UKTIS) and Institute of Genetic Medicine, Newcastle University, Newcastle upon Tyne, UK

*Corresponding author luke.vale@newcastle.ac.uk

Background: Nausea and vomiting in pregnancy (NVP) affects up to 85% of all women during pregnancy, but for the majority self-management suffices. For the remainder, symptoms are more severe and the most severe form of NVP – hyperemesis gravidarum (HG) – affects 0.3–1.0% of pregnant women. There is no widely accepted point at which NVP becomes HG.

Objectives: This study aimed to determine the relative clinical effectiveness and cost-effectiveness of treatments for NVP and HG.

Data sources: MEDLINE, EMBASE, Cumulative Index to Nursing and Allied Health Literature, Cochrane Central Register of Controlled Trials, PsycINFO, Commonwealth Agricultural Bureaux (CAB) Abstracts, Latin American and Caribbean Health Sciences Literature, Allied and Complementary Medicine Database, British Nursing Index, Science Citation Index, Social Sciences Citation Index, Scopus, Conference Proceedings Index, NHS Economic Evaluation Database, Health Economic Evaluations Database, China National Knowledge Infrastructure, Cochrane Database of Systematic Reviews and Database of Abstracts of Reviews of Effects were searched from inception to September 2014. References from studies and literature reviews identified were also examined. *Obstetric Medicine* was hand-searched, as were websites of relevant organisations. Costs came from NHS sources.

Review methods: A systematic review of randomised and non-randomised controlled trials (RCTs) for effectiveness, and population-based case series for adverse events and fetal outcomes. Treatments: vitamins B6 and B12, ginger, acupressure/acupuncture, hypnotherapy, antiemetics, dopamine antagonists, 5-hydroxytryptamine receptor antagonists, intravenous (i.v.) fluids, corticosteroids, enteral and parenteral

feeding or other novel treatment. Two reviewers extracted data and quality assessed studies. Results were narratively synthesised; planned meta-analysis was not possible due to heterogeneity and incomplete reporting. A simple economic evaluation considered the implied values of treatments.

Results: Seventy-three studies (75 reports) met the inclusion criteria. For RCTs, 33 and 11 studies had a low and high risk of bias respectively. For the remainder ($n = 20$) it was unclear. The non-randomised studies ($n = 9$) were low quality. There were 33 separate comparators. The most common were acupuncture versus placebo ($n = 12$); steroid versus usual treatment ($n = 7$); ginger versus placebo ($n = 6$); ginger versus vitamin B6 ($n = 6$); and vitamin B6 versus placebo ($n = 4$). There was evidence that ginger, antihistamines, metoclopramide (mild disease) and vitamin B6 (mild to severe disease) are better than placebo. Diclectin® [Duchesnay Inc.; doxylamine succinate (10 mg) plus pyridoxine hydrochloride (10 mg) slow release tablet] is more effective than placebo and ondansetron is more effective at reducing nausea than pyridoxine plus doxylamine. Diclectin before symptoms of NVP begin for women at high risk of severe NVP recurrence reduces risk of moderate/severe NVP compared with taking Diclectin once symptoms begin. Promethazine is as, and ondansetron is more, effective than metoclopramide for severe NVP/HG. I.v. fluids help correct dehydration and improve symptoms. Dextrose saline may be more effective at reducing nausea than normal saline. Transdermal clonidine patches may be effective for severe HG. Enteral feeding is effective but extreme method treatment for very severe symptoms. Day case management for moderate/severe symptoms is feasible, acceptable and as effective as inpatient care. For all other interventions and comparisons, evidence is unclear. The economic analysis was limited by lack of effectiveness data, but comparison of costs between treatments highlights the implications of different choices.

Limitations: The main limitations were the quantity and quality of the data available.

Conclusion: There was evidence of some improvement in symptoms for some treatments, but these data may not be transferable across disease severities. Methodologically sound and larger trials of the main therapies considered within the UK NHS are needed.

Study registration: This study is registered as PROSPERO CRD42013006642.

Funding: The National Institute for Health Research Health Technology Assessment programme.

Contents

List of tables	xvii
List of figures	xxi
List of boxes	xxiii
List of abbreviations	xxv
Plain English summary	xxvii
Scientific summary	xxix
Chapter 1 Introduction and background	1
Background	1
Aetiology	1
Impact on patients	2
Assessment and diagnosis	2
Current interventions for nausea and vomiting in pregnancy/hyperemesis gravidarum	3
<i>Patient-initiated first-line interventions</i>	4
<i>Clinician-prescribed second-line interventions</i>	6
<i>Clinician-prescribed third-line interventions</i>	6
<i>Interventions presented in the report but not routinely used to treat nausea and vomiting in pregnancy</i>	7
Current guidance and use of therapies within the NHS	7
Aims and objectives	8
Structure of the report	8
Chapter 2 Methods for the systematic review of effectiveness	9
General methodology	9
Inclusion criteria	9
<i>Types of studies</i>	9
<i>Types of participants</i>	9
<i>Types of interventions and comparators</i>	9
<i>Types of outcome measures</i>	10
Search strategy	10
First exclusion process	12
Second exclusion process	12
Data extraction	13
Risk of bias in included studies and quality assessment	13
Data synthesis	14
Chapter 3 Clinical effectiveness: overview of included studies	15
Studies identified	15
Quality of included studies	16
<i>Randomised controlled trials</i>	16
<i>Case series studies</i>	21
Interventions and comparators	22
Participants and symptom severity	26

Outcome measures	26
Additional sources of outcome data on medications	26
Meta-analysis of included randomised controlled trials	29
Structure of individual results chapters	29
Chapter 4 Clinical effectiveness: ginger	31
Introduction	31
Ginger capsules versus placebo capsules	31
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	31
<i>Author-defined symptom severity/relief scales</i>	31
<i>Nausea outcomes</i>	38
<i>Vomiting outcomes</i>	38
<i>Retching outcomes</i>	39
<i>Safety outcomes</i>	39
Ginger syrup versus placebo syrup	39
<i>Combined severity score</i>	39
<i>Nausea outcomes</i>	39
<i>Vomiting outcomes</i>	39
<i>Retching outcomes</i>	39
<i>Safety outcomes</i>	40
Ginger biscuit versus placebo biscuit	40
<i>Author-defined symptom severity/relief scales</i>	40
<i>Nausea outcomes</i>	40
<i>Vomiting outcomes</i>	40
<i>Retching outcomes</i>	40
<i>Safety outcomes</i>	40
Ginger versus vitamin B6	40
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	40
<i>Pregnancy-Unique Quantification of Emesis and Nausea scale</i>	41
<i>Author-defined symptom severity/relief scales</i>	41
<i>Nausea outcomes</i>	41
<i>Vomiting outcomes</i>	41
<i>Retching outcomes</i>	41
<i>Safety outcomes</i>	42
Ginger capsules versus acupressure	42
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	42
<i>Nausea outcomes</i>	42
<i>Vomiting outcomes</i>	42
<i>Retching outcomes</i>	42
<i>Safety outcomes</i>	43
Ginger versus doxylamine–pyridoxine	43
<i>Combined severity score</i>	43
<i>Nausea outcomes</i>	43
<i>Vomiting outcomes</i>	43
<i>Retching outcomes</i>	43
<i>Safety outcomes</i>	43
Ginger versus antihistamine (dimenhydrinate) capsules	43
<i>Combined severity score</i>	43
<i>Nausea outcomes</i>	43
<i>Vomiting outcomes</i>	44
<i>Retching outcomes</i>	44
<i>Safety outcomes</i>	44
Ginger versus metoclopramide	44
Summary	44

Chapter 5 Clinical effectiveness: acupressure, acupuncture and nerve stimulation	45
Introduction	45
Acupressure versus placebo	45
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	45
<i>McGill Nausea Questionnaire</i>	63
<i>Nausea outcomes</i>	63
<i>Vomiting outcomes</i>	64
<i>Retching outcomes</i>	64
<i>Safety outcomes</i>	64
Acupressure versus vitamin B6	64
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	64
<i>Nausea outcomes</i>	64
<i>Vomiting outcomes</i>	65
<i>Retching outcomes</i>	65
<i>Safety outcomes</i>	65
Acupressure (case series)	65
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	65
<i>Nausea outcomes</i>	65
<i>Vomiting outcomes</i>	65
<i>Retching outcomes</i>	65
<i>Safety outcomes</i>	65
Nerve stimulation versus placebo	65
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	66
<i>Author-defined scale</i>	66
<i>Nausea outcomes</i>	66
<i>Vomiting outcomes</i>	66
<i>Retching outcomes</i>	66
<i>Safety outcomes</i>	66
Acupuncture versus placebo	66
<i>Combined severity score</i>	66
<i>Nausea outcomes</i>	66
<i>Vomiting outcomes</i>	67
<i>Retching outcomes</i>	67
<i>Safety outcomes</i>	67
Acupuncture versus metoclopramide	68
<i>Combined severity score</i>	68
<i>Nausea outcomes</i>	68
<i>Vomiting outcomes</i>	68
<i>Retching outcomes</i>	68
<i>Safety outcomes</i>	68
Acupuncture versus Chinese herbal medicine versus barbiturates	68
<i>Author-defined scale</i>	68
<i>Nausea outcomes</i>	69
<i>Vomiting outcomes</i>	69
<i>Retching outcomes</i>	69
<i>Safety outcomes</i>	69
Summary	69
Chapter 6 Clinical effectiveness: aromatherapy	71
Introduction	71
Aromatherapy versus no aromatherapy	71
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	71
<i>Nausea outcomes</i>	71

<i>Vomiting outcomes</i>	71
<i>Retching outcomes</i>	71
<i>Safety outcomes</i>	73
Summary	73
Chapter 7 Clinical effectiveness: vitamin B6 (pyridoxine)	75
Introduction	75
Vitamin B6 versus placebo	75
<i>Pregnancy-Unique Quantification of Emesis and Nausea scale</i>	75
<i>Nausea outcomes</i>	75
<i>Vomiting outcomes</i>	75
<i>Retching outcomes</i>	75
<i>Safety outcomes</i>	79
Vitamin B6 and metoclopramide combination versus metoclopramide alone	79
<i>Combined severity score</i>	79
<i>Nausea outcomes</i>	79
<i>Vomiting outcomes</i>	79
<i>Retching outcomes</i>	79
<i>Safety outcomes</i>	79
Vitamin B6 versus serotonin antagonist	79
<i>Rhodes Index of Nausea, Vomiting and Retching</i>	79
<i>Nausea outcomes</i>	79
<i>Vomiting outcomes</i>	79
<i>Retching outcomes</i>	79
<i>Safety outcomes</i>	80
Summary	80
Chapter 8 Clinical effectiveness: pyridoxine/doxylamine combination	81
Introduction	81
Doxylamine/pyridoxine versus placebo	81
<i>Pregnancy-Unique Quantification of Emesis and Nausea scale</i>	81
<i>Nausea outcomes</i>	81
<i>Vomiting outcomes</i>	81
<i>Retching outcomes</i>	85
<i>Safety outcomes</i>	85
Doxylamine/pyridoxine versus ondansetron	85
<i>Combined severity score</i>	85
<i>Nausea outcomes</i>	85
<i>Vomiting outcomes</i>	85
<i>Retching outcomes</i>	85
<i>Safety outcomes</i>	85
Pre-emptive doxylamine/pyridoxine	85
<i>Pregnancy-Unique Quantification of Emesis and Nausea scale</i>	85
<i>Nausea outcomes</i>	86
<i>Vomiting outcomes</i>	86
<i>Retching outcomes</i>	86
<i>Safety outcomes</i>	86
Summary	86
Chapter 9 Clinical effectiveness: antihistamines	87
Introduction	87
Antihistamines versus placebo	87
<i>Author-defined symptom severity/relief scale</i>	87

<i>Nausea outcomes</i>	87
<i>Vomiting outcomes</i>	87
<i>Retching outcomes</i>	87
<i>Safety outcomes</i>	87
Antihistamines alone or in combination with Vitamin B6 versus control	89
<i>Author-defined scale</i>	89
<i>Nausea outcomes</i>	89
<i>Vomiting outcomes</i>	89
<i>Retching outcomes</i>	89
<i>Safety outcomes</i>	89
Summary	89
Chapter 10 Clinical effectiveness: dopamine antagonists	91
Introduction	91
Promethazine versus metoclopramide	91
<i>Combined severity score</i>	91
<i>Nausea outcomes</i>	91
<i>Vomiting outcomes</i>	91
<i>Retching score</i>	91
<i>Safety outcomes</i>	91
Summary	94
Chapter 11 Clinical effectiveness: serotonin antagonists (ondansetron)	95
Introduction	95
Ondansetron versus usual treatment	95
Ondansetron versus metoclopramide	95
<i>Combined severity score</i>	95
<i>Nausea outcomes</i>	95
<i>Vomiting outcomes</i>	100
<i>Retching outcomes</i>	100
<i>Safety outcomes</i>	100
Ondansetron versus antihistamines	100
<i>Author-defined relief scale</i>	100
<i>Nausea outcomes</i>	100
<i>Vomiting outcomes</i>	100
<i>Retching outcomes</i>	101
<i>Safety outcomes</i>	101
Summary	101
Chapter 12 Clinical effectiveness: intravenous fluids	103
Introduction	103
Dextrose saline versus saline only	103
<i>Combined severity score</i>	103
<i>Nausea outcomes</i>	103
<i>Vomiting outcomes</i>	103
<i>Retching outcomes</i>	103
<i>Safety outcomes</i>	103
Intravenous fluids versus intravenous fluids plus diazepam	103
<i>Combined severity score</i>	103
<i>Nausea outcomes</i>	105
<i>Vomiting outcomes</i>	105

<i>Retching outcomes</i>	105
<i>Safety outcomes</i>	105
Summary	105
Chapter 13 Clinical effectiveness: transdermal clonidine	107
Introduction	107
Transdermal clonidine versus placebo patch	107
<i>Pregnancy-Unique Quantification of Emesis and Nausea scale</i>	107
<i>Nausea outcomes</i>	107
<i>Vomiting outcomes</i>	107
<i>Retching outcomes</i>	107
<i>Safety outcomes</i>	107
Summary	107
Chapter 14 Clinical effectiveness: outpatient/day case management	109
Introduction	109
Outpatient management versus standard inpatient care	109
<i>Pregnancy-Unique Quantification of Emesis and Nausea scale</i>	109
<i>Nausea outcomes</i>	109
<i>Vomiting outcomes</i>	109
<i>Retching outcomes</i>	109
<i>Safety outcomes</i>	109
Day case management	109
Summary	109
Chapter 15 Clinical effectiveness: corticosteroids	111
Introduction	111
Corticosteroids versus placebo	111
<i>Combined severity score</i>	111
<i>Nausea outcomes</i>	111
<i>Vomiting outcomes</i>	111
<i>Retching outcomes</i>	111
<i>Safety outcomes</i>	117
Corticosteroids versus promethazine (Phenergan)	117
<i>Combined severity score</i>	117
<i>Nausea outcomes</i>	117
<i>Vomiting outcomes</i>	117
<i>Retching outcomes</i>	117
<i>Safety outcomes</i>	117
Corticosteroids versus metoclopramide	117
<i>Combined severity score</i>	117
<i>Nausea outcomes</i>	118
<i>Vomiting outcomes</i>	118
<i>Retching outcomes</i>	118
<i>Safety outcomes</i>	118
Corticosteroids against usual treatment	118
<i>Combined severity score</i>	118
<i>Nausea outcomes</i>	118
<i>Vomiting outcomes</i>	118
<i>Retching outcomes</i>	118
<i>Safety outcomes</i>	118
Summary	118

Chapter 16 Clinical effectiveness: nasogastric enteral/jejunostomy feeding	119
Introduction	119
Nasogastric enteral feeding	119
<i>Combined severity score</i>	119
<i>Nausea outcomes</i>	119
<i>Vomiting outcomes</i>	119
<i>Retching outcomes</i>	119
<i>Safety outcomes</i>	119
Jejunostomy feeding	119
<i>Combined severity score</i>	121
<i>Nausea outcomes</i>	121
<i>Vomiting outcomes</i>	121
<i>Retching outcomes</i>	121
<i>Safety outcomes</i>	121
Summary	121
Chapter 17 Clinical effectiveness: gabapentin	123
Introduction	123
<i>Author-defined scale</i>	123
<i>Nausea outcomes</i>	123
<i>Vomiting outcomes</i>	123
<i>Retching outcomes</i>	123
<i>Safety outcomes</i>	123
Summary	123
Chapter 18 Economic analysis	125
Introduction	125
Systematic review of economic evaluations	125
<i>Methods</i>	125
<i>Results</i>	126
<i>Discussion</i>	126
Economic modelling	127
<i>Introduction</i>	127
<i>Methods</i>	127
Economic evaluation	129
<i>Introduction</i>	129
<i>Estimation of costs of interventions</i>	130
<i>Estimation of total cost of care</i>	130
<i>Methods</i>	140
<i>Results</i>	140
Summary	151
Chapter 19 Issues of importance to patients	153
Introduction	153
Background	153
Relating what patients want to review findings	153
Summary	155

Chapter 20 Issues of importance to practitioners	157
Introduction	157
Findings	157
<i>First-line 'over-the-counter' interventions and alternative therapies</i>	157
<i>Second-line interventions prescribed by general practitioners in primary care settings</i>	158
<i>Second- and third-line interventions delivered in secondary care settings</i>	159
Summary	160
Chapter 21 Discussion	161
Clinical effectiveness and harms	161
<i>First-line 'over-the-counter' interventions</i>	162
<i>Second-line interventions prescribed by general practitioners in primary care settings</i>	162
<i>Second- and third-line interventions delivered in secondary care settings</i>	163
Cost-effectiveness	164
Strengths and limitations	165
Chapter 22 Conclusions	167
Introduction	167
Implications for women and for practitioners	167
Recommendations for research	168
<i>Trajectory of research</i>	168
<i>Research recommendations</i>	168
Acknowledgements	171
References	173
Appendix 1 Examples of hyperemesis gravidarum/nausea and vomiting in pregnancy assessment tools	183
Appendix 2 Data abstraction form: clinical effectiveness	185
Appendix 3 Risk of bias for randomised controlled trials	195
Appendix 4 Quality of case series studies	199
Appendix 5 Included papers	207
Appendix 6 Excluded papers and reasons for exclusion	217
Appendix 7 UK Teratology Information Service enquiries and follow-ups relating to hyperemesis gravidarum/nausea and vomiting in pregnancy medication	227
Appendix 8 Secondary outcome data	231
Appendix 9 Systematic review of published economic evaluations: inclusion criteria	241
Appendix 10 Cost of drug interventions and recommended daily doses	243

List of tables

TABLE 1 Tools used to measure the severity of NVP	3
TABLE 2 Secondary outcome measures	4
TABLE 3 List of search terms	11
TABLE 4 Risk of bias summary: review authors' judgements about each risk of bias item for included RCTs	17
TABLE 5 Study quality summary: review authors' judgements about each risk of bias item for each included case series or non-randomised study	22
TABLE 6 Number of studies by intervention and comparator	23
TABLE 7 Validated and non-validated symptom severity measures employed by each included study	27
TABLE 8 Results for ginger-based interventions for NVP	32
TABLE 9 Results for acupuncture, nerve stimulation and acupuncture interventions NVP	46
TABLE 10 Results for aromatherapy interventions for NVP	72
TABLE 11 Results for vitamin B6 interventions for NVP	76
TABLE 12 Results for pyridoxine-doxylamine interventions for NVP	82
TABLE 13 Results for antihistamine interventions for NVP	88
TABLE 14 Results for dopamine antagonist interventions for NVP	92
TABLE 15 Results for serotonin antagonist interventions for NVP/HG	96
TABLE 16 Results for i.v. fluid interventions for NVP and HG	104
TABLE 17 Results for transdermal clonidine interventions for HG	108
TABLE 18 Results for day case/outpatient management for NVP and HG	110
TABLE 19 Results for corticosteroid interventions for NVP and HG	112
TABLE 20 Results for nasogastric enteral/jejunostomy feeding for HG	120
TABLE 21 Results for gabapentin interventions for HG	124
TABLE 22 Weekly costs of pharmacological interventions	131
TABLE 23 Non-pharmacological costs	132

TABLE 24 Cost of patient-initiated first-line interventions	133
TABLE 25 Cost of patient-initiated first-line interventions following a GP visit	133
TABLE 26 Cost of clinician-prescribed second-line interventions following a GP visit	134
TABLE 27 Cost of clinician-prescribed second-line interventions if attending hospital as a 'day case'	136
TABLE 28 Cost of clinician-prescribed second-line interventions if admitted as an inpatient	138
TABLE 29 Cost of clinician-prescribed second-line interventions × 2 if admitted as an inpatient	138
TABLE 30 Cost of clinician-prescribed third-line interventions if admitted as an inpatient	138
TABLE 31 Cost of day case management compared with inpatient management	141
TABLE 32 Cost comparisons of patient-initiated first-line interventions	142
TABLE 33 Cost comparisons of patient-initiated first-line interventions following a GP visit	143
TABLE 34 Cost comparisons of clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit	143
TABLE 35 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case'	144
TABLE 36 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient	147
TABLE 37 Cost comparison of clinician-prescribed second-line interventions × 2 if admitted as an inpatient	149
TABLE 38 Cost comparison of clinician-prescribed third-line interventions if admitted as an inpatient	149
TABLE 39 Cost comparison of 2-day day case management with 2-day inpatient management	149
TABLE 40 Examples of bad HG/NVP practice by HCPs	154
TABLE 41 Examples of good HG/NVP practice by HCPs	155
TABLE 42 UK Teratology Information Service enquires relating to HG/NVP medications	229
TABLE 43 Weekly cost of all interventions	243
TABLE 44 Recommended dose and unit cost for all pharmacological interventions	244

TABLE 45 Cost of patient-initiated first-line interventions	246
TABLE 46 Cost of patient-initiated first-line interventions following a GP visit	247
TABLE 47 Cost of clinician-prescribed second-line interventions following a GP visit	247
TABLE 48 Cost of clinician-prescribed second-line interventions if attending hospital as a 'day case'	248
TABLE 49 Cost of clinician-prescribed second-line interventions if admitted as an inpatient	250
TABLE 50 Cost of clinician-prescribed second-line interventions × 2 if admitted as an inpatient	252
TABLE 51 Cost of clinician-prescribed third-line interventions if admitted as an inpatient	252
TABLE 52 Cost of day case management compared with inpatient management	253
TABLE 53 Cost comparisons of patient-initiated first-line interventions	254
TABLE 54 Cost comparisons of patient-initiated first-line interventions following a GP visit	254
TABLE 55 Cost comparisons of clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit	255
TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case'	257
TABLE 57 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient	265
TABLE 58 Cost comparisons of clinician-prescribed second-line interventions × 2 if admitted as an inpatient	268
TABLE 59 Cost comparisons of clinician-prescribed third-line interventions if admitted as an inpatient	268
TABLE 60 Cost comparison of 2-day day case management with 2-day inpatient management	268

List of figures

FIGURE 1 Treatments for NVP	5
FIGURE 2 Flow chart of clinical effectiveness literature	15
FIGURE 3 Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included RCT studies	16
FIGURE 4 Network plot of range of interventions and comparisons for NVP/HG	25
FIGURE 5 Flow diagram showing study selection for economic evaluations review	126
FIGURE 6 Structure of the decision model	128

List of boxes

BOX 1 Characteristics of the cost-effectiveness analysis

127

List of abbreviations

5-HT ₃	5-hydroxytryptamine	OR	odds ratio
CI	confidence interval	P6	pericardium 6
ECG	electrocardiogram	PSS	Pregnancy Sickness Support
EPHPP	Effective Public Health Practice Project	PUQE	Pregnancy-Unique Quantification of Emesis and Nausea
GP	general practitioner	QoL	quality of life
GRADE	Grading of Recommendations Assessment, Development and Evaluation	QT	time between start of the Q wave and of the T wave in the heart's electrical cycle
HCP	health-care professional	RCT	randomised controlled trial
HG	hyperemesis gravidarum	RINVR	Rhodes Index of Nausea, Vomiting and Retching
i.m.	intramuscular	SD	standard deviation
IQR	interquartile range	SoF	summary of findings
i.v.	intravenous	TPN	total parenteral nutrition
MeSH	medical subject heading	UKTIS	UK Teratology Information Service
NVP	nausea and vomiting in pregnancy	VAS	visual analogue scale
NVPI	Nausea and Vomiting of Pregnancy Instrument		

Plain English summary

Up to 85% of women suffer nausea and vomiting during the first half of pregnancy. Between 30% and 35% of these suffer symptoms that are severe. Hyperemesis gravidarum refers to the most severe form of nausea and vomiting and affects 0.3–1.0% of pregnant women.

There are medicinal and non-medicinal treatments for nausea and vomiting. Changes in diet or lifestyle are often the first treatments women might try. Similarly, women may buy vitamins B6 and B12, or ginger supplements. Other therapies may also be purchased or recommended by a health-care practitioner (e.g. acupuncture/acupuncture). Some interventions need to be prescribed such as antiemetic drugs. A small number of women with severe symptoms may receive intravenous fluids, corticosteroids and assisted feeding.

Our results suggest that ginger preparations, vitamin B6, antihistamines and metoclopramide were better than placebo for mild disease. Effectiveness of treatments in more severe disease is unclear and evidence limited. Antihistamines, metoclopramide and ondansetron appear to be effective for some women, but there is no strong evidence to say which is better than the other. The overall quality of the evidence was low or very low for all treatment comparisons due to clinical differences between studies, poor and incomplete reporting of outcomes and concerns regarding risk of bias. Of note, however, was the finding that symptoms tended to improve after a few days (even with placebo). Therefore, we inferred that if symptoms have not improved, or not improved sufficiently after a short time, a change of treatment could be considered.

Scientific summary

Background

Nausea and vomiting in pregnancy (NVP) is one of the commonest symptoms of pregnancy affecting 50–85% of women during the first half of pregnancy. Symptoms usually start between 6 and 8 weeks, and most resolve by 20 weeks. Most women (65–70%) self-manage, but for the remainder symptoms are more severe. The most severe form – hyperemesis gravidarum (HG) – affects 0.3–1.0% of pregnant women and is characterised by intractable vomiting, dehydration, electrolyte imbalance, nutritional deficiencies and weight loss. There is no widely accepted point at which NVP becomes HG. A number of different treatments are available grouped as (1) first-line interventions, usually initiated by women before seeking medical care and tend to be used in less severe NVP; (2) second-line interventions, typically prescribed when a woman presents to medical care [initially this may be a general practitioner (GP) but it may involve referral of women with more severe symptoms to hospital care]; and (3) third-line interventions, reserved for women in hospital with intractable symptoms, despite second-line therapies.

Aims

This study aimed to:

- review systematically the clinical effectiveness and cost-effectiveness of each treatment for NVP/HG
- determine which therapies are most likely to be cost-effective for implementation into the NHS
- identify and prioritise future research needs.

Methods

Clinical effectiveness review

We conducted a systematic review of the clinical effectiveness of randomised controlled trials (RCTs) and non-randomised comparative studies. Population-based case series were also reviewed for evidence of estimates of rare adverse events and fetal outcomes.

We searched MEDLINE, EMBASE, Cumulative Index to Nursing and Allied Health Literature, Cochrane Central Register of Controlled Trials, PsycINFO, Commonwealth Agricultural Bureaux (CAB) Abstracts, Latin American and Caribbean Health Sciences Literature, Allied and Complementary Medicine Database, British Nursing Index, Science Citation Index, Social Sciences Citation Index, Scopus, Conference Proceedings Index, NHS Economic Evaluation Database, Health Economic Evaluations Database, China National Knowledge Infrastructure, Cochrane Database of Systematic Reviews and Database of Abstracts of Reviews of Effects from inception up to September 2014. References from included studies and literature reviews were also examined. *Obstetric Medicine* was hand-searched, alongside websites of relevant organisations. The search strategy was based around nausea, vomiting and HG, and pregnancy terms. Costs were obtained from NHS sources.

All pharmacological and non-pharmacological interventions including novel treatments relevant to the NHS were considered. These included dietary/lifestyle interventions; vitamins such as vitamins B6 and B12; ginger; acupressure/acupuncture; hypnosis; antiemetic drugs (such as antihistamines; dopamine antagonists; 5-hydroxytryptamine receptor antagonists); corticosteroids; intravenous (i.v.) fluids; enteral feeding; and total parenteral nutrition.

Primary outcomes were severity of symptoms [such as Pregnancy-Unique Quantification of Emesis and Nausea (PUQE)]. Secondary outcomes included duration of symptoms; study-specific measures of NVP; quality of life; health-care utilisation; patient satisfaction; maternal weight; fetal outcomes [fetal or neonatal death, congenital abnormalities, low birthweight (< 2.5 kg), preterm birth (before 37 weeks' gestation) or small for gestational age (birthweight < 10th centile)]; adverse events, for example pregnancy complications (as reported in the study); costs; and cost-effectiveness. Both fixed- or random-effect model meta-analysis and a Bayesian mixed-treatment comparison were planned but were not performed due to heterogeneity in interventions, trial populations, reporting and definitions of outcome measures and methods. Thus, data on effectiveness, fetal outcomes and adverse events were tabulated and narratively reviewed.

Cost-effectiveness

The cost-effectiveness of the different treatments was planned to be assessed in an economic model but, due to the limited evidence, a simpler analysis considered the intervention costs, the difference in effectiveness implied if a more costly intervention was used. The perspective for cost was a health services perspective and all costs were reported in Great British Pounds (prices correct in 2014).

Results

Clinical effectiveness

Seventy-five papers from 73 studies met the inclusion criteria. For RCTs, 33 studies had a low risk of bias and 11 had a high risk of bias, with the remainder ($n = 20$) unclear. The non-randomised studies ($n = 9$) were judged low quality. There were 33 separate comparators. The most common comparisons were acupressure versus placebo ($n = 12$); steroid versus usual treatment ($n = 7$); ginger versus placebo ($n = 6$); ginger versus vitamin B6 ($n = 6$) and vitamin B6 versus placebo ($n = 4$). A common finding was that symptoms in all arms (including placebo) improved from baseline.

Ginger

Use of ginger was explored in 16 RCTs. The evidence available was at high or unclear risk of bias, in all but three trials. Six studies comparing ginger preparations with placebo generally reported evidence of ginger as improving a range of symptoms. Considering low risk of bias studies only, ginger still looked promising in reducing symptoms but the findings are not conclusive. One trial compared ginger with acupressure. Ginger again looked promising but the evidence was not very conclusive. For the comparison of ginger with vitamin B6 there are some higher-quality studies, but little evidence of a difference in the severity of symptoms between groups. There were few data for the comparisons of ginger with doxylamine–pyridoxine or antihistamine or metoclopramide, and little evidence suggesting any difference between groups. Overall, ginger might be better than placebo in reducing the severity of symptoms, but these data are limited to less severe symptoms.

Acupressure, acupuncture and nerve stimulation

Use of either acupuncture or acupressure was explored in 18 RCTs and one case series study. The quality of evidence available varied between low to high risk of bias. Comparisons with placebo were equivocal: two studies involving acupressure (both had mild symptoms and low risk of bias) reported better outcomes, but the remainder found no evidence of a difference or did not report NVP symptoms. The evidence for nerve stimulation was also mixed. Comparisons of traditional Chinese acupuncture and herbal medicine with Western medicine were at high risk of bias and impossible to emulate within the NHS. Overall, acupressure may reduce symptoms of nausea and retching in women with mild–moderate symptoms, but data were limited and inconclusive.

Aromatherapy

The evidence from two trials available for aromatherapy was at unclear risk of bias. There was no evidence of a difference compared with placebo or routine antenatal care.

Vitamin B6

Five studies considered the effectiveness of pyridoxine (vitamin B6), and they were at low risk of bias or risk of bias was unclear. Participants in all studies had mild to moderate symptoms at baseline. Comparisons of vitamin B6 preparations with placebo generally reported evidence of reduced symptoms of nausea, especially for women with more severe symptoms, and vomiting. Higher doses of vitamin B6 resulted in a greater improvement in symptoms. There was no evidence to suggest that vitamin B6 and metoclopramide as a combination treatment had an advantage over metoclopramide alone. Overall, there is a suggestion that vitamin B6 might be better than placebo in reducing the severity of symptoms especially at higher doses.

Vitamin B6 (pyridoxine)/doxylamine combination

Four studies compared the effectiveness of vitamin B6 and antihistamine with either placebo or ondansetron (and placebo). Two trials were at low risk of bias and two were at unclear risk of bias. Diclectin® [Duchesnay Inc.; doxylamine succinate (10 mg) plus pyridoxine hydrochloride (10 mg) slow release tablet] (vitamin B6 and antihistamine combination) appears more effective than placebo. Ondansetron appears more effective at reducing nausea than pyridoxine plus doxylamine, but with equivocal evidence for vomiting. Pre-emptive treatment with Diclectin before symptoms of NVP begin in women at high risk of severe NVP recurrence appears to result in a reduced risk of moderate–severe NVP compared with women who take Diclectin once symptoms begin.

Antihistamines

Of the three studies, two were at high risk of bias whereas one was at low risk. Participants in all studies had mild symptoms. Use of antihistamines resulted in an improvement compared with placebo or no treatment over a range of symptoms. The addition of vitamin B6 does not appear to improve effectiveness.

Dopamine antagonists

Dopamine antagonists were used in one trial (low risk of bias) and one poor-quality non-randomised study. There is limited evidence suggesting that promethazine is as effective as metoclopramide in reducing the symptoms of NVP.

Serotonin antagonists

Five trials and one case series study compared serotonin antagonists (ondansetron) against a range of alternatives. Three trials tested ondansetron against metoclopramide; symptoms were classified as mild to moderate in two trials and severe in one trial. The remaining two trials compared ondansetron with antihistamines with symptoms being moderate to severe. Only one trial was at low risk of bias. The studies comparing ondansetron with metoclopramide had mixed results, with both drugs improving symptoms. A study with low risk of bias found ondansetron more effective at reducing vomiting compared with metoclopramide after 4 days. Both ondansetron and antihistamine improve symptoms with no difference between effects. Overall, ondansetron reduces the severity of symptoms.

Intravenous fluids

Two studies were identified. One compared different compositions of i.v. solution (dextrose saline vs. saline only), which was at low risk of bias and one compared i.v. fluids containing vitamins with diazepam. I.v. fluid improves reported symptoms. Dextrose saline may be more effective at improving nausea over time for those with moderate nausea. Diazepam appears to be more effective than i.v. fluids alone at reducing nausea on day 2, but there was no evidence post treatment for those with moderate–severe nausea.

Transdermal clonidine

Evidence from one study with unclear risk of bias suggests that the use of transdermal clonidine patches looks promising for the treatment of severe HG.

Outpatient/day case management

The two studies of day case management were at low and high risk of bias respectively. Day case management of women with moderate to severe symptoms is feasible and acceptable, and as effective as inpatient management for some women.

Corticosteroids

The evidence available for corticosteroids was at low (three trials), unclear/high risk of bias (three studies) or weak (one case study). There was no evidence of a difference between either placebo or promethazine, but corticosteroids appeared to reduce vomiting episodes when compared with Phenergan® (Sanofi-Aventis) suppositories or metoclopramide.

Nasogastric/central/jejunoscopy feeding

Two case series studies of nasogastric and jejunoscopy feeding were identified for treatment of severe HG. Both were poor quality. Enteral feeding may be an effective but extreme method of supporting women suffering from very severe symptoms.

Gabapentin

One very small study which examined gabapentin therapy in women with HG was identified. Given the reported cases of congenital anomalies among the seven exposed infants, more research is needed.

Cost-effectiveness

No relevant economic studies were identified by the systematic review and the economic analysis was limited by lack of data. Estimates of costs for each therapy (both pharmacological and non-pharmacological) were derived and used to illustrate the benefits that would be implied if a more costly treatment was chosen over a less costly one. These data were set against the limited evidence base. For treatments initiated by women themselves, weekly costs of treatment ranged from £0.12 (vitamin B6) to £90 (hypnotherapy). For care prescribed by clinicians as third-line interventions, costs of treatment ranged from £1994 to £2115 (depending on combination of antiemetics and steroids used) if patients were admitted as inpatients. The total cost data were used to estimate the implied value for the benefits of treatment should a decision be made to adopt one treatment over another. The implied valuation showed the additional benefits that a more expensive treatment would need to provide in order to be considered a worthwhile use of resources. For patient-initiated interventions, the implied valuations ranged from 1.01 : 1.00 (vitamin B12 vs. vitamin B6) to 41 : 1 (hypnotherapy vs. ginger). For vitamin B12 versus vitamin B6, the interpretation is that vitamin B12 would need to provide at least 1% more in benefits to be considered cost-effective. Implied values were calculated for all comparators and related to evidence on clinical effectiveness, where available. These simple data on costs may be of use to stakeholders when judging what treatments to use.

Strengths and limitations

The main strength of the review was the comprehensively systematic approach to identifying studies investigating NVP and HG, which allowed us to identify all relevant studies across all levels of severity. This is a departure from the preplanned inclusion criteria of severe nausea and vomiting only, but it reflects the very limited evidence on severe symptoms and the fact that the overall quality of the evidence is either low or very low for all of the treatment comparisons made in the review for all severities. Quality was downgraded due to clinical heterogeneity, imprecision, a sparseness of data, or a combination of these factors. There was considerable variation as to how nausea and vomiting outcomes were recorded and considerable variation reporting of severity. This prevented the conduct of the planned meta-analysis and economic modelling. Another major limitation was the lack of comparisons of interventions of relevance to the NHS. Thus, we were restricted to a narrative review that, at best, was able to consider direction of effect.

Conclusions

Implications for health care

There appears to be evidence that some treatments (ginger, vitamin B6, antihistamines, metoclopramide) were better than placebo for mild symptoms, but there is little on the effectiveness of treatments in more severe NVP/HG. Evidence on differences in effectiveness were available for few other comparisons. Of note, however, was the finding that symptoms tended to improve after a few days (even with placebo). Therefore, if symptoms have not improved or not improved sufficiently after a short time, a change of treatment could be considered. Also of note was that day case management for moderate–severe symptoms is feasible, acceptable and as effective as inpatient care for some women. Overall, uncertainty exists about most of the estimates reported in the review and further research is very likely to have an important impact on our confidence in the findings of the review. The findings from the review provide little other evidence to help inform recommendations on the use of treatments for severe NVP/HG.

Recommendations for research

The main gaps in the evidence base are the lack of direct comparative studies of relevant treatments to the UK NHS and the consequent lack of robust data to estimate cost-effectiveness. The key research recommendations in order of importance are:

1. A RCT including economic evaluation to determine which second-line, hospital-prescribed therapy (i.v. rehydration with antihistamines, dopamine receptor antagonists or serotonin receptor antagonists) should be adopted as mainstream provision in the UK NHS.
2. A RCT including economic evaluation to test the use of subsequent treatments, such as steroids, as a third-line therapy. This could examine indication, effectiveness and dose of corticosteroids versus serotonin receptor antagonists (ondansetron).
3. A RCT including economic evaluation to determine which second-line, GP-prescribed therapy (e.g. vitamin B6–antihistamine combination vs. dopamine receptor antagonist) should be adopted in UK primary care.
4. In addition to the use of objective symptom scoring systems like the PUQE, consideration is needed as to what are the core outcomes of importance to women and further work.
5. The longer-term critical fetal and maternal outcomes (death, congenital abnormality) of all therapies used in the NHS need to be monitored and analysed to guide further research into stratified care.

Study registration

This study is registered as PROSPERO CRD42013006642.

Funding

Funding for this study was provided by the Health Technology Assessment programme of the National Institute for Health Research.

Chapter 1 Introduction and background

Background

Nausea and vomiting in pregnancy (NVP) is one of the most common symptoms of pregnancy affecting 50–85% of all women during the first half of pregnancy.¹ Symptoms usually start between 6 and 8 weeks of gestation, rise to a peak before the end of the first trimester and, in the majority of women, resolve by 20 weeks.² Most women (65–70%) self-manage their symptoms with avoidance of dietary triggers and oral hydration.² However, in the remainder, symptoms are more severe and/or protracted, leading to physical and psychosocial sequelae. These can include reduced quality of life (QoL), lost work time and negative effects on relationships with family and friends.³

The most severe form of NVP is referred to as hyperemesis gravidarum (HG), and is reported to affect 0.3–1.0% of pregnant women.¹ It is characterised by intractable vomiting, dehydration, ketosis, electrolyte imbalance, nutritional deficiencies and weight loss (usually defined as > 5% of pre-pregnancy weight). However, there is no widely accepted point at which NVP becomes HG. Likewise, the distinction between studies of women with NVP and HG is generally not possible as the degree of dehydration and weight loss prior to the intervention are rarely reported. Furthermore, although some studies report baseline symptom severity using a validated scale, this is insufficient to make a diagnosis of HG. For these reasons, study populations are seldom described as having HG, and are more frequently defined in terms of the severity of NVP. Therefore, for the purposes of this review, studies on interventions for both NVP and HG have been included.

Aetiology

The underlying pathophysiology of NVP/HG is poorly understood but is thought to involve a combination of biological, physiological, psychological and sociocultural factors.⁴

Genetic factors increase the risk of occurrence: results from a Norwegian study which included over 500,000 women found that the risk of HG was 15.2% in the second pregnancy of women who had a previous history of HG compared with 0.7% in women who did not [odds ratio (OR) 26.4, 95% confidence interval (CI) 24.2 to 28.7].⁵ The risk of developing HG is also increased threefold in the daughters of women who suffered from HG (unadjusted OR 2.9, 95% CI 2.4 to 3.6).⁶

Endocrine factors, especially higher levels of human chorionic gonadotropin, as is the case in multiple or molar pregnancies, have been associated with more severe forms of NVP/HG. A recent observational study found that free human chorionic gonadotropin and pappalysin-1 (also known as pregnancy-associated plasma protein A) were higher in women suffering from HG than in non-sufferers.⁷

Gestational transient thyrotoxicosis, has been reported in 60% of women suffering from HG⁸ and thyroid-stimulating hormone levels are raised in women with HG.⁹ Certain human chorionic gonadotropin subtypes can stimulate thyroid-stimulating hormone receptors and so contribute to the hyperthyroidism. The degree of hyperthyroidism has been found to correlate with the severity of NVP/HG.¹⁰ Higher levels of oestrogen, progesterone and leptin, and lower levels of adrenocorticotrophic hormone and prolactin have also been associated with HG.¹¹

Delayed gastric emptying related to relaxation of smooth muscle during pregnancy may influence NVP symptoms. Furthermore, higher rates of *Helicobacter pylori* infection have been noted in women suffering from HG:¹¹ in a meta-analysis of 25 studies investigating the association of *H. pylori* and HG, 14 studies demonstrated an increased risk of HG in infected women (with OR between 2.42 and 109.3), and 11 studies found no association.¹²

Lack of a definitive physiological trigger for HG has, in the past, led to numerous psychosomatic and psychological theories such as resentment or ambivalence towards the pregnancy, immaturity, conversion disorder, symptom of hysteria, neurosis or depression.¹³ It is now more commonly accepted that psychological afflictions are a consequence of the condition rather than a cause.^{4,9}

Impact on patients

Severe NVP causes emotional and psychological distress and can have a profound effect on a women's QoL, behavioural and cognitive function, affecting work capacity, household activities and interaction with children.^{14–16} Women with HG report feeling isolated, depressed and lonely, unable to cope with routine daily interactions or simple tasks. Two recent observational studies found higher incidences of depression, anxiety and stress in women diagnosed with HG compared with controls.^{17,18} Following cessation of symptoms the depression, anxiety and stress scores took several weeks to resolve,¹⁸ not returning to control values until the third trimester.¹⁷ However, in some women these psychological symptoms do not fully resolve and can result in post-traumatic stress disorder.¹⁹

As a result women make greater use of health-care resources. Based on Hospital Episode Statistics data for England, there were nearly 26,000 admissions for NVP/HG in 2010–11 with an average length of stay of 2 days.²⁰ These NHS costs are likely to underestimate the full costs as women may purchase a variety of products over the counter, pay for alternative therapies, receive treatment in primary care settings or as a hospital outpatient, and may incur extra child care, living costs and lost earnings. In addition, the associated increased risk of cognitive, behavioural and emotional dysfunction in pregnancy¹⁸ may prompt the use of further services and resources.

In the absence of a definitive cause, management of NVP/HG tends to focus on the alleviation of symptoms and prevention of serious morbidity. Typically, women are admitted to hospital, prescribed intravenous (i.v.) fluid therapy and antiemetic medication, but there is little time spent dealing with their psychological, social and emotional needs or providing information and guidance about the condition. The result is that women can feel unsupported, dissatisfied with care and experience negative interpersonal interactions with health-care providers.²¹

Finally, severe NVP/HG has implications for offspring. A recent systematic review and meta-analysis reported that women with HG were more likely to deliver preterm (OR 1.32, 95% CI 1.04 to 1.68) and to have a baby that was small for gestational age (OR 1.28, 95% CI 1.02 to 1.60), although there was no evidence of an association with congenital anomalies (pooled results from three studies: OR 1.17, 95% CI 0.68 to 2.03) or perinatal death (OR 0.92, 95% CI 0.61 to 1.41).²² A large Swedish birth cohort reported that women with HG who had their first admission in the second trimester were at increased risk of preterm pre-eclampsia (OR 2.09, 95% CI 1.38 to 3.16), placental abruption (OR 3.07, 95% CI 1.88 to 5.00) and to have a baby that was small for gestational age (OR 1.39, 95% CI 1.06 to 1.83), suggesting an association between HG and placental-mediated disease.²³

Assessment and diagnosis

The diagnosis of NVP/HG is made after excluding differential diagnoses, including gastrointestinal disorders, urinary tract infection, metabolic and endocrine disorders, drugs, psychological disorders (such as eating disorders) and other pregnancy-associated conditions (in particular molar pregnancy). However, there is currently no widely accepted approach to measuring the severity of symptoms in women. The most commonly used tools for the assessment of NVP/HG severity are presented in *Table 1*, with actual examples of the tools provided in *Appendix 1*.

TABLE 1 Tools used to measure the severity of NVP

Tool	Description
PUQE score	<p>Three questions regarding nausea, vomiting and retching during previous 12 hours (PUQE-24 = previous 24 hours)</p> <p>For each component: 0 = no symptoms, 5 = worst possible symptoms</p> <p>Maximum score = 15</p> <p>Scores of ≥ 13 indicate severe symptoms²⁴⁻²⁶</p>
RINVR	<p>Contains total of eight questions about duration/amount, frequency and distress caused by symptoms of nausea, vomiting and retching</p> <p>For each component: 0 = no symptoms, 5 = worst possible symptoms</p> <p>Maximum score = 40</p> <p>Scores of ≥ 33 indicated severe symptoms²⁷⁻²⁹</p>
McGill Nausea Questionnaire (measures nausea only)	<p>Contains a nausea rating index (nine sets of words which describe sensory, affective, evaluative and miscellaneous afferent feelings related to nausea that patients rank)</p> <p>An overall nausea index (0–5, where 0 = no symptoms, 5 = excruciating symptoms)</p> <p>Plus a VAS: 0 cm = no nausea, 10 cm = extreme nausea^{30,31}</p>
NVPI	<p>Three questions relating to nausea, retching and vomiting over the past 7 days</p> <p>For each component: 0 = no symptoms, 5 = worst possible symptoms</p> <p>Maximum score = 15</p> <p>A score of ≥ 8 indicates severe symptoms^{32,33}</p>
VAS	<p>Patients rate their symptoms on a scale of 0–10, where 0 = no symptoms, 10 = extreme symptoms</p>

NVPI, Nausea and Vomiting of Pregnancy Instrument; PUQE, Pregnancy-Unique Quantification of Emesis and Nausea; RINVR, Rhodes Index of Nausea, Vomiting and Retching; VAS, visual analogue scale.

However, although the measurement of NVP/HG symptom severity is the main aim for women and practitioners, other wider outcomes are also relevant when assessing the broader effectiveness of interventions. Thus, key secondary outcomes in studies to date have included both measures related to maternal physical and psychosocial health, and fetal or neonatal outcomes (*Table 2*).

Current interventions for nausea and vomiting in pregnancy/hyperemesis gravidarum

For the purposes of this report, interventions are considered in three broad groups:

- First-line interventions, usually initiated by women before seeking medical care and hence tend to be used in less severe NVP.
- Second-line interventions, typically prescribed when a women presents to medical care. Initially this is likely to be a general practitioner (GP) in primary care but may involve referral of women with more severe symptoms for inpatient, outpatient or day case care in hospital.
- Third-line interventions, reserved for women in hospital with persistent or recurrent symptoms despite second-line therapies.

TABLE 2 Secondary outcome measures

Maternal: physical	Maternal: psychosocial	Fetal/neonatal
Admission/readmission rate	QoL (SF-12 or SF-36 score)	Congenital abnormality
Length of hospital stay	General Health Questionnaire	Low birthweight (< 2.5 kg)
Antiemetic/other medication use	Pregnancy-specific QoL measure	Small for gestational age (< 10th centile)
Amount/duration i.v. fluid administration	NVP specific questionnaire	Preterm birth (before 37 weeks' gestation)
Enteral/TPN	Satisfaction with care	5-minute APGAR score
Side effects	Direct costs to woman/family	Stillbirth/intrauterine death
Economic costs (hospital/medical care)	Time lost from work	Neonatal death
Adverse pregnancy outcomes	Edinburgh Postnatal Depression Scale	Spontaneous miscarriage
Weight loss		Admission to special care baby unit
Therapeutic termination of pregnancy		Infant development outcomes

APGAR, American Pediatric Gross Assessment Record; SF-12, Short Form questionnaire-12 items; SF-36, Short Form questionnaire-36 items; TPN, total parenteral nutrition.

The relationship between these three intervention groups is described in *Figure 1*, with key individual interventions described in detail in the following sections.

Patient-initiated first-line interventions

When first experiencing the symptoms of NVP, women often access information, advice and services from a variety of sources. Information is readily available regarding simple lifestyle changes, dietary modifications and alternative therapies via the internet and in pregnancy magazines. 'Self-help' interventions also include a range of supplements that are available 'over the counter'. Many women try one or more of these before seeking medical advice.

Dietary/lifestyle interventions

Women report using a range of dietary/lifestyle interventions (e.g. increasing oral fluid intake, eating small frequent meals, eating bland foods/protein-predominant meals and avoiding spicy, odorous and fatty foods, and stopping iron-containing multivitamins).^{2,34}

Vitamins

Vitamins are vital nutrients. They are available over the counter as single vitamin or multivitamin preparations.

Vitamin B6 (pyridoxine) A water-soluble vitamin essential for many metabolic processes within the body. Usually taken in doses of 10–50 mg up to four times daily to treat NVP.

Vitamin B12 (cyanocobalamin) A water-soluble vitamin essential for normal function of the nervous system, red blood cell formation and many other metabolic processes.

Ginger

Ginger (*Zingiber officinale*) is considered a food supplement (not a drug) and is available in several preparations; powdered fresh root, tablets, capsules and syrup. Its anti-nausea properties were first described in traditional Chinese medicine.³⁵

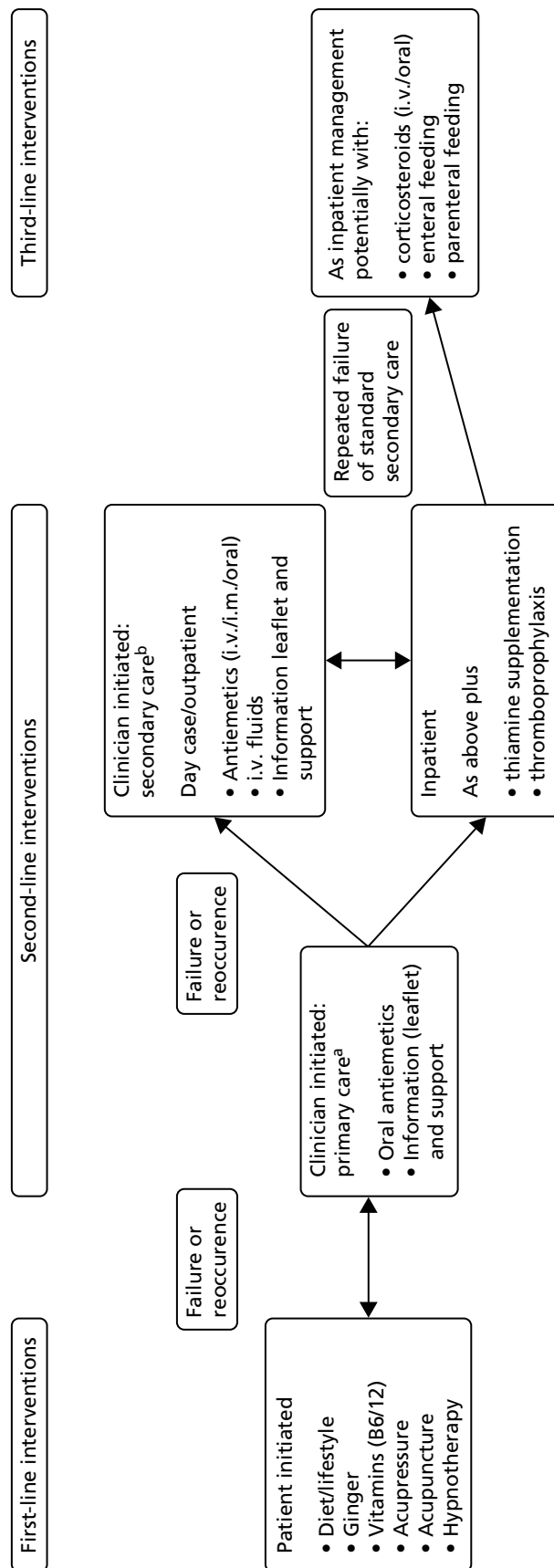


FIGURE 1 Treatments for NVP. a, Care may also involve urine \pm blood tests, weight and maternal observations; and b, care will involve urine and blood tests, weight, maternal observations and pelvic ultrasound. i.m., intramuscular.

Acupressure/acupuncture

Acupressure involves the application of physical pressure to specific acupuncture points; with respect to NVP this involves the pericardium 6 (P6) point near the wrist.

Acupuncture involves the manipulation of thin needles inserted into acupuncture points in the skin.

Hypnotherapy

Hypnotherapy employs direct suggestion of symptom removal with the subject under hypnosis.

Aromatherapy

Aromatherapy was first used by ancient civilisations for cosmetics, perfumes and drugs. It involves the use of plant materials, aromatic plant and essential oils to alter mood, cognitive, psychological or physical well-being. Oils can either be applied topically via massage, via inhalation or via emersion mixed with water. Common uses include stress and anxiety relief, to uplift mood or counter depression. Evidence surrounding efficacy and safety remains unclear for some treatments.

Clinician-prescribed second-line interventions

Second-line interventions tend to be used for more severe symptoms either instead of or, less frequently, in addition to, first-line interventions. These may be initiated either in primary care by the GP or in a secondary care hospital setting.

Antiemetic drugs

Antiemetic drugs include antagonists to histamine, acetylcholine, dopamine and 5-hydroxytryptamine (5-HT₃) receptors in the chemoreceptor trigger zone, vestibular apparatus and visceral afferents. Dyspepsia symptoms which often accompany NVP are also often treated with H₂ receptor blockers (e.g. ranitidine) or proton pump inhibitors (e.g. omeprazole).²

Antihistamines (H₁ receptor blockers) are probably the most widely used antiemetics and include doxylamine, meclizine, diphenhydramine, hydroxyzine, dimenhydrinate and cyclizine. Doxylamine is sometimes used in combination with vitamin B6 (pyridoxine). This combined therapy could be used as a treatment option in countries where Diclectin® (Duchesnay Inc.; delayed release doxylamine, 10 mg, plus pyridoxine, 10 mg, available in Canada and the USA but not the UK) is not available.³⁶

Dopamine antagonists are known to stimulate gastrointestinal motility, so encouraging the transit of substances through the stomach. They also work centrally by antagonising the action on D₂ receptors in the chemoreceptor trigger zone. Several phenothiazines including promethazine and prochlorperazine have been used to treat NVP/HG. Other drugs in this class used to treat NVP/HG include metoclopramide, domperidone, droperidol and trimethobenzamide.

5-HT₃ receptor antagonists (selective serotonin receptor antagonists) are commonly used to treat chemotherapy and post-operative induced nausea and vomiting which is caused by release of 5-HT₃ from the upper small intestine. The action of 5-HT₃ receptor antagonists are mediated through the central chemoreceptor trigger zone and peripheral (intestinal and spinal) 5-HT₃ receptors.

Intravenous fluids

Administration of i.v. fluids treats the consequences of NVP/HG rather than the symptoms. Women who are severely dehydrated and ketotic need hospital admission and i.v. fluid and electrolyte replacement. This is routinely carried out in either a day care 'outpatient' setting or on an inpatient ward.

Clinician-prescribed third-line interventions

Third-line interventions are reserved for women who have severe and persisting symptoms and associated weight loss and dehydration (although latter may have been corrected). Although commenced while the women are in hospital, some of these interventions may be continued on an outpatient basis.

Corticosteroids

Steroids are being increasingly used in refractory cases of NVP/HG which have been unresponsive to other treatments (i.v. hydrocortisone 100 mg twice daily, followed by oral prednisolone 40–50 mg, reducing to a maintenance dose).

Enteral feeding and total parenteral nutrition

Enteral feeding refers to the delivery of nutrients directly into the stomach, duodenum or jejunum. For women who cannot tolerate enteral nutrition, the use of total parenteral nutrition (TPN) has been reported in case series but use is associated with significant maternal morbidity.³⁷

Interventions presented in the report but not routinely used to treat nausea and vomiting in pregnancy

Diazepam

Diazepam is a benzodiazepine drug used to treat, for example, anxiety, panic attacks, insomnia and seizures. It enhances the effects of the neurotransmitter gamma-aminobutyric acid which leads to central nervous system depression. Its use results in sedation, long-term use results in physical dependence.

Clonidine

Clonidine is a centrally acting α_2 adrenergic agonist and imidazoline receptor agonist. It is usually used to treat hypertension, attention deficit hyperactivity disorder and, less commonly, anxiety disorders, withdrawal, migraine and certain chronic pain conditions. Observational data suggests that it may be effective in the treatment of refractory nausea and vomiting.³⁸

Gabapentin

Gabapentin was originally synthesised to mimic the action of the neurotransmitter gamma-aminobutyric acid, but acts on various brain receptors. It is generally used to treat seizures and neuropathic pain, with less common uses including the treatment of generalised anxiety disorders, restless leg syndrome and itching caused by various aetiologies. It has previously been associated with improvements in refractory nausea in a small study of breast cancer patients.³⁹

Current guidance and use of therapies within the NHS

Currently there are no national guidelines within the NHS pertaining to NVP/HG; however, the Royal College of Obstetricians and Gynaecologists are in the process of producing said guidelines which should be published in early 2016. Initially, GPs may try different antiemetics before referring women to hospital. Traditionally, secondary care would involve admission to either an antenatal or gynaecology ward for treatment with i.v. fluids, antiemetics and vitamin supplements. Oral intake would gradually be resumed followed by discharge back into the community. Resumption of symptoms would result in readmission and a repeat of previous care, possibly trying different antiemetics or a combination thereof.

Increasingly, more obstetric and gynaecology units are using 'day case' management as the first option for initial referrals. Care usually involves some form of rapid rehydration and treatment with an i.v. antiemetic, followed by discharge with oral antiemetics, ideally with advice, support and guidance regarding self-help measures. However, assessing symptom severity, as well as the packages of care, vary substantially and lack a strong evidence base.

When available, day case, outpatient management does result in fewer admissions to hospital. Consequently, women who are admitted tend to be suffering from more severe symptoms. These women are likely to have had repeated hospital attendances and to have tried a number of different combinations of interventions. This latter group of women are likely to experience persistent severe symptoms, weight loss, electrolyte imbalance and failure to cope. In some of these women corticosteroid therapy may be

considered an appropriate option when more conventional options have failed. In rare circumstances where this proves unsuccessful, enteral or parenteral nutrition may be instigated and, as a last resort, some women will opt for termination of pregnancy.

Aims and objectives

This study aimed to:

- review systematically the evidence of the clinical effectiveness and cost-effectiveness of each treatment for NVP/HG
- determine which therapies are most likely to be cost-effective for implementation into the UK NHS
- identify and prioritise future research needs.

Structure of the report

The following chapter (see *Chapter 2*) describes the methods employed for the systematic review and synthesis of evidence for interventions for HG and/or NVP. *Chapter 3* provides an overview of the identified evidence, including the quality of the included studies, and a brief discussion of the issues that arose in attempting to synthesise the emergent data. *Chapters 4–17* detail the findings for each individual intervention, focussing on the evidence for their effectiveness in terms of nausea, vomiting and retching. *Chapter 18* presents the methods and results of the economic evaluation. Key issues considered likely to be important from the perspective of both patients and health-care practitioners are described in *Chapters 19* and *20*. The implications of the results of this review are discussed in depth in *Chapter 21*, with the final conclusions outlined in *Chapter 22*.

Chapter 2 Methods for the systematic review of effectiveness

General methodology

The systematic review followed the approach suggested by the Evidence for Policy and Practice Information and Co-ordinating Centre at the Institute of Education, London. The review protocol was registered with PROSPERO, the International Prospective Register of Systematic Reviews,⁴⁰ and it aimed to systematically appraise and summarise the evidence on available interventions for NVP/HG within the three broad groups described in *Chapter 1*:

- first-line interventions
- second-line interventions
- third-line inpatient interventions.

The review examined the evidence for these groups of interventions in relation to their clinical effectiveness and associated adverse events, and their cost-effectiveness.

Inclusion criteria

Types of studies

Randomised controlled trials (RCTs), non-randomised comparative studies and population-based case series were deemed eligible for inclusion. The latter design was included primarily to facilitate calculation of estimates of rare adverse events and fetal outcomes, and for treatments reserved for the most severe cases such as TPN.

Types of participants

Participants were women experiencing nausea, vomiting and/or retching in pregnancy where recruitment to a trial took place before 20 weeks' gestation. As HG is difficult to differentiate from severe or intractable NVP, two approaches were used initially to identify relevant populations of women. First, studies were selected where their study samples were reported as suffering severe symptoms using published scales and cut-points for severity [e.g. Pregnancy-Unique Quantification of Emesis and Nausea (PUQE)²⁵ ≥ 13 , the Rhodes Index of Nausea, Vomiting and Retching (RINVR)²⁷ ≥ 33]. These cut-off points are well correlated.²⁵ For studies of mixed levels of severity, the study was included if $> 80\%$ of participants exceeded these cut-offs. Second, studies were selected if, using the authors' definition, women in the study sample were defined as having severe symptoms. Similarly, studies were included if $> 80\%$ of the sample met this definition. However, due to the inconsistent application of severity scales both within and across studies, and to ensure completeness, a broader selection of studies was deemed eligible for inclusion than originally anticipated. Details of the method used by authors to define severity were recorded for all eligible studies.

Types of interventions and comparators

All pharmacological and non-pharmacological interventions relevant to the NHS for delivery in the community, and in hospital, either as an inpatient or an outpatient, were deemed relevant for this evidence synthesis. These interventions included dietary/lifestyle interventions; vitamins such as vitamin B6 and vitamin B12; ginger; acupressure/acupuncture; hypnosis; antiemetic drugs [such as antihistamines; dopamine antagonists, hydroxytryptamine (5-HT₃) receptor antagonists]; corticosteroids; i.v. fluids; enteral feeding; and TPN. Studies were included that had a comparative group for assessment of relative effectiveness. This was either a no treatment group, a treatment as usual group or an alternative intervention group. For the treatment as usual

group we endeavoured to clearly define what this entailed. For rare fetal or adverse outcomes and for studies investigating treatments for women with the most severe symptoms (e.g. TPN), no comparator group was defined as the target studies were population-based series.

Types of outcome measures

Primary outcomes

Severity of symptoms [such as PUQE,²⁵ RINVR,²⁷ McGill Nausea Questionnaire,³⁰ Nausea and Vomiting of Pregnancy Instrument (NVPI) and³⁴ visual analogue scales (VASs)^{41–43}] (see *Table 1*).

Secondary outcomes

Duration of symptoms (reported period of symptoms, date of symptom relief); study-specific measures of NVP; health-related QoL; health-care utilisation (including admission and length of stay of the woman, readmission to hospital of the women, admission and length of stay on special care baby units); patient satisfaction; maternal weight; fetal outcomes [fetal or neonatal death, congenital abnormalities, low birthweight (< 2.5 kg), preterm birth (before 37 weeks' gestation) or small for gestational age (birthweight < 10th centile)]; adverse events, for example pregnancy complications (as reported in the study), but including haemorrhage, hypertension, pre-eclampsia and proteinuria; costs (as defined by the study authors); and cost-effectiveness (as defined by the study authors) (see *Table 2*).

Search strategy

The search strategy was designed and executed by an experienced information specialist in collaboration with the rest of the research team. The original protocol stated that the search strategy would combine the two main conditions of pregnancy and NVP/HG with associated interventions and QoL outcomes. However, given both the extensive array of interventions used to address NVP/HG, and the relatively small available literature in this field, it was subsequently decided not to restrict the scope of the review by including key interventions or outcomes as search terms. Although such a strategy increases the number of papers to be reviewed; it minimises the risk of missing any relevant studies. The search was therefore structured around two core concepts: (1) nausea, vomiting and HG; and (2) pregnancy. Key words for both concepts were coupled with relevant medical subject heading (MeSH) and thesaurus terms. The search strategy was designed in MEDLINE and translated as appropriate to the other databases. All terms were truncated as appropriate and variant spellings were used. In order to reduce the number of studies returned, search filters for the relevant study types (RCTs or case series studies) were applied where possible. No time or language limit was set.

The full list of search terms for MEDLINE is presented in *Table 3*.

We searched the following electronic bibliographic databases on the dates described below, with update searches executed between the 11 and 16 September 2014 unless otherwise stated. As our initial scoping search highlighted the number of complementary medicine interventions for NVP/HG, the search was extended to include additional key databases of non-English-language studies⁴⁴ (Latin American and Caribbean Health Sciences Literature and China National Knowledge Infrastructure).

- MEDLINE (Ovid) 1946–November 2013, searched 10 December 2013.
- EMBASE (Ovid) 1980–2013 week 50, searched 12 December 2013.
- Cochrane Central Register of Controlled Trials (Wiley) issue 11 2013, searched 16 December 2013.
- Cochrane Database of Systematic Reviews (Wiley) issue 11 2013, searched 16 December 2013.
- Database of Abstracts of Reviews of Effects (Wiley) issue 11 2013, searched 16 December 2013.
- Cumulative Index to Nursing and Allied Health Literature (EBSCOhost) 1981–November 2013, searched 17 December 2013.

TABLE 3 List of search terms

A. HG	B. Pregnancy	C. Nausea/vomiting	D. Exclusions
exp Morning Sickness/	Pregnancy/	Nausea/	Animals/ not humans
	Pregnancy Complications/	Vomiting/	Letter/
		Sialorrhea/	Editorial/
			News/
			exp Historical Article/
			Anecdotes as Topic/
			Comment/
(morning sickness or hyperemesis gravidarum).ti,ab.	(pregnan\$ adj5 (sick or sickness or nause\$ or vomit\$ or retch\$ or dry heave or heaving or emesis or hyperemesis or ptyalism or hypersalivat\$ or sialorrh\$ or spitting)).ti,ab.		

The search carried out was [A or (B and C)] not D.
 /= MeSH heading (translated in other databases where possible).
 exp = explode the MeSH heading.
 ti = term in the title field.
 ab = term in the abstract field.
 \$ = truncation.
 adjx = proximity, terms must be within x words of each other.

- British Nursing Index (NHS Healthcare Databases) 1992–January 2014, searched 22 January 2014.
- PsycINFO (Ovid) 1806–December week 2 2013, searched 17 December 2013.
- Commonwealth Agricultural Bureaux (CAB) Abstracts (Ovid) 1910–2013 week 49, searched 18 December 2013.
- Latin American and Caribbean Health Sciences Literature (<http://regional.bvsalud.org>), searched 18 December 2013.
- Allied and Complementary Medicine Database (NHS Healthcare Databases) 1985–January 2014, searched 22 January 2014.
- Science Citation Index (Web of Knowledge) 1970–November 2013, searched 18 December 2013.
- Social Science Citation Index (Web of Knowledge) 1970–November 2013, searched 18 December 2013.
- Scopus, searched 8 January 2014.
- Conference Proceedings Index – Science (Web of Knowledge) 1990–November 2013, searched 18 December 2013.
- ClinicalTrials.gov searched 8 January 2014.
- NHS Economic Evaluation Database (Wiley) issue 11 2013, searched 16 December 2013.
- Health Economic Evaluations Database (Wiley), searched 30 January 2014.
- China National Knowledge Infrastructure (<http://eng.cnki.net/grid2008/index.htm>), searched 14 March 2014.

In addition, the reference lists of included papers and key relevant literature reviews identified during the search process were also examined for additional relevant studies, and *Obstetric Medicine* vol. 1(1) (September 2008) and vol. 7(2) (June 2014) were hand-searched. Furthermore, the following websites of relevant organisations were also searched in order to source as much unpublished literature as possible:

- Motherisk (URL: www.motherisk.org/women/drugs.jsp; accessed September 2014).
- American Congress of Obstetricians and Gynecologists (URL: www.acog.org/; accessed September 2014).

- Pregnancy Sickness Support (PSS) (URL: www.pregnancysicknesssupport.org.uk; accessed September 2014).
- National Institute for Health and Care Excellence Clinical Knowledge Summaries (URL: <http://cks.nice.org.uk/nauseavomiting-in-pregnancy>; accessed September 2014).
- Hyperemesis Education and Research (URL: www.helper.org/health-professionals/treatments/index.php; accessed September 2014).
- UK Teratology Information Service (UKTIS) (URL: www.uktis.org/) including checking the references of the following relevant documents:
 - treatment of NVP (December 2013)
 - use of promethazine in pregnancy (October 2010)
 - use of vitamin B12 in pregnancy (September 2013)
 - use of pyridoxine (vitamin B6) in pregnancy (January 2011)
 - use of ginger in pregnancy (March 2013).
- European Medicines Agency (URL: www.ema.europa.eu/ema/; accessed September 2014).
- ProQuest Dissertations and Theses – UK & Ireland (URL: www.theses.com/; accessed September 2014).
- e-thesis online service (URL: <http://ethos.bl.uk/>; accessed September 2014).
- Trip (URL: www.tripdatabase.com/; accessed September 2014).
- System for Information on Grey Literature in Europe (URL: www.opengrey.eu/; accessed September 2014).
- Google Scholar (URL: <https://scholar.google.co.uk/>; accessed September 2014).

First exclusion process

All records were imported into a bibliographic referencing software programme (EndNote v.X7; Thomson Reuters, CA, USA). Duplicate records were identified and deleted. The remaining references were assessed for relevancy by two independent investigators on the basis of the title and abstract (or title only if abstract not available). Papers were considered relevant to the systematic review if they met the inclusion criteria detailed in *Inclusion criteria*.

All of the titles and abstracts of all references were read by both investigators and classified as potentially eligible, not eligible or unclear within EndNote. Reconciliation of the resultant EndNote databases was conducted via Microsoft Excel 2013 (Microsoft Corporation, Redmond, WA, USA) and any discrepancies were discussed. In case of doubt, papers went through to the next stage of the exclusion process. Full-text copies of those papers identified as of potential relevance were obtained.

Second exclusion process

All full-text English-language papers obtained were assessed by two investigators independently and classified as relevant, not relevant or unclear on the basis of the same inclusion criteria. Any disagreements at this stage were resolved by discussion between the two researchers. Full-text papers published in languages other than English (German, French, Portuguese, Arabic, Chinese, Korean, Danish and Spanish) were assessed by native speakers of the relevant languages, working alongside one of the two investigators to ensure consistency and adequate compliance against the specified inclusion criteria. Tables of studies excluded at this stage were prepared, detailing reasons for exclusion.

Data extraction

Information from all papers identified as meeting the specified inclusion criteria was extracted using a structured data abstraction form. Key data extracted from eligible papers included:

- i. study characteristics (bibliographic details, setting, intervention type, study population including definition of severity)
- ii. methodology and reporting
- iii. quantitative findings and conclusions.

For English-language papers, data extraction was carried out by one researcher and checked by another. For papers published in languages other than English, data extraction was carried out by a native speaker working alongside an investigator to identify and translate the relevant information. The data abstraction form for clinical effectiveness is presented as *Appendix 2*.

Risk of bias in included studies and quality assessment

The quality of the included studies was evaluated in accordance with the comprehensive approach advised by the Grading of Recommendations Assessment, Development and Evaluation (GRADE) Working Group.⁴⁵ The risk of bias of included RCTs was assessed with the Cochrane Collaboration's tool⁴⁶ (see *Appendix 3* for full details). This included assessment of sequence generation; allocation concealment; blinding; selective reporting of outcomes; incomplete outcome data; and other possible sources of bias. For the incomplete outcome data item, we coded the satisfactory level of loss to follow-up for each outcome as 'low risk of bias', if fewer than 20% of patients were lost to follow-up and reasons for loss to follow-up were similar in all study arms. Disputes were resolved by discussion with another member of the review team. The risk of bias for case series studies was assessed using the component-based tool developed by the Effective Public Health Practice Project (EPHPP), Canada⁴⁷ (see *Appendix 4*), which possesses a relatively high degree of inter-rater reliability in comparison to alternative tools.^{48,49} For English-language papers, quality assessment was conducted independently by two investigators, with any disputes resolved by discussion. For papers published in languages other than English, quality assessment was carried out by a native speaker working alongside an investigator.

We had initially used an objective approach to make decisions about overall risk of bias; if at least one item was adjudged as being at high risk of bias then the trial was given a high risk of bias overall rating. Similarly, a trial that had no items scored as being at high risk of bias, but at least one was at an unclear risk of bias, then the trial was deemed to be at an unclear risk of bias. This meant that only trials that were scored at low risk of bias for all six risk of bias items could get an overall low risk judgement. Instead, we used a robust approach to assess overall risk of bias in a study, and which did not simply rely on a vote counting approach. This approach gave careful consideration to all six risk of bias item judgements, and the impact of unclear and high risk of bias in individual items, given the scope and context of the trial. For example, if a study had adequately addressed five of the six individual risk of bias items, but a placebo-controlled trial was unblinded in some capacity (either patients, personnel or both), then we scored this trial as being at high overall risk of bias, as this flaw in the conduct of such a trial could seriously impact on the results. However, in some trials blinding was not possible due to the type of interventions being compared so although the individual 'blinding' risk of bias item was scored as being at high risk of bias, this did not necessarily mean the overall risk of bias in the study would follow. If other items were generally at low risk of bias, and we deemed blinding to be of little relevance, then we scored the study as being at low overall risk of bias. Justification for overall decisions has been provided where necessary.

In addition, two researchers independently assessed all included studies for the potential for imprecision, inconsistency and indirectness of results, using GRADE guidelines.⁵⁰⁻⁵⁴ Summary of findings (SoF) tables were not presented due to the narrative nature of the review and the heterogeneity of the outcomes.

Data synthesis

First, the range of interventions, populations and outcomes that have been studied were described. The direction and size of the reported effects from effectiveness studies were presented overall, as well as grouped according to population, intervention type, outcomes and study design. Results are summarised in tables. Groups of studies using similar definitions of severity were identified, based on the data extracted and expert opinion. A coding frame was developed for the different definitions used, which was checked by the second systematic reviewer. Two clinical specialists within the research team then grouped the studies into the coding frame. The grouping produced was compared and any discrepancies, including definitions that did not fit into the coding frame, were resolved by discussion.

Meta-analysis was considered to be inappropriate due to heterogeneity, as will be illustrated in the overview of included studies chapter (see *Chapter 3*). This judgement was made after consideration of interventions, trial populations, and especially the reporting and definitions of outcome measures and methods. We also explored whether data from different studies could be transformed on to a common scale (e.g. symptom severity might be recoded into number no longer experiencing severe symptoms) using imputation and subject to sensitivity analysis and methods, but this was not possible. Therefore it was not necessary for the team to investigate the validity of performing mixed-treatment (indirect) comparisons, using appropriate methods to compare interventions that have not been compared directly with each other.^{55,56} Heterogeneity could not be assessed by visualisation of results or, where relevant in statistical terms, by the chi-squared test for homogeneity and the I^2 statistic as specified a priori. Instead, a narrative synthesis was conducted, with the effects split into numerous categories and classes of intervention comparison described in the following chapter (see *Chapter 3, Interventions and comparators* and *Table 6*). Trials with more than two randomised groups may appear in more than one category depending on the comparisons made.

Data on effectiveness, fetal outcomes and adverse events were tabulated and described narratively, including variation in the form, setting, study population and delivery of the interventions. Given the inconsistencies in the application of both published and author-defined severity scales, studies were recategorised by clinical experts on the review team according to whether the participants were predominantly suffering from mild, moderate or severe NVP/HG. The effects are generally presented in terms of whether or not there were statistically significant differences between randomised groups at the last time point at which outcomes were assessed. However, where possible, magnitude of effects was reported such as a mean difference or risk ratio with corresponding 95% CIs. All studies are included in the narrative synthesis, irrespective of their risk of bias but the weight of evidence is discussed accordingly. Where necessary, comments are made in the text to advise caution for serious methodological shortcomings as well as applying the GRADE approach⁵⁰⁻⁵⁴ in the overall assessment of the quality of the evidence, although SoF tables were not constructed (see *Risk of bias in included studies and quality assessment*).

Owing to limited available data, it was not possible to examine publication bias using funnel plots as specified in the original review protocol. We were also unable to conduct subgroup analyses to explore the variation with pre-determined factors (e.g. the setting in which the intervention was applied and the severity and duration of symptoms at baseline) or sensitivity analyses to explore the impact of study design, including variation in definitions of outcomes, on measures of effectiveness due to lack of suitable data.

Chapter 3 Clinical effectiveness: overview of included studies

Studies identified

A flow chart of the studies is shown in *Figure 2*. In total, 11,830 papers were identified from the combination of standard electronic databases ($n = 11,659$), specialist Chinese databases ($n = 102$) and various sources of grey literature ($n = 69$). Of these, 5152 duplicate papers were identified and deleted (5150 from the standard electronic databases, and two from the grey literature).

The deletion of duplicate papers left 6678 individual papers for assessment. After screening titles and abstracts, 322 papers were identified as of potential relevance and full-text copies of 309 papers were obtained (with the remainder unobtainable). Of these, 96 were judged ineligible for the effectiveness review and immediately excluded (narrative overviews, systematic literature reviews or economic evaluations). After the second exclusion process, comprising more detailed reading of each full-text paper, a further 138 papers were judged not to meet the inclusion criteria of the review and were also excluded.

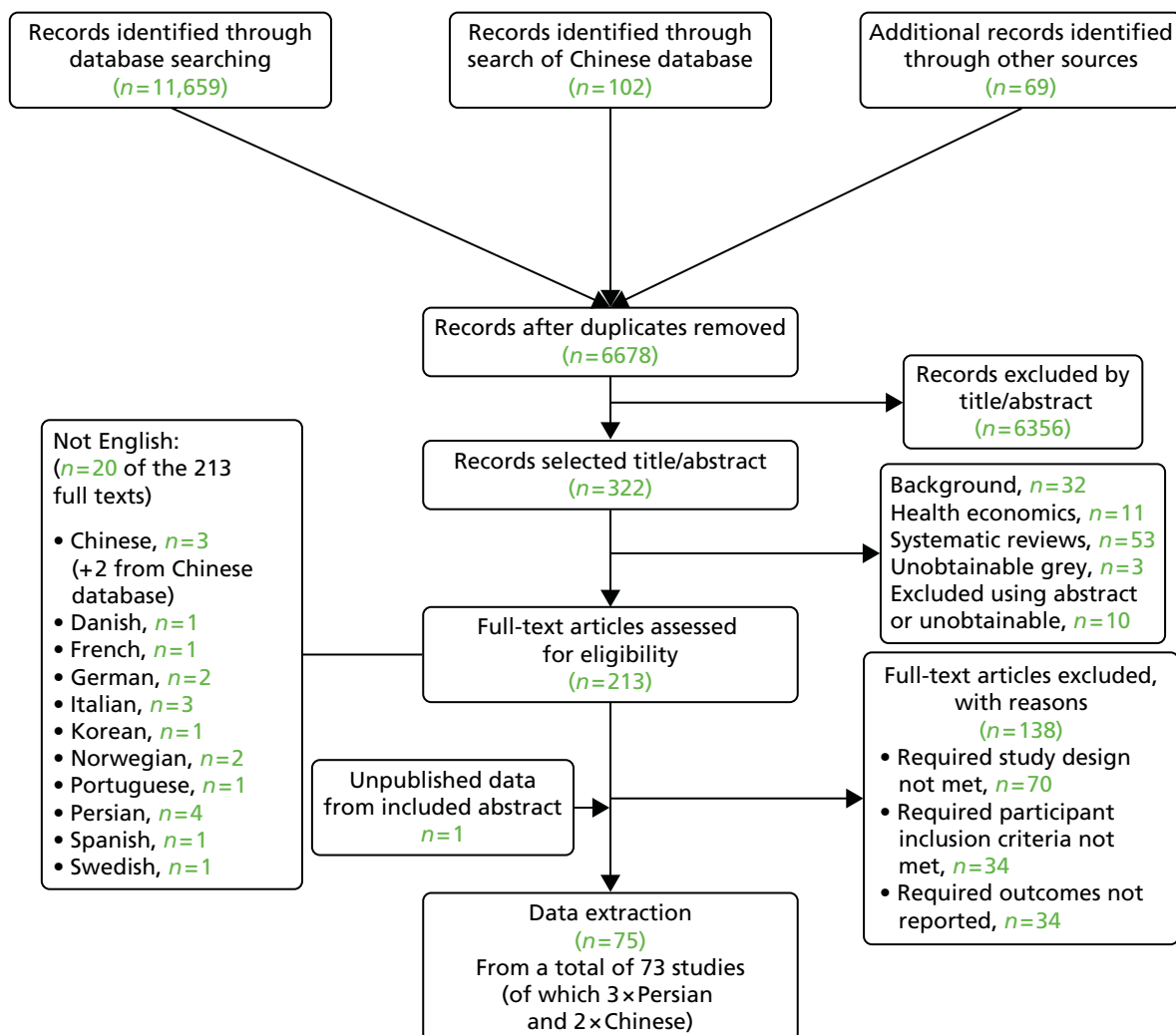


FIGURE 2 Flow chart of clinical effectiveness literature.

Key reasons for exclusion were duplicate paper already included; participant inclusion criteria for the identified study judged not relevant to our review; did not include any of the pre-specified outcomes; or ineligible study design (no comparator group).

As a result, 75 papers were identified for data extraction, from a total of 73 separate studies. A full list of included studies is provided as *Appendix 5*. A table of excluded studies detailing reasons for exclusion is provided as *Appendix 6*.

Quality of included studies

Randomised controlled trials

Overall risk of bias

The results of the quality assessment procedure for the 64 included RCTs (reported in 66 papers) are displayed in *Figure 3* and *Table 4*. There was variation both in terms of the quality of the studies and the quality of the reporting. In a large number of papers, there was insufficient detail provided to permit clear judgement of risk of bias in a range of key areas. Overall, 33 RCTs were classed as having low within-study risk of bias, 11 RCTs were classed as having high within-study risk of bias, and the remainder ($n = 20$) were classed as unclear in this respect. The high proportion of studies at unclear risk of bias was due to poor reporting and a lack of detail, particularly in the methods section. There were also a number of publications in abstract form only. As an unclear judgement was often due to poor reporting rather than specific methodological concerns, it was not judged appropriate to categorise studies with those deemed at high risk of bias as a result of more serious methodological flaws. Our robust approach to the assessment of the overall risk of bias within individual studies is described in more detail in *Chapter 2, Risk of bias in included studies and quality assessment*. More detail is provided below to illustrate the range in quality in terms of each individual component of the Cochrane’s risk of bias tool.⁴⁶

Random sequence generation

The risk of bias arising from the method of generation of the allocation sequence was low in 39 of the included RCTs.^{13,41,57,60,62–64,66,67,69,70,76,80–83,85–89,91,93,94,98–104,106–108,110,112–115} Methods employed included random number tables, computer-generated sequence generation^{61,63,64,81–83,88,93,94,98,99,106,108,112,114} and randomised block design.^{62,67,80,87,89,91,101,102,107,113,115,117} One trial was classed as high risk because women were asked to draw an envelope from a box with the same appearance but with different contents.¹¹¹ It was categorised as unclear in the remaining 24 RCTs due to insufficient information provided by the authors to permit judgement either way.^{42,43,58,59,65,68,71–75,77–79,84,90,92,95–97,104,105,109,116}

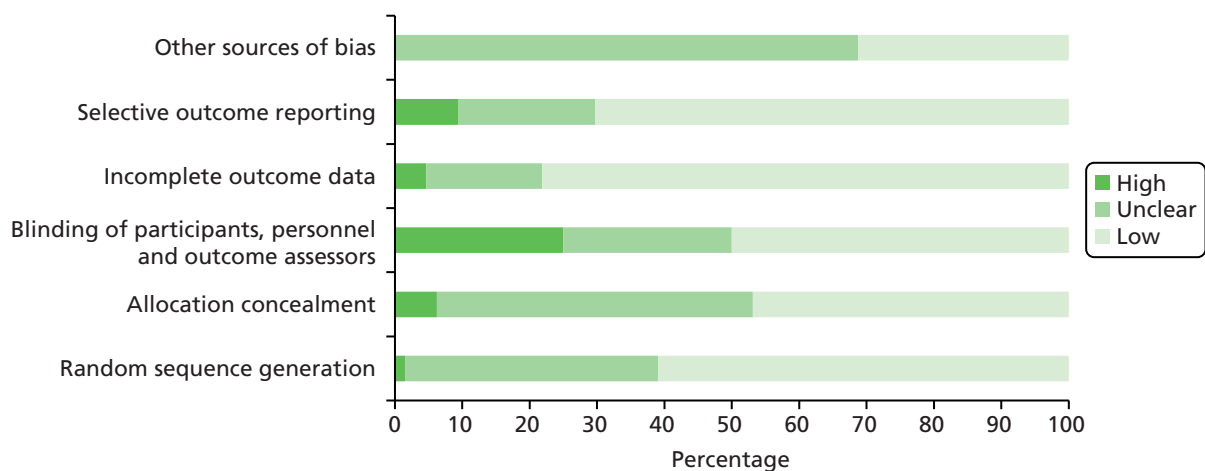


FIGURE 3 Risk of bias graph: review authors’ judgements about each risk of bias item presented as percentages across all included RCT studies.

TABLE 4 Risk of bias summary: review authors' judgements about each risk of bias item for included RCTs

Study	Random sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective outcome reporting	Other sources of bias	Within study risk of bias	Comments
Abas 2014 ⁵⁷	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Adamczak 2007 ⁵⁸	?	?	✗	?	?	?	Unclear	Abstract form only, blinding of participants and personnel not possible
Babaei 2014 ⁵⁹	?	?	✓	✓	✓	?	Unclear	Double-blind RCT, but methods of sequence generation and allocation concealment not reported
Basirat 2009 ⁶⁰	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Bayreuther 1994 ⁶¹	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Belluomini 1994 ⁶²	✓	?	✓	✗	✓	✓	Low	Most women lost to follow-up and attrition between groups was similar
Biswas 2011 ⁶³	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Bondok 2006 ⁶⁴	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Can Gurkan 2008 ⁶³	?	?	?	?	✓	?	Unclear	Study at unclear risk of bias
Capp 2014 ⁶⁵	?	?	?	✓	✓	?	Unclear	Study at unclear risk of bias
Carlsson 2000 ⁶⁶	✓	✓	✗	?	✓	?	Low	Not possible to blind personnel to acupuncture and acupuncture placebo
Chittumma 2007 ⁶⁷	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Diggory 1962 ⁶⁸	?	?	✗	✗	?	?	High	Study at high risk of bias
Ditto 1999 ⁶⁹	✓	?	?	?	?	?	Unclear	Study at unclear risk of bias
Ensiyeh 2009 ⁷⁰	✓	?	?	✓	✓	✓	Low	Study at low risk of bias
Erez 1971 ⁷¹	?	?	✓	✓	✓	✓	Low	Labelled as double-blind crossover RCT and predates 1996 CONSORT statement
Eftekhari 2013 ⁷²	?	✗	✓	✓	✓	?	Unclear	Study at unclear risk of bias

continued

TABLE 4 Risk of bias summary: review authors' judgements about each risk of bias item for included RCTs (continued)

Study	Blinding of participants, personnel and outcome assessors							Comments
	Random sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective outcome reporting	Other sources of bias	Within study risk of bias	
Evans 1993 ⁷³	?	✓	?	?	?	?	Unclear	Study at unclear risk of bias
Fischer-Rasmussen 1991 ⁷⁴	?	?	✓	✓	✓	✓	Low	Labelled as double-blind crossover RCT and predates 1996 CONSORT statement
Ghahiri 2011 ⁷⁵	?	?	✗	✓	✓	?	Unclear	Study at unclear risk of bias
Ghani 2013 ⁷⁶	✓	✓	?	✓	✓	?	Low	Study at low risk of bias overall, but did not report control results
Haji Seid Javadi 2013 ⁷⁷	?	?	✗	?	?	?	High	Vitamin B6 tablets and ginger capsules given so trial unblinded
Heazell 2006 ⁷⁸	?	✓	✗	✓	✓	?	Unclear	Study at unclear risk of bias
Hsu 2003 ⁷⁹	?	?	✓	?	?	?	Unclear	Study at unclear risk of bias
Jamigorn 2007 ⁸⁰	✓	✓	✗	✓	✓	?	Low	Not possible to blind personnel to acupuncture and acupuncture placebo
Kashifard 2013 ⁸¹	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Keating 2002 ⁸²	✓	✓	✓	✓	?	?	Low	Study at low risk of bias
Knight 2001 ⁸³	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Koren 2010 ⁸⁴	?	✓	✓	✓	✓	?	Low	Study at low risk of bias
Maina 2014 ⁸⁵	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Maltepe 2013 ⁸⁶	✓	?	✓	✓	✓	?	Low	Study at low risk of bias
Mao 2009 ⁸⁷	✓	✗	✗	✓	✗	?	High	Study at high risk of bias
McParlin 2008 ⁸⁸	✓	✓	✗	✓	✓	✓	Low	Not possible to blind patients or personnel to interventions in this trial
Mohammadbeigi 2011 ⁸⁹	✓	✓	✗	✓	✓	?	High	Personnel member gave medicines to patients and conducted randomisation
Monias 1957 ⁹⁰	?	?	✓	?	✗	?	High	Unclear whether or not this was truly a RCT and outcomes selectively reported

Study	Blinding of participants, personnel and outcome assessors							Comments
	Random sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective outcome reporting	Other sources of bias	Within study risk of bias	
Naeimi Rad 2012 ⁹¹	✓	?	✗	✓	✓	?	Low	Not possible to blind personnel, but patients were blinded to sham devices
Narenji 2012 ⁹²	?	✗	✗	✓	✓	?	High	Study at high risk of bias
Nelson-Piercy 2001 ⁹³	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Neri 2005 ⁹⁴	✓	?	?	✓	✗	?	High	Selective reporting of outcomes and just 2/6 core risk of bias items satisfied
Oliveira 2013 ⁹⁵	?	?	?	?	?	?	Unclear	Study at unclear risk of bias
Ozgoli 2009 ⁹⁶	?	?	✗	✓	✓	✓	Unclear	Unclear details on whether or not trial was truly randomised, personnel unblinded
Pasha 2012 ⁹⁷	?	?	?	✓	✓	?	Unclear	Study at unclear risk of bias
Pongrojapaw 2007 ⁴²	?	?	?	✓	?	✓	Unclear	Study at unclear risk of bias
Rosen 2003 ⁹⁸	✓	✓	?	✓	✓	?	Low	Not possible to blind personnel, but patients were blinded to sham devices
Saberi 2013 ¹³	✓	?	✗	✓	✓	✓	Low	Trial of acupuncture, ginger and no treatment so blinding not possible
Safari 1998 ⁹⁹	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Sahakian 1991 ¹⁰⁰	✓	?	✓	?	?	?	Unclear	Study at unclear risk of bias
Smith 2002 ¹⁰¹	✓	✓	✗	✓	✓	✓	Low	Not possible to blind personnel to sham control, but possible to blind patients
Smith 2004 ¹⁰²	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Sripromote 2003 ¹⁰³	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Steele 2001 ¹⁰⁴	?	?	?	✓	✓	?	Unclear	Study at unclear risk of bias

continued

TABLE 4 Risk of bias summary: review authors' judgements about each risk of bias item for included RCTs (continued)

Study	Random sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective outcome reporting	Other sources of bias	Within study risk of bias	Comments
Sullivan 1996 ¹⁰⁵	?	✓	✓	✓	?	?	Unclear	If no change in nausea or emesis was observed after 48 hours of medication and hydration, patient was considered a treatment failure, but unclear if included in final analyses
Tan 2010 ¹⁰⁶	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Tan 2009 ¹⁰⁷	✓	✓	?	✓	✓	?	Unclear	Placebo used Tic Tac [®] (Ferrero UK Ltd, Greenford, UK) which could potentially lead to unblinding
Tan 2013 ¹⁰⁸	✓	✓	✓	✓	✓	?	Low	Study at low risk of bias
Veciana 2001 ¹⁰⁹	?	?	?	?	?	?	Unclear	Study at unclear risk of bias
Vutyavanich 1995 ⁴¹	✓	?	✓	✓	✓	✓	Low	Study at low risk of bias
Vutyavanich 2001 ¹¹⁰	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Werntoft 2001 ¹¹¹	✗	?	?	✗	?	?	High	Study at high risk of bias
Wibowo 2012 ¹¹²	✓	✓	✓	✓	✓	✓	Low	Study at low risk of bias
Willetts 2003 ¹¹³	✓	?	✓	✓	✗	?	High	Selective reporting of outcomes and just 3/6 core risk of bias items satisfied
Yost 2003 ¹¹⁴	✓	?	?	✓	✗	?	High	Selective reporting of outcomes and just 2/6 core risk of bias items satisfied
Zhang 2005 ¹¹⁵	✓	✗	✗	✓	✗	?	High	Study at high risk of bias
Ziaei 2004 ¹¹⁶	?	?	✓	✓	✓	?	Unclear	Study at unclear risk of bias

✗, high risk of bias; ?, unclear risk of bias; ✓, low risk of bias; CONSORT, Consolidated Standards of Reporting Trials

Allocation concealment

Thirty studies employed allocation concealment methods judged to carry low risk of bias, such as the use of sequentially numbered sealed opaque envelopes containing allocation assignment.^{57,60,61,63,64,66,67,73,76,78,80–85,88,89,93,98,99,102,103,105–108,112,118} Thirty studies did not provide sufficient information to allow a judgement of low or high risk and were therefore classed as unclear.^{13,41–43,58,59,62,65,68–71,74,75,77,79,90,91,94–97,100,104,109,111,113,114,116,117} The remaining four RCTs were judged as having high risk of allocation concealment bias.^{72,87,92,115} For example, one study stated that patients were randomly divided into two groups by those involved in the study,⁷² or the nature of the intervention being tested meant it was not possible to conceal allocation.^{87,92,115}

Blinding of participants, personnel and outcome assessors

Of the included RCT studies, 32 were judged to have low risk of bias in relation to the blinding of participants and other personnel involved in the trial,^{41,57,59–64,67,71,72,74,79,81–85,90,93,99,100,102,103,105,106,108,110,112,113,116,117} generally through the provision of medication in identical formats for both active and placebo. Sixteen studies were judged to have high risk of bias in this respect, for example due to clear differences in either the appearance, dosage rates or mode of delivery between intervention and placebo comparator, or as a result of evidence that the research staff involved were aware of allocation status.^{13,58,66,68,75,77,78,80,87–89,91,92,96,101,115} In some instances, however, despite lack of blinding, the nature of the intervention meant that this was not relevant; for example, in McParlin and colleagues⁸⁸ where blinding of participants and staff was not possible as the packages of care delivered to the intervention and control groups varied in content. However, it is important to highlight that although it might not have been possible to blind patients or clinicians, outcome assessors and analysts handling the resultant data may nevertheless have been blinded. The remaining 16 studies did not provide sufficient information to permit a judgement of low or high bias, often due to imprecise, poor reporting, and were thus classed as unclear.^{42,43,65,69,70,73,76,94,95,97,98,104,107,109,111,114}

Incomplete outcome data

Most studies ($n = 50$) were judged as carrying low risk of bias in relation to this component.^{13,41,42,57,59–61,63–65,67,70–72,74–76,78,80–85,87–89,91–94,96–99,102–108,110,112–118} Although published protocols were rarely available, all data for the primary outcomes pre-specified in the paper were reported for all randomised participants, or rates of drop-out were either sufficiently low ($< 20\%$), or proportionately comparable between groups, so that it was not considered likely to result in a clinically relevant bias. Three studies displayed a high risk of bias in this regard, all as a result of high numbers of participant drop-outs.^{62,68,111} The remainder (11 studies in total) were judged as unclear due to lack of sufficient information.^{43,58,66,69,73,77,79,90,95,100,109}

Selective outcome reporting

Six studies were judged as having high risk of bias in terms of selective outcome reporting, due to either not reporting data for pre-specified outcomes, or for reporting data in the results that were not pre-specified in either the original study protocol or methods section.^{87,90,94,113–115} Forty-five studies were classed as having low risk of bias, with all outcomes specified and subsequently reported.^{13,41,43,57,59–67,70–72,74–76,78,80,81,83–85,88,89,91–93,96–99,101–104,106–108,110,112,116,117} Risk of bias was judged as unclear for the final 13 studies.^{42,58,68,69,73,77,79,82,95,100,105,109,111}

Other sources of bias

Twenty of the included RCT studies were judged as having low risk of bias in this area.^{13,41,42,61,62,64,67,70,71,74,81,83,88,96,101–103,106,110,112} However, a substantial number ($n = 44$) were classed as unclear, due to lack of sufficient information in the paper to permit detailed assessment of whether or not an important risk of bias existed, or due to insufficient rationale or evidence that an identified problem had introduced serious levels of bias to the study.^{43,57–60,65,66,68,69,72,73,75–80,82,84,85,87,89–95,97–100,104,105,107–109,111,113–117,119} For example, in one paper,⁷⁶ lack of reporting of full results for the control group resulted in an unclear judgement in this area.

Case series studies

The nine case series or non-randomised studies were quality assessed using the component-based EPHPP tool,⁴⁷ which appraises studies on the basis of six core components, rated 1–4 (where 1 is deemed to be

the highest quality of study). These areas are selection bias; strength of overall study design; extent to which confounders were identified and controlled for in the study; blinding of participants and/or research personnel; approach to data collection; and rate of withdrawals/drop-outs from study. As shown in the *Table 5*, all studies were judged as weak in terms of quality (which corresponds to a high risk of bias judgement using the standard Cochrane approach for RCTs).

Interventions and comparators

The included studies were grouped into the three broad groups of interventions outlined in *Chapter 1*: patient-initiated first-line interventions; clinician-prescribed second-line interventions; and clinician-prescribed third-line interventions. It should be noted that, for patient-initiated first-line interventions, the only studies identified that could be classified as lifestyle interventions were those which trialled ginger preparations and/or vitamin B6. No studies of dietary- or hypnotherapy-based interventions were identified. However, studies of a number of novel therapies not covered by our original review protocol were identified, namely the use of aromatherapy, transdermal clonidine and gabapentin. The studies comprising the evidence base for each group of interventions are detailed in *Table 6*. Note that all studies are two-arm RCTs unless otherwise stated.

In addition, the network plot (*Figure 4*) shows the range of interventions from all comparative studies included in the review. Individual interventions have been grouped where appropriate.

The size of the nodes in the network plot is proportional to the frequency of the intervention in the review, and the width of the lines indicates the frequency of the comparisons made between two interventions. These nodes and lines, however, do not represent the weight of evidence in the review as this would also be influenced by sample size and the precision of estimates, as well as other factors. The plot did not include a trial on pre-emptive treatment of doxylamine/pyridoxine combination, outpatient versus inpatient care^{117,127} or two four-arm trials,^{68,101} which would have over-reported the number of comparisons in the network plot. These interventions included dietary instructions only, or together with either placebo, antihistamines or antihistamine/vitamin B6 combination in one trial⁶⁸ and traditional acupuncture, P6 acupuncture, placebo or no acupuncture in another trial.¹⁰¹ Ginger, vitamin B6, antihistamines, acupuncture, metoclopramide, corticosteroids, doxylamine/pyridoxine combination and the

TABLE 5 Study quality summary: review authors' judgements about each risk of bias item for each included case series or non-randomised study

Study	Selection bias	Study design	Confounders	Blinding	Data collection method	Withdrawals and drop-outs	Overall rating
Alalade 2007 ¹²⁰	2	3	3	3	1	4	Weak
Ashkenazi-Hoffnung 2013 ³⁶	2	3	3	2	3	1	Weak
Einarson 2004 ¹²¹	1	3	3	3	1	1	Weak
Ferreira 2003 ¹²²	2	3	3	3	1	1	Weak
Guttuso 2010 ³⁹	1	3	3	N/A	1	1	Weak
Hsu 1996 ¹²³	2	3	3	3	1	1	Weak
Markose 2004 ¹²⁴	2	3	3	3	1	3	Weak
Moran 2002 ¹²⁵	3	3	3	3	3	1	Weak
Saha 2009 ¹²⁶	3	3	3	2	2	4	Weak

N/A, not applicable.

TABLE 6 Number of studies by intervention and comparator

Intervention/comparator	Number of studies	Studies
Patient-initiated first-line intervention		
Ginger vs. placebo	7 ^a	Basirat 2009 ⁶⁰ (biscuit) Fischer-Rasmussen 1991 ⁷⁴ Keating 2002 ⁸² (syrup) ^a Mohammadbeigi 2011 ⁸⁹ (also ginger vs. metoclopramide) Ozgili 2009 ⁹⁶ Vutyavanich 2001 ¹¹⁰ Willets 2003 ¹¹³
Ginger vs. acupressure	1	Saberi 2013 ¹³
Ginger vs. vitamin B6	6	Chittumma 2007 ⁶⁷ Ensiyeh 2009 ⁷⁰ Narenji 2012 ⁹² Haji Seid Javadi 2013 ⁷⁷ Smith 2004 ¹⁰² Sripramote 2003 ¹⁰³
Ginger vs. doxylamine/pyridoxine	1	Biswas 2011 ⁶³
Ginger vs. antihistamine	1	Pongrojapaw 2007 ⁴²
Ginger vs. metoclopramide	1 ^a	^a Mohammadbeigi 2011 ⁸⁹ (also ginger vs. placebo)
Vitamin B6 vs. placebo	3	Tan 2009 ¹⁰⁷ [metoclopramide ± vitamin B6 (dopamine receptor antagonist)] Sahakian 1991 ¹⁰⁰ Vutyavanich 1995 ⁴¹
High- vs. low-dose vitamin B6	1	Wibowo 2012 ¹¹² (high- vs. low-dose vitamin B6)
Antihistamine ± vitamin B6	2	Babaei 2014 ⁵⁹ Diggory 1962 ⁶⁸ (four-arm RCT)
Aromatherapy	2	Ghani 2013 ⁷⁶ Pasha 2012 ⁹⁷
Acupressure vs. nocebo	8	Bayreuther 1994 ⁶¹ Belluomini 1994 ⁶² Can Gurkan 2008 ⁴³ Heazell 2006 ⁷⁸ Hsu 2003 ⁷⁹

continued

TABLE 6 Number of studies by intervention and comparator (continued)

Intervention/comparator	Number of studies	Studies
		Naeimi Rad 2012 ⁹¹
		Steele 2001 ¹⁰⁴
Acupressure	1	Werntoff 2001 ¹¹¹ (three-arm RCT)
Acupressure vs. vitamin B6	1	Markose 2004 ¹²⁴ (case series)
Nerve stimulation vs. placebo	3	Jamigorn 2007 ⁸⁰
		Evans 1993 ⁷³
		Rosen 2003 ⁹⁸
Acupuncture vs. placebo	3	Veciana 2001 ¹⁰⁹
		Carlsson 2000 ⁶⁶
		Knight 2001 ⁸³
Acupuncture vs. metoclopramide	1	Smith 2002 ¹⁰¹ (four-arm RCT)
Acupuncture ± moxibustion vs. Chinese herbal medicine vs. Western medicine	2	Neri 2005 ⁹⁴
		Mao 2009 ⁸⁷
		Zhang 2005 ¹¹⁵
Clinician-prescribed second-line intervention		
Doxylamine/pyridoxine (Diclectin) vs. placebo	1	Koren 2010 ⁸⁴
Pre-emptive Diclectin vs. Diclectin once symptoms begin	1	Maltepe 2013 ¹¹⁷ (pre-emptive therapy)
		Koren 2013 ¹²⁷
Doxylamine/pyridoxine vs. metoclopramide	1	Ashkenazi-Hoffnung 2013 ³⁶ (cohort study)
Doxylamine/pyridoxine vs. ondansetron	2	Oliveira 2013 ⁹⁵
		Capp 2014 ⁶⁵
i.v. fluids (D-Saline vs. N-Saline)	1	Tan 2013 ¹⁰⁸
i.v. fluids ± diazepam	1	Ditto 1999 ⁶⁹
Antihistamine vs. placebo	1	Erez 1971 ⁷¹
Antihistamine + vitamin B6 vs. placebo	1	Monias 1957 ⁹⁰
Droperidol/antihistamine combination vs. other medication (dopamine receptor antagonist)	1	Ferreira 2003 ¹²² (cohort study)
Metoclopramide vs. antihistamine (phenothiazine) (dopamine receptor antagonist)	1	Tan 2010 ¹⁰⁶
Serotonin antagonist (ondansetron) vs. antihistamines	2	Eftekhari 2013 ⁷²
		Sullivan 1996 ¹⁰⁵
Serotonin antagonist (ondansetron) vs. metoclopramide	3	Abas 2014 ⁵⁷
		Ghahiri 2011 ⁷⁵
		Kashifard 2013 ⁸¹
Serotonin antagonist (ondansetron) vs. other	1	Einarson 2004 ¹²¹ (cohort study)
Transdermal clonidine vs. placebo patch	1	Maina 2014 ⁸⁵
Out patient management vs. routine inpatient care	1	McParlin 2008 ⁸⁸ and McParlin unpublished
Out patient management	1	Alalade 2007 ¹²⁰ (case series)

TABLE 6 Number of studies by intervention and comparator (continued)

Intervention/comparator	Number of studies	Studies
Clinician-prescribed third-line intervention		
Steroids vs. 'treatment as usual'	7	Adamczak 2007 ⁵⁸ Bondok 2006 ⁶⁴ Moran 2002 ¹²⁵ (case series) Nelson-Piercy 2001 ⁹³ Safari 1998 ¹²⁸ Yost 2003 ¹¹⁴ Ziaei 2004 ¹¹⁶
Nasogastric feeding	1	Hsu 1996 ¹²³ (case series)
Jejunostomy	1	Saha 2009 ¹²⁶ (case series)
Gabapentin	1	Guttuso 2010 ³⁹ (case series)

D-Saline, dextrose saline; N-Saline, normal saline.

a Results from a three-arm RCT, ginger vs. placebo vs. metoclopramide; therefore, this study appears twice in the comparator table.

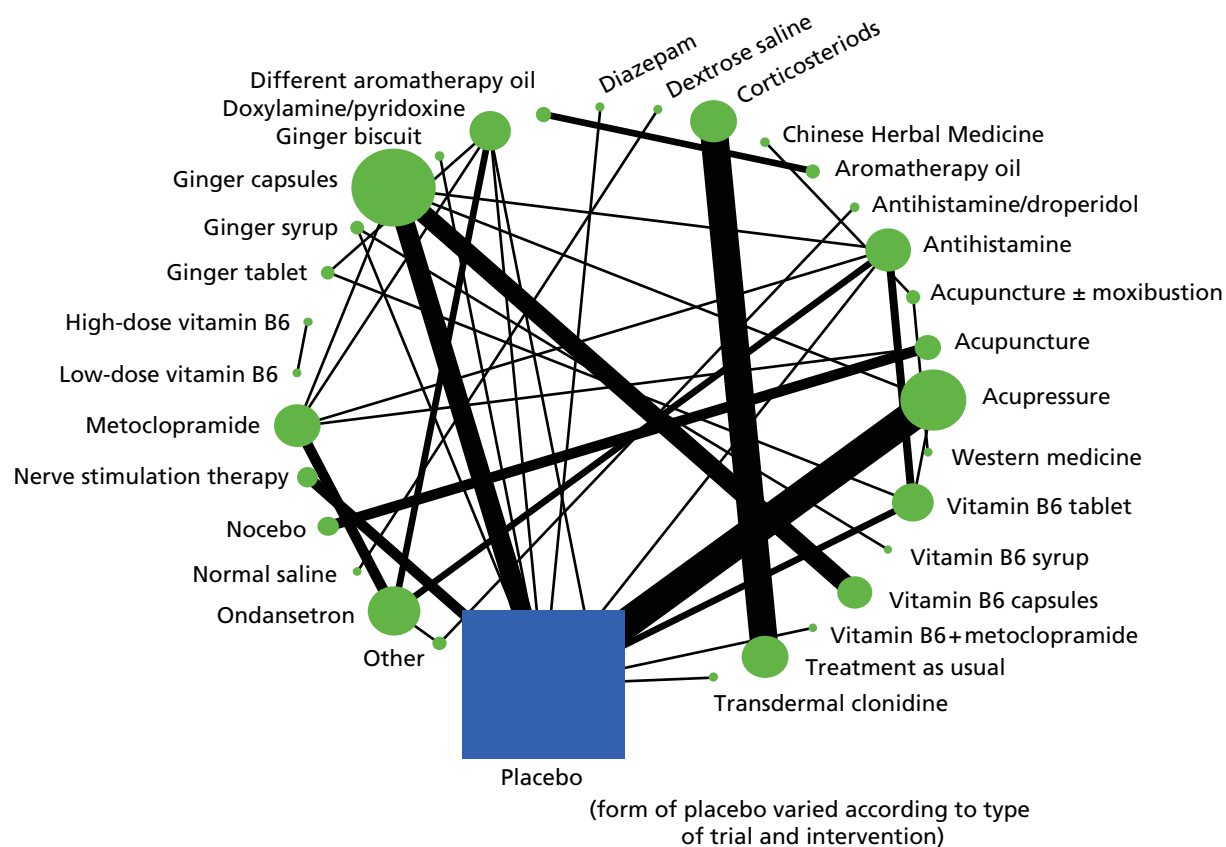


FIGURE 4 Network plot of range of interventions and comparisons for NVP/HG. Size of node is proportional to frequency of intervention and width of line to frequency of comparisons between two interventions. Plot does not include one pre-emptive trial,¹¹⁷ outpatient care trial¹²⁷ and two four-arm trials.^{68,101}

serotonin antagonist ondansetron are more widely reported than other interventions, but there is also information on interventions such as acupuncture, nerve stimulation therapy and aromatherapy oils which have been considered as treatments for NVP/HG. Evidence on the effects of interventions such as Chinese herbal medicine, dextrose saline, transdermal clonidine and diazepam is very limited and in most cases is reported in single trials. As expected, placebo interventions are most widely reported as comparators, and so this has the biggest node on the network plot (emphasised by the square node). The most commonly reported treatment comparisons are ginger capsules versus placebo; acupressure versus placebo; ginger capsules versus vitamin B6 capsules; corticosteroids versus 'treatment as usual'; metoclopramide versus ondansetron; and acupuncture versus nocebo (nocebo is an inert intervention that creates comparable side effects/harmful effects in a patient, as opposed a placebo, which is an inert substance that creates either a beneficial response or no response in a patient).

Participants and symptom severity

In addition to substantial variation in terms of the range of interventions and comparators evident within the literature, it is also important to highlight the heterogeneity of symptom severity found among patient populations.

It was initially intended that as part of this review only studies that recruited women with severe NVP or HG would be included. However, assessment of symptom severity varied within and across studies, and it was not possible to easily place every participant population into categories. We therefore attempted to categorise the symptom severity of participants for each study, using the description of severity in the inclusion criteria and, if available, any severity score given at baseline. These two items of information were assessed by two independent assessors (CMP and SCR) to assign severity as mild, moderate, severe or unclear. Agreement was reached for all but one study, which was classified as unclear.

This classification was then used in each results chapter to describe symptoms and outcomes in terms of severity.

Outcome measures

Finally, and linked to the issues discussed above, the identified literature in this field was also characterised by the range of symptom severity scales employed from study to study to assess intervention outcomes. Out of the 73 included studies (reported in 75 papers), only 23 used validated NVP/HG assessment scales such as PUQE (10 studies), RINVR (11 studies) or the McGill Nausea Questionnaire (one study). Thirty-one studies assessed nausea and/or vomiting severity using a 10-point VAS. Twenty-one studies employed either a study-specific, non-validated author-defined assessment scale (including, for example, numbers of episodes of vomiting combined with the use of a Likert scale to assess subjective feelings of symptom severity among participants), or used the various proxy measures of symptom severity outlined in our protocol [e.g. percentage weight loss, length of hospital stay, or hospital (re-)admission episodes]. *Table 7* illustrates the primary symptom severity outcome measures employed by each included study.

Additional sources of outcome data on medications

The UKTIS is currently commissioned by Public Health England to provide advice to UK health professionals on the fetal effects of therapeutic, poisoning and chemical exposures in pregnancy, and to conduct surveillance of known and emerging teratogens. The UKTIS database currently contains a record of just under 60,000 enquiries dating back to 1978, of which 320 relate to use of specific drugs in the treatment of HG (period of enquiry 18 June 1978 to 18 March 2014). Surveillance data collected by the UKTIS are reviewed periodically and published in UKTIS monographs through the National Poisons Information

TABLE 7 Validated and non-validated symptom severity measures employed by each included study

Study	PUQE	RINVR	McGill Nausea Questionnaire	NVPI	VAS	Other scale/proxy measure
Abas 2014 ⁵⁷					✓	✓
Adamczak 2007 ⁵⁸						✓
Alalade 2007 ¹²⁰						✓
Ashkenazi-Hoffnung 2013 ³⁶						✓
Babaei 2014 ⁵⁹		✓				
Basirat 2009 ⁶⁰					✓	
Bayreuther 1994 ⁶¹					✓	
Belluomini 1994 ⁶²		✓				
Biswas 2011 ⁶³					✓	
Bondok 2006 ⁶⁴						✓
Can Gurkan 2008 ⁴³					✓	
Capp 2014 ⁶⁵					✓	✓
Carlsson 2000 ⁶⁶					✓	
Tan 2010 ¹⁰⁶					✓	
Tan 2013 ¹⁰⁸					✓	
Tan 2009 ¹⁰⁷					✓	
Chittumma 2007 ⁶⁷		✓				
Diggory 1962 ⁶⁸						✓
Ditto 1999 ⁶⁹					✓	
Einarson 2004 ¹²¹	✓					
Ensiyeh 2009 ⁷⁰					✓	
Erez 1971 ⁷¹						✓
Evans 1993 ⁷³						✓
Ferreira 2003 ¹²²						✓
Fischer-Rasmussen 1991 ⁷⁴					✓	
Ghahiri 2011 ⁷⁵					✓	
Ghani 2013 ⁷⁶		✓				
Guttuso 2010 ³⁹	✓					
Heazell 2006 ⁷⁸						✓
Hsu 1996 ¹²³						✓
Hsu 2003 ⁷⁹			✓			
Jamigorn 2007 ⁸⁰		✓				
Kashifard 2013 ⁸¹					✓	
Keating 2002 ⁸²					✓	
Knight 2001 ⁸³					✓	
Koren 2010 ⁸⁴	✓					
Maina 2014 ⁸⁵	✓				✓	
Maltepe 2013 ⁸⁶	✓					
Mao 2007 ⁸⁷						✓
Markose 2004 ¹²⁴		✓				

continued

TABLE 7 Validated and non-validated symptom severity measures employed by each included study (*continued*)

Study	PUQE	RINVR	McGill Nausea Questionnaire	NVPI	VAS	Other scale/proxy measure
McParlin 2008 ⁸⁸	✓					
Mohammadbeigi 2011 ⁸⁹		✓				
Monias 1957 ⁹⁰						✓
Moran 2002 ¹²⁵					✓	
Naeimi Rad 2012 ⁹¹					✓	
Narenji 2012 ⁹²					✓	
Nelson-Piercy 2001 ⁹³					✓	
Neri 2005 ⁹⁴						✓
Oliveira 2013 ⁹⁵					✓	
Ozgoli 2009 ⁹⁶					✓	
Pasha 2012 ⁹⁷					✓	
Pongrojpraw 2007 ⁴²					✓	
Rosen 2003 ⁹⁸		✓				
Saberi 2013 ¹³		✓				
Safari 1998 ⁹⁹						✓
Saha 2009 ¹²⁶						✓
Sahakian 1991 ¹⁰⁰					✓	
Haji Seid Javadi 2013 ⁷⁷	✓					
Smith 2004 ¹⁰²		✓				
Smith 2002 ¹⁰¹						✓
Sripramote 2003 ¹⁰³					✓	
Steele 2001 ¹⁰⁴		✓				
Sullivan 1996 ¹⁰⁵					✓	
Eftekhari 2014 ⁷²						✓
Veciana 2001 ¹⁰⁹		✓				
Vutyavanich 2001 ¹¹⁰					✓	
Vutyavanich 1995 ⁴¹					✓	
Werntoft 2001 ¹¹¹					✓	
Wibowo 2012 ¹¹²	✓					
Willets 2003 ¹¹³		✓				
Yost 2003 ¹¹⁴						✓
Zhang 2005 ¹¹⁵						✓
Ziaei 2004 ¹¹⁶					✓	

Service database (www.TOXBASE.org). Data collected by the UKTIS in relation to medications for NVP/HG, including specific monograph data on ginger, vitamin B6, vitamin B12, promethazine and olanzapine, are provided in *Table 43, Appendix 7* for information.

Meta-analysis of included randomised controlled trials

As highlighted in the previous sections, there was wide variation across studies. Specifically, there was considerable heterogeneity between interventions within each of the categories of comparisons, and in terms of how interventions were administered/delivered. The measurement of outcomes also differed substantially between trials reporting the same comparisons, so in most cases the trials were not directly comparable. In a meta-analysis it is important not to combine outcomes that are too diverse; even if it had been possible to extract data for a meta-analysis, such an analysis is likely to produce misleading results due to the considerable heterogeneity between studies.⁴⁶ Furthermore, many of these trials were extremely poorly reported and their conduct was often uncertain. In summary, clinical and methodological variations between studies were considerable, and the intervention effect was likely to be affected by the factors that varied across studies. Consequently, we have not conducted a meta-analysis of findings from the RCTs.

Structure of individual results chapters

The following chapters present more detailed findings from the evidence review for each individual intervention. As already indicated, given it was not possible to meta-analyse the data from individual studies for any group of interventions and comparators, the results are summarised in narrative form. The narrative content of each chapter focuses on the findings from the included studies in terms of their reported effectiveness for addressing our primary outcomes of interest, that is, the key symptoms associated with HG/NVP. Thus, where available, effectiveness is reported in terms of the validated overall HG/NVP assessment scales (PUQE, RINVR or McGill Nausea Questionnaire). Otherwise, the effectiveness of interventions is reported in relation to their impact on the three key symptoms: nausea, vomiting and retching. Data illustrating significant results in relation to these key symptoms are detailed in the narrative text; otherwise, results are described as not significant or not clear. Data for case series studies are not included in the narrative but available in the accompanying results tables for information. Additional secondary outcome data reported by included studies (see *Table 2* for a full list) are presented in *Appendix 8*.

Chapter 4 Clinical effectiveness: ginger

Introduction

Ginger was used as an intervention to treat HG, NVP or various forms of pregnancy sickness in a total of 16 RCTs.^{13,42,60,63,67,70,74,77,82,89,92,96,102,103,110,113} Heterogeneity was observed in relation to the clinical setting and patient populations in which the studies were conducted, as well as the interventions, comparators and outcomes reported in each trial. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*) given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses and have thus reported a narrative summary only for each intervention and comparator set, with additional outcome data presented in *Table 8*.

Of the included trials, half ($n = 8$) described the women participants as experiencing symptoms categorised at the mild end of the NVP severity scale.^{60,63,67,70,82,89,110,113} Six were classed as including participants with mild–moderate symptom severity.^{13,74,92,96,102,103} The information provided by the remainder was insufficient to categorise objectively severity, although the authors describe the women as experiencing pregnancy-related nausea and vomiting.^{42,77} Of the 16 studies, 4 were judged to carry a high risk of bias,^{77,89,92,113} 10 low risk,^{13,60,63,67,70,74,82,102,103,110} and the remainder, unclear due to lack of information.^{42,96}

Ginger capsules versus placebo capsules

Five trials^{74,89,96,110,113} compared ginger and placebo capsules for the treatment of NVP. The trial by Mohammadbeigi and colleagues⁸⁹ was a three-arm RCT which compared ginger, metoclopramide and placebo.⁸⁹ Only one trial⁷⁴ explicitly reported the effects of ginger in the treatment of HG (which they define as vomiting occurring during pregnancy, presenting prior to the 20th week and of sufficient severity to require hospital admission). However, it is important to note that this deviates from the standard definition of HG (see *Chapter 1, Background*). Three of these trials were judged to carry a low risk of bias, two high risk due to issues regarding blinding of personnel administering the treatment⁸⁹ or selective outcome reporting¹¹³ and the risk of bias in the remaining trial was unclear.⁹⁶

Rhodes Index of Nausea, Vomiting and Retching

In the Mohammadbeigi and colleagues⁸⁹ trial, the observed differences in the RINVR score on the second day (ginger: 26.41 ± 4.12 ; metoclopramide: 25.56 ± 5.51 ; placebo: 27.35 ± 3.36) and the fifth day (ginger: 18.71 ± 2.81 ; metoclopramide: 18.53 ± 5.18 ; placebo: 23.15 ± 4.03) compared with the first day of treatment (ginger: 31.68 ± 5.32 ; metoclopramide: 30.00 ± 8.29 ; placebo: 30.53 ± 4.64) were statistically significant in all groups ($p < 0.001$).⁸⁹ Although there was a reduction in scores in all three groups, the authors stated that the intensity of changes was different in ginger ($p < 0.01$) and metoclopramide ($p = 0.03$) groups compared with placebo, but did not differ between the ginger and metoclopramide groups ($p = 0.51$). However, it was not clear from the paper how differences in the intensity of changes were established.

Author-defined symptom severity/relief scales

Fischer-Rasmussen and colleagues⁷⁴ devised a study-specific severity/relief score based on the degree of nausea, duration and number of episodes of vomiting and change in body weight, and a subjective assessment of the extent to which these symptoms had improved, worsened or stayed the same. In this randomised crossover trial, the authors compared relief scores at day 5 and day 11, following 4 days of treatment with either ginger or placebo capsules. Fischer-Rasmussen and colleagues⁷⁴ reported that the mean severity scores decreased equally in the two groups. However, given the differences in gestational

TABLE 8 Results for ginger-based interventions for NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ginger vs. placebo									
Basirat 2009 ⁶⁰	Antenatal clinic, Babol University of Medical Sciences, Iran	Effect of ginger in a biscuit form on NVP, double-blind RCT	65 (I = 35, C = 35), 7–17 weeks	Author defined, women with nausea and vomiting of pregnancy	Nausea VAS score: I = 5.88 ± 1.83, C = 4.67 ± 1.97 Episodes of vomiting I = 1.63 ± 1.18, C = 1.3 ± 1.3 (MILD)	Ginger biscuits (0.5 mg of ginger, five biscuits daily for 4 days)	Non-ginger containing biscuits (five biscuits daily for 4 days)	VAS for nausea Episodes of vomiting	Nausea: average change in scores over 4 days in the ginger group = 2.57 ± 1.77, placebo group = 1.39 ± 1.62; <i>p</i> = 0.01 Vomiting: average change in episodes over 4 days in ginger group = 0.96 ± 0.2, placebo group = 0.62 ± 0.19; <i>p</i> = 0.243
Fischer-Rasmussen 1991 ⁷⁴	Obstetrics and Gynaecology Department, University Hospital of Copenhagen, Denmark	Effectiveness of ginger capsules on the symptoms of HG, double-blind randomised crossover trial	30 (I = 15, C = 15), I = 11 weeks (7–17), C = 10.8 weeks (7–16)	Women admitted to the hospital with HG before 20th week of gestation and with symptoms persisting after 2 days	70% of participants reported constant nausea at baseline 66.6% of women reported vomiting two to six times per day 33.3% of women reported vomiting seven or more times per day (MILD–MODERATE)	Ginger capsules (250 mg of powdered root ginger) four times daily for 4 days, followed by a 2-day wash out before alternate treatment	Placebo capsules (250 mg of lactose) four times daily for 4 days, followed by a 2-day wash out before alternate treatment	Author-defined severity score for nausea (1–3) Author-defined relief score for nausea and vomiting (–3 to 3) Episodes of vomiting	Mean severity scores decreased equally in the two groups Mean relief scores highlighted greater relief of symptoms after ginger treatment compared to placebo (<i>p</i> = 0.035)
Keating 2002 ⁸²	Department of Obstetrics and Gynaecology, University of South Florida, FL, USA, private practice office	Effectiveness of ginger syrup mixed in water in treating NVP in the first trimester of pregnancy, double-blind RCT	26 (I = 14, C = 12), 7–11 weeks	Women complaining of nausea with and without vomiting either at a planned or ad hoc obstetric visit and were not taking a prescribed or over-the-counter antiemetic	NR (MILD)	250 mg ginger + honey and water. One table spoon mixed with 4–8 ounces water taken four times a day for 2 weeks. Both groups received verbal and written dietary advice	Placebo syrup = water, honey and lemon oil. One table spoon mixed with 4–8 ounces water taken four times a day for 2 weeks	VAS for nausea Episodes of vomiting	Ten women in ginger group had ≥ 4-point improvement on VAS by day 9. Placebo group: two women had same improvement by day 9 Eight women in the ginger group stopped vomiting by day 6. Only two women in the placebo group stopped by day 6 No statistical analysis

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ozgoli 2009 ⁹⁶	Isfahan City Hospitals, Iran	Effects of ginger in nausea and vomiting of pregnancy, single-blind RCT	70 (I = 35, C = 35)	Women with mild and moderate nausea, with or without vomiting	Nausea intensity moderate I = 56% C = 54%; mild I = 18.7% C = 25.7%; severe I = 8% C = 7% (MILD-MODERATE)	Capsules containing 250 mg of ginger extract taken four times daily for 4 days. Dietary advice given. Twice daily VAS score completed	Identical placebo capsules containing lactose taken following same regime. Dietary advice given. Twice daily VAS score completed	VAS for vomiting symptoms VAS for nausea intensity	Nausea intensity: ginger group had a higher rate of improvement (85% vs. 56%; $p < 0.01$) Vomiting: 50% decrease in the ginger group vs. 9% in the placebo group; $p < 0.05$
Vutyavanich 2001 ¹¹⁰	Antenatal clinic, Maharaj Nakorn Chiang Mai University Hospital, Chiang Mai, Thailand	Effectiveness of ginger for the treatment of NVP, double-masked RCT	70 (I = 32, C = 38), I = 10.3 ± 2.6	First attended the clinic before 17 weeks' gestation and had nausea of pregnancy, with or without vomiting	Baseline nausea scores (cm): I = 4.7 ± 2.1, C = 5.4 ± 2.1 (MILD)	Capsules containing 250 mg of ginger taken three times/day following meals and another before bed for 4 days. Dietary advice given	Identical placebo capsules following the same regime. Dietary advice given	VAS for nausea Episodes of vomiting	Decrease in nausea score: ginger group (2.1 ± 1.9) vs. placebo group (0.9 ± 2.2; $p = 0.014$) Decrease in vomiting episodes: ginger group (1.4 ± 1.3) vs. placebo group (0.3 ± 1.1; $p < 0.001$)
Willetts 2003 ¹¹³	Antenatal clinic, Royal Hospital for Women, Sydney, NSW, Australia	Effect of a ginger extract (EV.EXT35) on the symptoms of morning sickness, double-blind RCT	120 (I = 60, C = 60)	Women who have experienced morning sickness daily for at least a week which had failed to respond to dietary measures	Nausea experience score presented as a figure, no numerical values available (MILD)	Capsules containing 125 mg of ginger extract taken four times daily for 4 days. RINVR score completed 1 hour later	Identical placebo capsules containing soya oil taken following same regime	RINVR score	Nausea experience score: significantly less for the ginger extract group relative to the placebo group after the first day and each subsequent treatment day Retching: reduced by the ginger extract although to a lesser extent Vomiting: no significant effect observed

continued

TABLE 8 Results for ginger-based interventions for NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ginger vs. vitamin B6									
Chittumma 2007 ⁶⁹	Antenatal clinic, Vajira Hospital, Bangkok, Thailand	Effectiveness of ginger in comparison with vitamin B6 for treatment of NVP, double-blind RCT	126 (I = 63, C = 63), I = 12 (SD 2) C = 11 (SD 2)	Women who had nausea with or without vomiting and required treatment	RINVR score: I = 8.7 (SD 2.2), C = 8.3 (SD 2.5) (MILD)	Ginger capsules, 2 x 325 mg four times daily for 4 days. Dietary advice given	Vitamin B6 capsules, 2 x 12.5 mg four times daily for 4 days. Dietary advice given	RINVR score	Symptoms reduced in both groups. Ginger from 8.7 ± 2.2 to 5.4 ± 2.0 and vitamin B6 from 8.3 ± 2.5 to 5.7 ± 2.3 (p < 0.05). The mean score change with ginger was greater than with vitamin B6 (3.3 ± 1.5 vs. 2.6 ± 1.3) (p < 0.05)
Ensiyeh 2009 ⁷⁰	Antenatal clinic, Fatemeh Hospital, Hamedan, Iran	Effectiveness of ginger in comparison with vitamin B6 for the treatment of NVP, double-blind RCT	70 (I = 35, C = 35), NR	Women's first attendance at clinic and experienced nausea, with or without vomiting	VAS for nausea: I = 5.4 cm ± 2.6, C = 4.6 cm ± 2.3 (MILD)	Ginger capsules, 500 mg, twice daily for 4 days. VAS completed three times daily	Vitamin B6 capsules, 20 mg twice daily for 4 days. VAS completed three times daily	VAS to assess nausea Episodes of vomiting	Mean change in nausea score over 4 days: ginger 2.2 ± 1.9 compared with vitamin B6 group 0.9 ± 1.7 (p = 0.024) Vomiting episodes decreased in both groups, but no difference between groups. Ginger mean change 0.6 ± 0.7, vitamin B6 mean change 0.5 ± 1.1 (p = 1.101)
Haji Seid Javadi 2013 ⁷⁷	Health centres, University of Medical Sciences of Qazvin, Iran	Effectiveness of ginger in comparison with vitamin B6 for the treatment of NVP, RCT	95 (I = 47, C = 48), ginger 9 (SD 1.1) weeks, vitamin B6 9 (SD 1.6) weeks	Women suffering from pregnancy induced nausea	NR (NOT CLEAR)	Ginger tablets, 250 mg, 6-hourly for 4 days	Vitamin B6 tablets, 40 mg, 12-hourly for 4 days	PUQE score	Mean change of PUQE score over 4 days: ginger group 8.32 (SD 2.19), vitamin B6 group 7.77 (SD 1.80); p = 0.172 Retching in vitamin B6 group had a greater, non-significant reduction; p = 0.333
Narenji 2012 ⁹²	Antenatal clinic, Arak University of Medical Sciences, Iran	Effectiveness of ginger syrup in comparison with vitamin B6 for the treatment of NVP, RCT	100 (I = 50, C = 50) states < 17 weeks	Women attending clinic suffering from at least 24 hours of nausea	VAS mean (SE): nausea I = 5.6 (1.94), C = 6.04 (2.55); vomiting I = 5.6 (1.94), C = 6.66 (2.41) (MILD-MODERATE)	Given i.v. fluids to rehydrate. Ginger syrup (mix of ginger and honey), one teaspoon twice daily for 4 days	Given i.v. fluids to rehydrate, vitamin B6 capsules, 40 mg twice daily for 4 days	VAS to assess nausea Episodes of vomiting	Nausea score: vitamin B6 mean change 0.7 (SD 1.99), ginger mean change 1.0 (SD 1.32); p = 0.8 Vomiting decreased in both groups but no significant difference between groups

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Smith 2004 ¹⁰²	University, Women's and Childrens Hospital, Adelaide, SA, Australia, referrals from GPs and health-care providers	Comparison of ginger with pyridoxine hydrochloride (vitamin B6) in the treatment of NVP, randomised, controlled equivalence trial	301 (I = 146, C = 145), I median = 8.5, range 8–15; C median = 8.6, range 8–15	Women with nausea or vomiting	Use of antiemetics: I = 33 (44%), C = 42 (56%) (MILD–MODERATE)	Ginger capsules, (350 mg) three times a day for 3 weeks. (Participants could also take any other antiemetics they wanted)	Vitamin B6 capsules (25 mg) three times a day for 3 weeks. (Participants could also take any other anti-emetics they wanted)	RINVR score	Nausea: equivalent reduction in score for both treatments. Mean difference 0.2 (90% CI -0.3 to 0.8) Retching: mean difference 0.3 (90% CI 0.0 to 0.6) Vomiting: mean difference 0.5 (90% CI 0.0 to 0.9), averaged over time, with no evidence of different effects at the three time points (7, 14 and 21 days)
Siripramote 2003 ¹⁰³	Antenatal clinic, Vajira Hospital, Thailand, and Obstetrics and Gynaecology Department, Bangkok Metropolitan Administration Medical College, Bangkok	Efficacy of ginger in comparison with vitamin B6 in the treatment of NVP, double-blind RCT	138 (I = 68, C = 70), I = 10.1 (SD 2.74), C = 10.3 (SD 2.95)	Women with nausea of pregnancy, with or without vomiting and requesting antiemetics	Nausea scores: I = 5.0 cm (SD 1.99), C = 5.3 cm (SD 2.08). Episodes of vomiting in last 24 hours: I = 1 (range 0–10), C = 1 (range 0–10) (MILD–MODERATE)	Ginger capsules, 500 mg, three times daily for 3 days. Dietary advice given. VAS for nausea completed three times daily, plus record of vomiting episodes	Vitamin B6 capsules, 10 mg, following the same regime	VAS for nausea Episodes of vomiting	Nausea: decreased in both groups Ginger: 5.0 (SD 1.99) to 3.6 (SD 2.48) Vitamin B6: 5.3 (SD 2.08) to 3.3 (SD 2.07); <i>p</i> < 0.001 Vomiting: reduced in both groups Ginger: 1.9 (SD 2.06) to 1.2 (SD 1.75) Vitamin B6: 1.7 (SD 1.81) to 1.2 (SD 1.50); <i>p</i> < 0.01

continued

TABLE 8 Results for ginger-based interventions for NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ginger vs. acupressure									
Saberi 2013 ¹³	Antenatal clinic, Naghvi Hospital, Kashan, Iran	Effectiveness of acupressure in comparison with ginger in the treatment of NVP, three-arm RCT	159 (I = 53, C = 53, control = 53), ginger = 8.78 (SD 2.32), acupressure = 9.32 (SD 2.38), control = 9.11 (SD 0.18)	Women having mild to moderate nausea and/or vomiting	RINVR score: ginger = 17.91 (SD 6.11); acupressure = 17.91 (SD 5.90), C = 17.90 (SD 5.30) (MILD-MODERATE)	Ginger capsules, 250 mg, three times daily for 4 days. n = 53 acupressure: sea-bands given, to be worn continuously for 4 days	C group received no I	RINVR score	Mean difference in improvement pre and post I: ginger 8.61 (SD 5.24), acupressure 4.17 (SD 5.53), C -0.84 (SD 3.72) p < 0.001 for differences in total score and for each component (nausea, vomiting and retching)
Ginger vs. vitamin B6 combination									
Biswas 2011 ⁶³	Seth Sukhlal Karnani Memoria Hospital, Kolkata, India and R. G. Kar Medical College Hospitals, Kolkata, India	Efficacy and tolerability of ginger extract in comparison with a combination of doxylamine 10 mg + pyridoxine 10 mg, single-blind RCT	78 (I = 42, C = 36), I = 10.25 (SD 2.8), C = 9.3 (SD 3.1)	Women who sought treatment for the symptoms of morning sickness between 6 and 16 weeks of pregnancy without having received any previous treatment	Median (IQR) VAS score nausea: I = 34.5 mm (IQR 25.0), C = 30.4 mm (IQR 34.0). Vomiting: I = 14.5 mm (IQR 33.0), C = 22.0 mm (IQR 35.0) (MILD)	Ginger tablets, 150 mg, three daily	Doximate (Sigma Laboratories, PVT LTD, Mumbai) (doxylamine 10 mg plus pyridoxine 10 mg) three times daily	VAS for nausea Episodes of vomiting	Both groups had a significant decline in severity of nausea and vomiting from baseline to week 2 Nausea persisted, but with reduced severity median values tending towards zero at study end No significant difference between groups
Ginger vs. antihistamine									
Pongrojapaw 2007 ⁴²	Hospital antenatal clinic, Thammasat University Hospital, Thailand	To study the efficacy of ginger and dimenhydrinate in the treatment of NVP, double-blind RCT	170 (I = 85, C = 85), I = 10.25 (SD 2.8), C = 9.3 (SD 3.1)	Pregnant women with nausea and vomiting less than 16 weeks of gestation	NR (NOT CLEAR)	Ginger capsules, 500 mg, twice daily for 1 week	Dimenhydrinate capsules, 50 mg twice daily for 1 week	VAS to assess nausea Episodes of vomiting	Nausea: mean score on days 1-7 decreased in both groups. Daily score between groups not statistically different (p > 0.05) Vomiting episodes: on days 1-7 decreased in both groups. Dimenhydrinate significantly more effective on days 1-2 (p < 0.05). Days 3-7 scores similar

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ginger vs. metoclopramide									
Mohammadbeigi 2011 ^{6b}	B'esar Hospital, Kurdistan, Iran	To study the effects of ginger NVP with metoclopramide, three-arm RCT	102 (I = 34, C = 34, placebo = 34), ginger 9.5 ± 2.02, metoclopramide 10.03 ± 1.99, placebo 10.32 ± 2.25	Women with inefficacy of food regimens in controlling vomiting and nausea	NR (MILD)	Ginger capsules, 200 mg, three times daily for 5 days	Metoclopramide capsules, 10 mg, three times daily, and placebo (200 mg flour) for 5 days	RINVR score	Trend in symptom improvement to the 5th day in all three groups Nausea: ginger ($p = 0.003$) and metoclopramide ($p = 0.001$) had significantly better improvement than placebo, but no difference between ginger and metoclopramide ($p = 0.683$) Vomiting: ginger ($p = 0.046$) and metoclopramide ($p = 0.018$) had significantly better improvement than placebo. No difference between ginger and metoclopramide group ($p = 0.718$)

C, control; I, intervention; IQR, interquartile range; NR, not reported; SD, standard deviation.

age at baseline, the scores at points 1 and 11 were not considered directly comparable, and therefore no statistical analysis was carried out. Mean relief scores showed greater relief of symptoms after ginger treatment than after placebo ($p = 0.035$), with analyses of single-item components demonstrating a particular reduction in vomiting episodes and nausea in the ginger group.

Vutyavanich and colleagues¹¹⁰ used 5-item Likert scales to assess patients' subjective response to treatment in terms of whether general nausea and vomiting symptoms had worsened, improved or stayed the same. Of the 67 women surveyed, 28 out of 32 (88%) treated with ginger reported that their symptoms improved, compared with only 10 out of 35 (29%) in the placebo group ($p < 0.001$).

Nausea outcomes

Willets and colleagues¹¹³ examined the effect of a ginger extract on pregnancy-induced nausea.¹¹³ The RINVR was used to assess nausea experience in terms of frequency, duration and distress (in addition to vomiting and retching) 1 hour after treatment was administered. For both the ginger extract and placebo groups, there was a reduction in the overall nausea experience score from baseline to day 1 of treatment, which then remained at this reduced level until day 4 of treatment. The effect of ginger extract relative to placebo on nausea experience was calculated as a difference parameter for 4 consecutive days of treatment post baseline. On days 1, 2 and 4, the authors state that the difference parameter was significantly less than zero, and that results showed a significant effect for ginger on nausea experience by/on day 4. However, the authors did not report p -values and added a note of caution as results were considered likely to be confounded by a regression-to-the-mean effect.

Mohammadbeigi and colleagues⁸⁹ also measured nausea severity with the RINVR. The observed difference in nausea severity between the second to the fifth days of the intervention and the first day was significant in all three groups ($p < 0.001$). However, there were differences observed in the intensity of changes in the ginger ($p < 0.01$) and metoclopramide ($p = 0.01$) groups compared with the placebo group. The difference in intensity was not different between the ginger and metoclopramide groups ($p = 0.68$).

Ozgoli and colleagues⁹⁶ investigated the effect of ginger capsules on NVP.⁹⁶ They measured nausea intensity twice a day for 4 days using a 10-point VAS, which was converted into author-defined categories of severe, moderate, mild and no nausea (where 10 = most severe and 0 = absence of nausea). Based on these categories, nausea improved in the ginger group compared with the placebo group: after treatment, 28% of women who had taken ginger capsules had no nausea compared with 10% in the placebo group ($p < 0.05$). There were also more women who reported severe nausea in the placebo group (15%) than in the ginger group (9%) ($p < 0.05$).

Vutyavanich and colleagues¹¹⁰ also used a VAS to assess severity of nausea. The mean change in nausea scores (baseline minus average post-therapy nausea scores for all women) was greater in the ginger group than in the placebo group ($p = 0.014$). To account for three missing patients in the placebo group, the authors assumed that their nausea scores changed as much as subjects with the best improvement for the purposes of intent-to-treat analysis. These results were sensitive to plausible extreme value sensitivity analysis and differences in nausea scores were no longer significant ($p = 0.08$). When the mean change in nausea scores were compared at baseline and at the end of the trial at day 4, there was a greater reduction in nausea scores in the ginger group than in the placebo group ($p = 0.035$ and $p = 0.005$ in analyses that did and did not impute for three missing placebo values respectively).

Vomiting outcomes

In the Willets and colleagues¹¹³ trial there was no difference between ginger extract and placebo groups for vomiting symptoms assessed using the RINVR.

Mohammadbeigi and colleagues⁸⁹ also measured vomiting severity with the RINVR.⁸⁹ Vomiting severity decreased in the ginger, metoclopramide and placebo arms between the second to fifth days of intervention compared with the first day ($p < 0.01$). There was an observed difference between the ginger

($p < 0.05$) and metoclopramide ($p = 0.02$) groups compared with the placebo group. However, there was no difference between the ginger and metoclopramide groups ($p = 0.72$).

In the trial of Ozgoli and colleagues,⁹⁶ there was a 50% decrease in the number of vomiting episodes in the ginger group post-treatment period ($p < 0.05$). In the control group, a 9% reduction was reported, which was not statistically significant. The between-group rate of reduction in number of vomiting episodes was also significant ($p < 0.05$).

After 4 days of treatment, the proportion of women in the Vutyavanich and colleagues trial¹¹⁰ who had vomiting was lower in the ginger compared with the placebo group (37.5% vs. 68.0%; $p = 0.021$). Similarly, a greater reduction in vomiting episodes was found in the ginger group compared with placebo (based on baseline number of episodes minus average number of vomiting episodes over the 4 days of treatment) ($p < 0.001$).

Retching outcomes

In the Willetts and colleagues¹¹³ trial, using the RINVR to assess symptoms, ginger extract was shown to lower retching symptom scores compared with the placebo group for the first 2 of 4 consecutive days of treatment.

Safety outcomes

Fetal outcomes and maternal adverse events were reported in four out of five included trials.^{74,89,96,110} Fischer-Rasmussen and colleagues⁷⁴ reported one miscarriage and one termination in trial participants. These data are summarised in the text following but it should be noted that given the anticipated rarity of these events, small trials are likely to provide unreliable estimates. No congenital anomalies were reported and the mean birthweight/gestation at delivery were within normal ranges. No adverse effects were reported in the trial of Ozgoli and colleagues⁹⁶ Vutyavanich and colleagues¹¹⁰ reported one miscarriage in the intervention ginger and three in placebo group, delivery at term in the majority of pregnancies in both groups (placebo = 91.4%, ginger = 96.9%; no p -value reported) and no congenital anomalies. See *Appendix 8, Secondary outcome data* for full details, with additional UKTIS ginger data provided in *Appendix 7*.

Ginger syrup versus placebo syrup

The trial of Keating and Chez⁸² reported on the effect of ginger syrup versus non-ginger-containing placebo syrup on nausea and vomiting in early pregnancy. This trial was judged as carrying a low risk of bias.

Combined severity score

No combined score reported.

Nausea outcomes

Nausea severity was measured using the 10-point VAS: 10 out of 13 (77%) women who received ginger had at least a 4-point improvement on the 10-point nausea scale by day 9; whereas only 2 out of 10 (20%) of the women in the placebo group had the same improvement. Conversely, no woman in the ginger group, but seven (70%) in the placebo group, had a ≤ 2 -point improvement on the nausea scale at both 9 and 14 days. It was not reported whether or not these differences were statistically significant.

Vomiting outcomes

Eight out of 12 women in the ginger group who were vomiting daily at the beginning of treatment stopped vomiting by day 6 compared with 2 out of 10 (20%) women in the placebo. Again, it was not reported whether or not this difference was statistically significant.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No safety-related outcome data were reported by Keating and Chez.⁸² See *Appendix 7* for additional UKTIS ginger data.

Ginger biscuit versus placebo biscuit

We identified one trial,⁶⁰ judged at low risk of bias, which reported on the effect of ginger biscuits on nausea and vomiting in early pregnancy.

Author-defined symptom severity/relief scales

Basirat and colleagues⁶⁰ used a 5-item Likert scale to assess patients' subjective response to treatment in terms of whether symptoms had worsened, improved or remained the same. They found that a higher proportion of women consuming ginger compared with placebo biscuits reported improved symptoms: 28 out of 32 (88%) compared with 21 out of 30 (70%) ($p = 0.043$).

Nausea outcomes

Nausea severity was assessed via a 10-point VAS. The average improvement in nausea scores (baseline minus average post-therapy nausea scores of day 1–4 for all subjects) in the ginger group was significantly greater than that in the placebo group. The nausea score [standard deviation (SD)] on day 4 in the placebo and ginger groups decreased to 3.03 (SD 2.47) from 4.67 (SD 1.97) and 3.03 (SD 2.19) from baseline score of 5.88 (SD 1.83), respectively ($p = 0.01$).

Vomiting outcomes

The average reduction in the number of vomiting episodes (baseline minus average post-therapy nausea scores of day 1–4 for all subjects) was greater in the ginger biscuit group (0.96 ± 0.21) than in the placebo biscuit group (0.62 ± 0.19). However, this difference was not significant ($p = 0.243$). After 4 days of treatment, the proportion of women who had no vomiting in the ginger group (11/32, 34%) was greater than that in the placebo group (6/30, 20%). It was not reported whether or not this difference was statistically significant.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No safety-related outcome data were reported by Basirat and colleagues.⁶⁰ See *Appendix 7* for additional UKTIS ginger data.

Ginger versus vitamin B6

Six trials compared ginger with vitamin B6 capsules for the treatment of NVP in pregnancy.^{67,70,77,92,102,103} Most of these trials were judged as being at a low risk of bias.^{67,70,102,103} However, risk of bias in the trial of Haji Seid Javari and colleagues⁷⁷ was judged as unclear, and Nanreji and colleagues⁹² was judged as being at a high risk of bias due to concerns with regard to the blinding process and allocation concealment.

Rhodes Index of Nausea, Vomiting and Retching

Chittumma and colleagues⁶⁷ used a modified version of the RINVR to assess change in nausea and vomiting severity (comprised episodes of nausea, duration of nausea and numbers of vomits). The trial found a reduction in symptoms in both groups [ginger from 8.7 (SD 2.2) to 5.4 (SD 2.0); vitamin B6 from 8.3 (SD 2.5) to 5.7 (SD 2.3); $p < 0.05$]. However, the mean score reduction (i.e. improvement in symptoms) with ginger was greater than with vitamin B6 (3.3 ± 1.5 vs. 2.6 ± 1.3) ($p < 0.05$).

Pregnancy-Unique Quantification of Emesis and Nausea scale

Haji Seid Javadi and colleagues⁷⁷ measured changes in symptoms using the PUQE scoring system (capturing numbers of nausea, vomiting and retching episodes, along with severity). The study found an improvement in PUQE scores between baseline and the subsequent 4 days of treatment [ginger 8.32 (SD 2.19), vitamin B6 7.77 (SD 1.80); $p < 0.001$]. However, this difference between groups was not significant ($p = 0.172$).

Author-defined symptom severity/relief scales

At the 1-week follow-up point, Ensiyeh and Sakineh⁷⁰ used a 5-point Likert scale (much worse, worse, same, better, much better) to assess overall treatment response. On day 7, 29 out of 35 (82.9%) women in the ginger group reported an improvement in general symptoms, compared with 23 out of 34 (67.6%) in the vitamin B6 group. However, this difference was not significant ($p = 0.52$).

Nausea outcomes

Sripramote and Lekhyananda¹⁰³ used a 10-point VAS to measure nausea severity at baseline (pre treatment) and days 1–3 (post treatment).¹⁰³ Comparing baseline and post treatment, there was a reduction in mean score change in both ginger (1.4, SD 2.22) and vitamin B6 groups (2.0, SD 2.19) ($p < 0.001$ for both groups). However, the difference between groups was not statistically significant ($p = 0.136$).

Ensiyeh and Sakineh⁷⁰ also used a 10-point VAS to assess nausea severity between baseline and the first 4 days of treatment. They found a greater mean change in nausea score over the 4 days in the ginger group (2.2 ± 1.9) compared with the vitamin B6 group (2.2 ± 1.9) ($p = 0.024$).

Smith and colleagues¹⁰² used the RINVR to assess differences in nausea severity between baseline and at days 7, 14 and 21. Equivalent reductions in mean differences over time were reported for both treatment groups (mean difference 0.2, 90% CI -0.3 to 0.8).

Narenji and colleagues⁹² used a 10-point VAS to assess nausea severity. However, although both groups reduced their mean nausea scores [vitamin B6 mean change 0.7 (SD 1.99), ginger mean change 1.0 (SD 1.32)], there was no difference between groups ($p = 0.8$).

Vomiting outcomes

Sripramote and Lekhyananda¹⁰³ assessed vomiting severity by the change in number of vomiting episodes. Both groups showed a reduction in the mean number of vomiting episodes [ginger group 0.7 (SD 2.18), $p = 0.003$; vitamin B6 group 0.5 (SD 1.44), $p = 0.008$]. However, the mean change difference between groups was not significant.

Ensiyeh and Sakineh⁷⁰ also assessed the change in number of vomiting episodes between baseline and the first 4 days of treatment. Although number of vomiting episodes decreased in both groups (ginger 0.6 ± 0.7 , vitamin B6 0.5 ± 1.1), this difference was not statistically significant (p -value reported as $p = 1.101$ in the paper, which we assume is a reporting error).

Narenji and colleagues⁹² also found a reduction in the number of episodes of vomiting in both groups. However, they reported no difference between groups (although did not provide a p -value).

Smith and colleagues¹⁰² used the RINVR to assess differences in vomiting severity between baseline and at days 7, 14 and 21. Equivalent reductions in mean differences over time were reported for both treatment groups (mean difference 0.5, 90% CI 0.0 to 0.9).

Retching outcomes

Smith and colleagues¹⁰² used the RINVR to assess differences in retching severity between baseline and at days 7, 14 and 21. Equivalent reductions in mean differences over time were reported for both treatment groups (mean difference 0.3, 90% CI -0.0 to 0.6).

In the trial of Haji Seid Javadi and colleagues,⁷⁷ changes in retching severity were assessed by PUQE scores. There was no difference in the reduction of retching between the vitamin B6 group and ginger groups ($p = 0.333$).

Safety outcomes

Fetal outcomes and maternal adverse events were reported in four out of the six included trials.^{67,70,102,103} These data are summarised in the text following but as noted previously, given the anticipated rarity of these events, small trials are likely to provide unreliable estimates.

No pregnancy outcomes were reported by either Sripramote and Leekhyananda¹⁰³ or Chittumma and colleagues,⁶⁷ and both trials found only minor side effects in both groups. Sripramote and Leekhyananda¹⁰³ found no difference in rates of sedation (ginger 26.6% vs. vitamin B6 32.8%; $p = 0.439$), or heartburn (ginger 9.4% vs. vitamin B6 6.3%; $p = 0.510$). In the trial of Chittumma and colleagues,⁶⁷ minor side effects such as sedation, heartburn and arrhythmia were reported, but the difference between groups was not significant (% experiencing side effects: ginger 25.4%, vitamin B6 23.8%; $p = 0.80$).

Ensiyeh and Sakineh⁷⁰ reported slight drowsiness in 7% of the hydroxyzine-treated patients. In terms of miscarriages, the ginger group reported two events and the vitamin B6 group reported one event. However, no congenital abnormalities or neonatal problems were reported in either group. Smith and colleagues¹⁰² reported no differences between groups in terms of live births, congenital abnormalities or birthweight ($p > 0.05$). See *Appendix 8, Secondary outcome data* for full details, with additional UKTIS ginger data provided in *Appendix 7*.

Ginger capsules versus acupressure

The trial of Saberi and colleagues¹³ reported a randomised comparison of the effectiveness of ginger capsules versus acupressure to relieve NVP. A control arm was also included where no intervention was performed during the 7 days of the trial. Analysis was performed on all participants. This study was judged as carrying low risk of bias.

Rhodes Index of Nausea, Vomiting and Retching

The authors calculated the mean difference in the overall RINVR scores between the three groups by subtracting the pre-intervention (3 days before intervention) from the post-intervention (4 days after treatment) scores. It was significantly greater ($p < 0.001$) in the ginger group (8.61 ± 5.32) than in the acupressure (4.17 ± 5.53) and control groups (-0.84 ± 3.72). The authors reported that the total RINVR scores was reduced by 49% in the ginger group and by 29% in the acupressure group. There was little change in the control group (raised to 0.06%). It was not reported if the difference between these three groups was significant.

Nausea outcomes

The difference in nausea severity (as assessed by the RINVR in the same way as above) was found to be lower in the ginger group (3.94 ± 2.58) compared with the acupressure (2.00 ± 2.37) or control groups (0.18 ± 1.24) ($p = 0.001$ for differences between groups). A similar trend was observed in the reduction percentage of scores (ginger 48%, acupressure 29%, control 0.03%), although it was not reported if this difference was significant.

Vomiting outcomes

The difference in vomiting severity (as assessed by the RINVR in the same way as above) was found to be lower in the ginger group (2.66 ± 2.64) compared with the acupressure (0.64 ± 2.14) or control groups (-0.17 ± 2.12) ($p = 0.001$). A similar trend was observed in the reduction percentage of scores (ginger, -52%; acupressure, 19%; control, -0.24%), although it was not reported if this difference was significant.

Retching outcomes

The difference in retching severity (assessed as above) was also found to be lower in the ginger group (2.01 ± 1.56) compared with the acupressure (1.52 ± 1.86) or control groups (0.31 ± 1.36) ($p = 0.001$ for

differences between groups). A similar trend was observed in the percentage reduction of scores (ginger 46%, acupuncture 37%, control -0.09%), although it was not reported if this difference was significant.

Safety outcomes

Saberi and colleagues¹³ did not report pregnancy-related outcomes and found no side effects in trial participants. See *Appendix 7* for additional UKTIS ginger data.

Ginger versus doxylamine–pyridoxine

Biswas and colleagues⁶³ reported a randomised, comparison of ginger extract tablets versus doxinate (doxylamine 10 mg plus pyridoxine 10 mg) in the treatment of nausea and vomiting during pregnancy. Patients were blinded to the treatment, whereas investigators were not. However, overall, this study was judged to have a low risk of bias.

Combined severity score

No combined score reported.

Nausea outcomes

Nausea was assessed in terms of severity (using the 10-point VAS) and number of spells. Both groups showed a significant decline in nausea severity median score {ginger from 34.5 [interquartile range (IQR) 0.0–91.0] to 0.00 [IQR 0.0–52.0]; doxylamine–pyridoxine from 30.4 [IQR 0.0–100.0] to 0.0 [IQR 0.0–65.0]} and number of nausea spells {ginger from 3.0 [IQR 1.0–10.0] to 0.43 [IQR 0.00–12.00]; doxylamine–pyridoxine from 4.0 [IQR 0.0–12.00(sic)] to 0.6 [IQR 0.0–7.7]} ($p = 0.001$). However, between group differences were not significant at any time point.

Vomiting outcomes

Vomiting was assessed in terms of severity (using the 10-point VAS) and number of episodes. Again, both groups showed a decline in severity [ginger from median 14.00 (range 0.00–73.00) to median 0.00 (range 0.00–30.00); doxylamine–pyridoxine from median 22.00 (range 0.00–87.00) to median 0.00 (range 0.00–47.00)] and vomiting episodes [ginger from median 0.14 (range 0.00–5.50) to median 0.15 (range 0.00–5.50); doxylamine–pyridoxine from median 2.00 (range 0.00–6.00) to median 0.00 (range 0.00–2.80)]. However, as above, between group differences were not significant at any time point.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No stillbirths, congenital anomalies, neonatal or fetal complications were reported in the trial of Biswas and colleagues⁶³ See *Appendix 7* for additional UKTIS ginger data.

Ginger versus antihistamine (dimenhydrinate) capsules

Pongrojpraw and colleagues⁴² examined the effect of ginger versus antihistamine (dimenhydrinate) capsules in the treatment of NVP. Risk of bias was unclear due to lack of information provided in this abstract.

Combined severity score

No combined score reported.

Nausea outcomes

The mean nausea score during days 1–7 of the treatment decreased in both groups. There was no significant difference in the daily mean nausea scores between both groups ($p > 0.05$).

Vomiting outcomes

The frequency of vomiting episodes during days 1–7 of the treatment decreased in both groups. There was no significant difference in the daily mean number of vomiting episodes between days 3–7 post treatment ($p > 0.05$), although the daily mean number of vomiting episodes in the dimenhydrinate group during days 1 and 2 of the treatment was less than in the ginger group ($p < 0.05$).

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes were reported; however, Pongrojpraw and colleagues⁴² found a significant difference in drowsiness after treatment (dimenhydrinate 77.6% compared with ginger 5.9%; $p < 0.01$). These data are summarised in the text following, but it should be noted that given the anticipated rarity of these events small trials are likely to provide unreliable estimates. See *Appendix 8, Secondary outcome data* for full details, noting the aforementioned caveat in relation to generalisability of this safety data, with additional UKTIS ginger data provided in *Appendix 7*.

Ginger versus metoclopramide

The 2011 trial of Mohammadbeigi and colleagues⁸⁹ randomised women to ginger, metoclopramide or placebo capsules for the treatment of pregnancy sickness. Details are reported in *Ginger capsules versus placebo capsules* concerning ginger versus placebo. In summary, both were found to be efficacious but there was no evidence that one treatment was superior to the other. Furthermore, as detailed above, the trial was judged as carrying a high risk of bias due to concerns over lack of blinding of trial personnel.

Summary

- The evidence available for ginger was predominantly at low risk of bias or the risk of bias was unclear, with four exceptions.^{77,89,92,113}
- Ginger is one of the most widely reported interventions but the evidence is limited and generalisability relates to women with less severe nausea and vomiting symptoms receiving ginger treatment.
- The quality of the evidence for ginger versus placebo is low and was downgraded due to clinical heterogeneity and imprecision. Further research is very likely to have an important impact on our confidence in the estimate of effect and may change the estimate. There was a sparseness of data in the other comparisons, so they were further downgraded to very low quality of evidence and we are very uncertain about the estimates.
- The identified studies comparing ginger preparations with placebo generally reported evidence of an improvement over a range of symptoms with the use of ginger.
- When consideration is restricted to those studies at low risk of bias, the results are less clear cut. Ginger looks promising in reducing symptoms, but the findings are not conclusive.
- For the comparison of ginger preparations with acupuncture, ginger again looks promising, but the evidence is very limited in both quantity and quality.
- For the comparison of ginger versus vitamin B6 there are some higher-quality studies, but in general there was little evidence of a difference in the severity of symptoms between groups. There were few data for the comparisons of ginger against doxylamine–pyridoxine or antihistamine or metoclopramide, and minimal evidence suggesting any difference between groups.
- Overall, there is a suggestion that ginger might be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required for this and all other comparators.

Chapter 5 Clinical effectiveness: acupressure, acupuncture and nerve stimulation

Introduction

Acupuncture or acupressure was used as an intervention for NVP in a total of 18 RCTs and one case series study. The majority ($n = 13$) of studies compared acupressure, acupuncture or nerve stimulation against placebo or sham treatment. However, heterogeneity was observed in relation to the clinical setting and patient populations in which the studies were conducted, as well as the interventions, comparators and outcomes reported in each trial. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*), given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses. We have thus reported a narrative summary only for each intervention and comparator set, with additional outcome data presented in *Table 9*.

Of the identified trials, eight were judged as being low risk of bias,^{61,62,66,80,83,91,98,101} with a further six unclear due to lack of sufficient information on the research process.^{43,73,78,79,104,109} Four were judged as carrying high risk of bias on the basis of incomplete outcome data and weak randomisation process,¹¹⁵ lack of blinding,⁸⁷ or selective outcome reporting.^{94,111,115} We also identified a case series which evaluated the effect of P6 acupressure by Markose and colleagues,¹²⁴ which was categorised as weak based on the EPHPP tool.

The severity of symptoms for patient participants in the studies comprising this group of interventions varied substantially. Three studies were classed as focussing on women with mild NVP,^{61,62,104} seven with mild to moderate,^{43,80,83,91,98,101,124} two with moderate severity,^{78,115} four with moderate to severe,^{66,87,94,111} and a final three for which severity was not possible to classify.^{73,79,109}

Acupressure versus placebo

Eight trials compared acupressure with placebo or sham treatment.^{43,61,62,78,79,91,104,111} Seven trials^{43,61,62,78,79,104,111} involved the comparison of acupressure administered to the P6 acupoint, which lies between the tendons of the palmaris longus and flexor carpi radialis muscles, four centimetres proximal to the wrist crease.¹²⁹ Six of these trials examined the application of acupressure to point P6 by the use of pressure bands worn by study participants,^{43,61,78,79,104,111} one comprised self-administered acupressure at the P6 acupoint.⁶² A final acupressure versus placebo trial compared acupressure self-administered with the two KID21 acupoints, which are located above the navel.⁹¹ The placebo comparison intervention came in a two formats. The vast majority of studies (7/8) compared the active acupressure intervention with the application of acupressure to a sham acupoint (i.e. at which there is no known acupressure point or meridian pathway).^{43,61,62,78,79,91,111} One final study compared the use of pressure bands with the wearing of placebo pressure bands which did not have a pressure button.¹⁰⁴ Three of these trials were judged as having low risk of bias^{61,62,91} and four were judged to be unclear.^{43,78,79,104} Werntoff and Dykes¹¹¹ was judged as having high risk of bias on the basis of incomplete outcome data and weak randomisation process.

Rhodes Index of Nausea, Vomiting and Retching

Belluomini and colleagues⁶² used the RINVR as a combined measure of change in nausea, vomiting and retching between the pre-treatment period (mean of days 1–3) and the post-treatment period (mean of days 5–7). The study found a significant reduction in combined scores for both groups (intervention from 12.64 ± 5.7 to 8.69 ± 5.0 ; $p \leq 0.001$; placebo from 11.47 ± 4.9 to 10.03 ± 4.6 ; $p = 0.019$).

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Acupressure vs. placebo									
Bayreuther 1994 ⁶¹	General practices, Southampton, UK	Effectiveness of acupressure at P6, RCT	n = 23 (I n = 11; comparator n = 12)	Patients presenting with early morning sickness	NR (MILD)	Sea-bands at P6 point for 7 consecutive days followed by 2 days no treatment	Sea-bands at placebo position for 7 consecutive days followed by 2 days no treatment	VAS for nausea Episodes of emesis	Mean levels of nausea between conditions/treatment difference: (1) paired t-test 1.69; (2) two-sample t-tests 1.67; (3) Wilcoxon signed-rank test 1.65; (4) Mann-Whitney U-test 1.61
Belluomini 1994 ⁶²	Physician and midwife practices, San Francisco, CA	Effectiveness of acupressure in reducing nausea and vomiting of pregnancy, RCT	n = 60 (I n = 30; comparator n = 30). At study entry: (1) 18.5 ± 1.4; (2) comparator 8.6 ± 1.4	Women complaining of nausea with or without emesis	Baseline nausea scores: I = 8.38 ± 2.2, C = 7.99 ± 2.5; baseline emesis scores: I = 2.09 ± 2.5, C = 1.83 ± 2.7 Total scores baseline: (1) 12.64 ± 5.7; (2) comparator 11.47 ± 4.9 (MILD)	3-day control period (no treatment) then self-administered acupressure for 10 minutes four times a day for 7 consecutive days at point PC-6	3-day control period (no treatment) followed by self-administered acupressure for 10 minutes four times a day for 7 consecutive days at a placebo point	RINVR	Nausea scores post treatment: (1) 15.80 ± 2.9 (p ≤ 0.001); (2) comparator 7.04 ± 2.6 (p ≤ 0.001) Emesis scores post treatment: (1) 1.28 ± 1.9 (p = 0.03); (2) comparator 1.63 ± 2.3 (p-value not reported) Total scores post treatment: (1) 18.69 ± 5.0 (p ≤ 0.001); (2) comparator 10.03 ± 4.6 (p = 0.019)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Can Gurkan 2008 ⁴³	Antenatal polyclinic of a maternity and child hospital in Istanbul, Turkey	Effect of acupressure on nausea and vomiting during pregnancy, RCT	n = 75 (I n = 26; C n = 25; placebo n = 24)	Women with a nausea severity of at least 50 using VAS, with or without vomiting in the last 24 hours	NR (MILD-MODERATE)	Three-step process: (1) days 1-3 no treatment; (2) days 4-6 acupressure wristbands applied to the P6 point to be only taken off at night when going to sleep and replaced before leaving their beds in the morning; and (3) days 7-9 no treatment (bands were not worn)	Three-step process: (1) days 1-3 no treatment; (2) days 4-6 acupressure wristbands applied to a placebo point (upper side of the wrist, being the opposite direction of the P6 point) to be only taken off at night when going to sleep and replaced before leaving their beds in the morning; and (3) days 7-9 no treatment (bands were not worn)	VAS for nausea Episodes of vomiting	Frequency of nausea: First 3 days compared with second 3 days: (1) $z = -3.35$ ($p < 0.001$); (2) placebo $z = -0.28$ ($p < 0.05$); (3) C $z = -0.92$ ($p > 0.05$) Second 3 days compared with third 3 days: (1) $z = -1.28$ ($p > 0.05$); (2) placebo $z = 10.18$ ($p > 0.05$); (3) C $z = -1.40$ ($p > 0.05$) Severity of nausea: First 3 days compared with second 3 days: (1) $z = -3.04$ ($p < 0.05$); (2) placebo $z = -2.96$ ($p < 0.05$); (3) C $z = -1.02$ ($p > 0.05$) Second 3 days compared with third 3 days: (1) $z = -1.48$ ($p > 0.05$); (2) placebo $z = -1.26$ ($p > 0.05$); (3) C $z = 0$ ($p > 0.05$) Intensity of discomfort felt from nausea: First 3 days compared with second 3 days: (1) $z = -3.7$ ($p < 0.001$); (2) placebo $z = -2.4$ ($p < 0.05$); (3) C $z = -1.2$ ($p > 0.05$) Second 3 days compared with third 3 days: (1) $z = -0.22$ ($p > 0.05$); (2) placebo $z = -1.4$ ($p > 0.05$); (3) C $z = -0.44$ ($p > 0.05$)

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
									Number of vomiting episodes:
									First 3 days compared with second 3 days: (1) $I z = -1.38$ ($p > 0.05$); (2) placebo $z = -1.85$ ($p > 0.05$); (3) C $z = -0.37$ ($p > 0.05$)
									Second 3 days compared with third 3 days: (1) $I z = -1.3$ ($p > 0.05$); (2) placebo $z = -0.7$ ($p > 0.05$); (3) C $z = -0.57$ ($p > 0.05$)
									Cross-comparison of symptoms in treatment group and placebo group on days 4–6:
									Number of nausea: (1) $I MR = 24.52$; (2) placebo group $MR = 26.56$, $p > 0.05$
									Severity of nausea: (1) $I MR = 24.9$, $U = 296.5$, $p > 0.05$; (2) placebo $MR = 26.15$, $U = 296.5$, $p > 0.05$
									Intensity of discomfort: (1) $I MR = 23.98$, $U = 272.5$, $p > 0.05$; (2) placebo $MR = 27.15$, $U = 272.5$, $p > 0.05$
									Number of vomiting: (1) $I MR = 22.67$, $U = 238.5$, $p > 0.05$; (2) placebo $MR = 28.56$, $U = 238.5$, $p > 0.05$

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Heazell 2006 ⁷⁸	Inner city secondary care centre, Manchester, UK	Efficacy of acupressure at the P6 point for the inpatient treatment of severe nausea and vomiting in early pregnancy, RCT	n = 80 (I n = 40; comparator n = 40). At study entry: (1) 1 – 8.5 (standard error of the mean 0.32, range 6–14); (2) comparator – 9.0 (standard error of the mean 0.36, range 5–14)	Women with NVP on their first inpatient admission (at least 2+ of ketonuria on urinalysis, an inability to tolerate oral fluids, and a requirement for antiemetic medication)	NR (MODERATE)	Acupressure bead placed at the P6 meridian point for 8 hours a day, from 9 a.m. to 5 p.m. In addition, patients were treated according to a standard protocol: 3 l of i.v. fluid in 24 hours and parenteral antiemetic medication while the patient was unable to tolerate oral fluids and thiamine (100 mg) that was taken orally once daily. When the patient could tolerate oral fluids, the antiemetic medication was administered orally. There was a defined antiemetic protocol that used cyclizine as a first-line agent, prochlorperazine as a second-line agent and metoclopramide, ondansetron or phenothiazine as a third-line agent	Patients assigned to the placebo group had the beads placed at a site on the dorsal aspect of the forearm which is not thought to be effective for 8 hours a day, from 9 a.m. to 5 p.m. In addition, patients were treated according to a standard protocol: 3 l of i.v. fluid in 24 hours and parenteral antiemetic medication while the patient was unable to tolerate oral fluids and thiamine (100 mg) that was taken orally once daily. When the patient could tolerate oral fluids, the antiemetic medication was administered orally. There was a defined antiemetic protocol that used cyclizine as a first-line agent, prochlorperazine as a second-line agent and metoclopramide, ondansetron or phenothiazine as a third-line agent	Not reported	Not reported

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Hsu 2003 ²⁹	Emergency department (triage), hospital setting, USA	Effect of acupressure wristband on pregnancy-related nausea/vomiting during the initial period of emergency department evaluation, RCT	n = 77 (I n = 38; comparator n = 39). At study entry: (1) I = 9 (range 3–18); (2) comparator = 9 (range 3–18)	Women with nausea/vomiting related to pregnancy	NR (NOT CLEAR)	Acupressure device placed over the P6 point for 60 minutes	Acupressure device placed over the sham site for 60 minutes	McGill Nausea Questionnaire	No difference between I and comparator was detected at baseline, 30 or 60 minutes for any of the indexes ($p < 0.2$ for all)
Naeimi Rad 2012 ³¹	Prenatology clinic, Rouhani Hospital, Babol University of Medical Science, Iran	Efficacy of KID21 point (Youmen) acupressure on nausea and vomiting of pregnancy, RCT	n = 80 (I = 40; comparator = 40). At study entry: (1) I 9.55 ± 1.81; (2) comparator 9.45 ± 2.02	Women with moderate to severe nausea and vomiting, but normal electrolytes	Nausea intensity before acupressure: median I = 8 (IQR 10–7); median C = 8 (IQR 9–7). Vomiting intensity before acupressure: median I = 2 (IQR 4–1); median C = 2 (IQR 3–1) (MILD–MODERATE)	Provision of routine advice to reduce nausea and vomiting via educational pamphlets. All women took vitamin B6 (40 mg twice daily). Acupressure to the KID21 points for 20 minutes for 4 consecutive days. Women were also advised to self-administer acupressure whenever they felt nausea and vomiting to the KID21 point	Provision of routine advice to reduce nausea and vomiting via educational pamphlets. All women took vitamin B6 (40 mg twice daily). Acupressure to a false point, gradually increasing in pressure, for 20 minutes for 4 consecutive days	VAS for nausea	Intensity of nausea, median VAS scores (IQR): (1) I first day = 7 (IQR 8–6), second day = 6 (IQR 7.75–4), third day = 5 (IQR 5–3), fourth day = 4 (IQR 5–2); (2) comparator first day = 7 (IQR 8–6), second day = 7 (IQR 8–6), third day = 7 (IQR 8–5), fourth day = 7 (IQR 8–5) Frequency of nausea, median VAS scores (IQR): (1) I first day = 1 (IQR 2–0), second day = 0 (IQR 1–0), third day = 0 (IQR 1–0), fourth day = 0 (IQR 0.75–0); (2) comparator first day = 1 (IQR 2–1), second day = 1 (IQR 2–0.25), third day = 1 (IQR 2–0), fourth day = 1 (IQR 2–0)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Steele 2001 ¹⁰⁴	Obstetric and gynaecology offices and clinics, Michigan, MI, USA	Effect of acupressure by sea-bands on nausea and vomiting of pregnancy, RCT	n = 110 (I = 68; comparator = 42)	Women who self-report of one or more episodes of pregnancy-related nausea and/or vomiting	NR (MILD)	Sea-bands with acupressure buttons on each wrist for 7 consecutive days (removed only when bathing) followed by a 72-hour no-treatment control period	Placebo sea-bands without acupressure buttons on each wrist for 7 consecutive days (removed only when bathing) followed by a 72-hour no-treatment control period	Author defined (four closed-ended questions on frequency and severity of nausea with responses recorded on a 5-point Likert scale)	Nausea frequency (days 1–4): (1) IMR = 40.30; (2) comparator MR = 72.82 Nausea severity (days 1–4): (1) IMR = 40.13; (2) comparator MR = 73.09 Vomiting frequency (days 1–4): (1) IMR = 41.51; (2) comparator MR = 70.94 Vomiting severity (days 1–4): (1) IMR = 39.28; (2) comparator MR = 73.65 Nausea frequency (days 5–7): (1) IMR = 47.18; (2) comparator MR = 58.47 Nausea severity (days 5–7): (1) IMR = 49.46; (2) comparator MR = 54.79 Vomiting frequency (days 5–7): (1) IMR = 50.44; (2) comparator MR = 53.22 Vomiting severity (days 5–7): (1) IMR = 46.63; (2) comparator MR = 58.24

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Werntoft 2001 ¹¹¹	Antenatal clinics, Sweden	Effect of acupressure on nausea and vomiting during pregnancy, three-arm RCT	n = 60 (I = 20; placebo = 20; no treatment = 20). At onset: (1) I = 5.8 (SD 2.3); (2) placebo = 5.6 (SD 1.2); (3) no treatment = 5.6 (SD 2.0). At study entry: (1) I = 9.8 (SD 1.9); (2) placebo = 9.6 (SD 1.6); (3) no treatment = 10.8 (SD 2.2)	Women described as experiencing NVP	VAS: P6 = 8.4 (SD 1.2); placebo = 8.4 (SD 1.4); no treatment = 8.0 (SD 1.5) (MODERATE-SEVERE)	Acupressure wristbands with button at P6 point worn for 2 weeks, removing it only when showering	Acupressure wristbands with button at placebo point on the upper side of the wrist, worn for 2 weeks, removing it only when showering. A third group received no treatment	VAS for nausea	Before pregnancy: (1) I 1.4 (SD 1.4); (2) placebo 1.1 (SD 0.9); (3) no treatment 1.5 (SD 2.4) Before treatment: (1) I 8.4 (SD 1.2); (2) placebo 8.4 (SD 1.4); (3) no treatment 8.0 (SD 1.5) After 1 day: (1) I 5.2 (SD 2.7); (2) placebo 5.6 (SD 2.5); (3) no treatment 7.6 (SD 1.6) After 3 days: (1) I 5.6 (SD 2.3); (2) placebo 5.5 (SD 2.8); (3) no treatment 7.2 (SD 1.3) After 6 days: (1) I 4.9 (SD 2.4); (2) placebo 6.3 (SD 2.4); (3) no treatment 6.9 (SD 2.0) After 14 days: (1) I 4.2 (SD 2.6); (2) placebo 5.9 (SD 2.4); (3) no treatment 6.5 (SD 2.2)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Nerve stimulation vs. placebo									
Evans 1993 ⁷³	Obstetrics clinic, University of California, CA, USA	Effect of stimulation of a region on the wrist by either surface pressure or micro-voltage electrical current on NVP, randomised crossover trial	n = 25	Women diagnosed with NVP by their physician	NR (NOT CLEAR)	An active battery-powered SAS wrist unit was provided to participants which they were told would produce a perceptible tingling sensation in the wrist and hand when active, while the inactive unit would produce only a pressure sensation on the wrist when stimulating the median nerve. Each subject was instructed to wear the device continually for 48 hours successively.	Not applicable	Author-defined subjective change in nausea and vomiting (improved, worsened, no change)	Average nausea score: (1) active unit 2.4; (2) placebo 2.7 ($p < 0.05$) Improvement in nausea: (1) active unit 15; (2) placebo 10 ($p < 0.05$)
Rosen 2003 ⁸⁸	Hospital clinics and physicians' private offices, Morristown, New Jersey, NJ, USA, Eastern Virginia Medical School, Norfolk, VA, USA, University of Arizona Health Sciences Centre, Arizona, AZ, USA, and New York University School of Medicine, New York, NY, USA	To evaluate the effectiveness of low-level nerve stimulation therapy over the volar aspect of the wrist at the P6 point to treat nausea and vomiting in early pregnancy, multicentre RCT	n = 230	Women with symptoms of mild to severe nausea and vomiting	RINVR score: I = 13.5 (SD 6.0); C = 12.0 (SD 5.3) (MILD-MODERATE)	Nerve stimulation for 3 weeks via a ReliefBand Model WB-R (Woodside Biomedical Inc., Carlsbad, CA, USA) [a non-invasive, portable (34 g), battery-powered, watch-like acustimulation device]. A rotary dial on the device allows users to select between five intensities (n = 117)	Participants given an identical non-stimulating device (n = 113)	RINVR	Mean change from baseline: I = 6.48 (95% CI 5.31 to 7.66), C = 4.65 (95% CI 3.67 to 5.63); $p = 0.02$

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Veciana 2001 ¹⁰⁹	Obstetrics and gynecology, Eastern Virginia Medical School, Norfolk, VA, USA	RCT	n = 460 (I = 230; comparator = 230). At study entry: all 6–12 weeks	Women with symptoms of mild to severe nausea and vomiting	NR (NOT CLEAR)	Nerve stimulation device worn at P6 point	Non-stimulating device worn at P6 point	RINVR	No significant difference in RINVR score between groups
Jamigorn 2007 ⁵⁰	Antenatal clinic, department of obstetrics and gynaecology, faculty of medicine, Chulalongkorn University, Bangkok, Thailand	Effectiveness of acupressure compared with vitamin B6 in reducing NVP, RCT	n = 66 (I = 33; comparator = 33) At onset: (1) 18.1 ± 1.7; (2) comparator 8.9 ± 3.5 At study entry: (1) 16.2 ± 1.0; (2) comparator 6.8 ± 1.5	Women who suffered from mild to moderate nausea and/or vomiting	RINVR score: I = 14.3 ± 3.3; C = 15.4 ± 3.0 (MILD–MODERATE)	Acupressure wristbands (sea-bands) with a button on P6 point worn continuously as possible on days 1–5 plus a placebo tablet to mimic vitamin B6. Also instructed to use oral dimenhydrinate (50 mg) every 6 hours when they had nausea and vomiting, to divide their meals into frequent small ones rich in carbohydrates and low fat and advised not to take any other medications	Sea-bands with a button on a dummy point and 50 mg-tablets of vitamin B6 were prescribed every 12 hours for 5 days. Also advised to divide their meals into frequent small ones rich in carbohydrates and low fat and not to take any other medications	RINVR	Significant improvement of nausea, retching and vomiting symptoms in both acupressure (p < 0.001) and vitamin B6 groups (p < 0.001) No statistically significant differences in scores between baseline and end of treatment (evening of the fifth day) between both groups (p > 0.05)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Acupuncture vs. metoclopramide/vitamin B12									
Neri 2005 ⁹⁴	Department of obstetrics and gynaecology, University of Modena and Reggio Emilia, University of Turin, Italy	Efficacy of acupuncture sessions plus acupressure compared with metoclopramide/vitamin B12 treatment, RCT	n = 81 (I = 43; C = 38). (Both groups less than 12 weeks' gestation)	Diagnosis of HG following the commonly accepted criteria of nausea and vomiting leading to clinical symptoms of dehydration and > 5% weight loss	NR. States no difference between groups at baseline (MODERATE-SEVERE)	Twice-weekly (for 2 weeks) acupuncture with 0.3 mm diameter sterile disposable steel needles (length 52 mm) to a depth of 10–30 mm to points PC6, CV12 and ST36. manipulated until the patient reported the characteristic irradiating sensation, then left in situ for 20 minutes without any further manual stimulation. Patients also advised to wear an acupressure device (sea-band) at the PC6 point for 6–8 hours a day at home	Metoclopramide infusion (20 mg/500 ml saline for 60 minutes) at the hospital twice a week for 2 weeks plus oral supplementation with vitamin B12 complex [pyridoxine, hydroxycobalamin, 30 mg/day (Benadon® Roche)] was prescribed at home	Author defined (intensity of nausea, number of vomiting episodes and rate of food intake)	Nausea intensity, number of cases improved (%): (1) I first session 1 (2.3%); second session 11 (25.5%); third session 19 (44.1%); (2) comparator first session 1 (2.3%); second session 9 (23.6%); third session 12 (31.5%) Vomiting episodes, number of cases improved: (1) I first session 7 (16.2%); second session 15 (34.8%); third session 24 (55.8%); (2) comparator first session 4 (10.5%); second session 12 (31.5%); third session 14 (36.8%)

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Acupuncture vs. placebo									
Carlsson 2000 ⁸⁶	University hospital, department of obstetrics and gynaecology in Uppsala, Sweden	Effect of acupuncture plus standard treatment on HG, randomised crossover study	n = 33 (group A n = 17; group B n = 16)	Women suffering from HG and not responding to conventional outpatient treatment	No numerical values for severity at baseline provided. However, 25 women had previously tried antihistamines (MODERATE-SEVERE)	Group A: on days 1 and 2, active acupuncture was performed at point PC6 for 30 minutes three times per day. No acupuncture was performed on days 3 and 4 (wash-out period). On days 5 and 6, sham acupuncture (needles were inserted only 1–2 mm in the skin) was performed for 30 minutes three times per day. No acupuncture was performed on days 3 and 4 (wash-out period). On days 5 and 6, active acupuncture (needles were inserted only 1–2 mm in the skin) was performed for 30 minutes three times per day. During the study, the women also received parenteral nutrition with 5% glucose and patients were discharged on day 7	Group B: on days 1 and 2, sham acupuncture (needles were inserted only 1–2 mm in the skin) was performed for 30 minutes three times per day. No acupuncture was performed on days 3 and 4 (wash-out period). On days 5 and 6, active acupuncture was performed at point PC6 for 30 minutes three times per day. During the study, the women also received parenteral nutrition with 5% glucose and patients were discharged on day 7	VAS for nausea Episodes of vomiting	Nausea in active acupuncture: (1) group A pre-acupuncture, post-VAS reduction = 4; (2) group B pre-acupuncture, post-VAS reduction = 3 Nausea in placebo acupuncture: (1) group B pre-placebo, post-VAS-reduction = 1.7; (2) group A pre-acupuncture, post-VAS reduction = 0.1 Vomiting occurrence: (1) 17 out of 17 women were still vomiting after 2 acupuncture days; (2) comparator 12 out of 16 women were still vomiting after 2 placebo treatment days

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Knight 2001 ⁸³	Maternity unit, Royal Devon and Exeter Hospital, Devon, UK	Effectiveness of acupuncture compared with placebo treatment of nausea of pregnancy, RCT	n = 55 (n = 28; comparator n = 27) (1) 17.8 ± 1.0; (2) comparator 8.0 ± 1.0	Women complaining of nausea, with or without vomiting, who approached a community midwife	Nausea scores, median (IQR): I = 85.5 (IQR 71.25–89.75); sham = 87.0 (IQR 73.0–90.0) (MILD–MODERATE)	At the first visit a traditional Chinese medical diagnosis was made and each woman was allocated one of three categories and treated with acupuncture to a combination of the following points: stomach 36; conception vessel 12; spleen 4; P6; stomach 44; and stomach 34 and P6. 40 x 0.25-mm needles (Seirin, Japan) inserted to a depth of 0.5–1.0 cm with the assistance of a guide tube, then manipulated. Needles were left in place for 15 minutes, given twice in the first week and once weekly for 2 weeks	Sham treatment consisted of tapping a blunt cocktail stick, supported by a plastic guide tube, over a bony prominence in the region of each acupuncture point. Sham needles were left in place for 15 minutes, given twice in the first week and once weekly for 2 weeks	VAS for nausea Episodes of vomiting	Median nausea score (IQR): (1) 1 day 1 85.5 (IQR 71.25–89.75), 3 days after session one 63.0 (IQR 50.75–86.5), 3 days after session two 65.0 (IQR 36.25–79.5), 3 days after session three 44.0 (IQR 29.0–77.25), 3 days after session four 47.5 (IQR 29.25–69.5); (2) comparator day 1 87.0 (IQR 73.0–90.0), 3 days after session one 69.0 (IQR 45.0–87.0), 3 days after session two 61.0 (IQR 30.0–80.0), 3 days after session three 53.0 (IQR 25.0–80.0), 3 days after session four 48.0 (IQR 14.0–80.0)

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Smith 2002 ¹⁰¹	Women's and children's hospital in Adelaide, SA, Australia	Effectiveness of acupuncture (traditional and P6) compared with sham (placebo) acupuncture or no acupuncture on NVP, RCT	I n = 148; comparator n = 148; sham n = 148; C n = 149	Women with symptoms of nausea or vomiting	RINVR score, I = nausea 8.3 (SD 2.5), dry retching 2.5, (SD 1.9) vomiting 2.3 (SD 2.7); C = nausea 8.2 (SD 2.6), dry retching 2.5 (SD 2.2), vomiting 2. (SD 2.8) (MILD-MODERATE)	(1) Serin (Japan) 0.2 x 30 mm needles were inserted to a depth 0.5-1.0 cm using a guide tube (maximum of six needles per session) then manipulated and left for a 20-minute period. Treatment was based on a traditional Chinese medicine diagnosis. Treatment points included stomach points 19, 20, 21; kidney points 21, 20; and conception vessel points 14, 13, 12, 11, or 10 plus points to treat the traditional Chinese medicine diagnosis	(2) comparator – women allocated to the P6 study group received acupuncture to this single point only for a 20-minute period (twice during the first week and then weekly for 3 further weeks); (3) sham – women allocated to the sham acupuncture group received acupuncture needles inserted into an area close to, but not on, acupuncture points over similar time points; (4) C – a standardised information sheet was made available about advice on diet, lifestyle and the use of vitamin B6 during the 4-week study period. Women in this group received a weekly 10-minute telephone call from the study investigator to assess their general sense of well-being and to encourage compliance with participating in the trial	RINVR	Nausea: (1) I – day 7: 5.0 (SD 3.0), day 14: 4.6 (SD 3.1), day 21: 3.8 (SD 3.1), day 26: 3.4 (SD 3.0); (2) comparator – day 7: 5.4 (SD 3.3), day 14: 4.8 (SD 3.6), day 21: 4.3 (SD 3.3), day 26: 4.0 (SD 3.3); (3) sham – day 7: 5.7 (SD 2.8), day 14: 5.0 (SD 3.0), day 21: 4.4 (SD 2.7), day 26: 3.7 (SD 2.8); (4) C – day 7: 6.1 (SD 2.9), day 14: 6.0 (SD 3.1), day 21: 5.8 (SD 3.1), day 26: 5.0 (SD 3.0) (p-values – day 7 p = 0.05, day 14 p = 0.01, day 21 p = 0.001, day 26 p = 0.001) Dry retching: (1) I – day 7: 1.3 (SD 1.4), day 14: 0.9 (SD 1.3), day 21: 0.9 (SD 1.4), day 26: 0.8 (SD 1.4); (2) comparator – day 7: 1.6 (SD 1.7), day 14: 1.3 (SD 1.5), day 21: 0.9 (SD 1.3), day 26: 0.9 (SD 1.3); (3) sham – day 7: 1.5 (SD 1.8), day 14: 1.3 (SD 1.7), day 21: 0.9 (SD 1.3), day 26: 0.9 (SD 1.4); (4) C – day 7: 1.7 (SD 1.7), day 14: 1.3 (SD 1.7), day 21: 1.6 (SD 1.7), day 26: 1.6 (SD 1.7) (p-values – day 7 p = ns, day 14 p = 0.01, day 21 p = 0.001, day 26 p = 0.001)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
									<p>Vomiting: (1) 1 – day 7: 1.4 (SD 2.0), day 14: 1.1 (SD 1.8), day 21: 0.9 (SD 1.6), day 26 0.9 (SD 1.5); (2) comparator – day 7: 1.2 (SD 2.0), day 14: 1.3 (SD 2.2), day 21: 1.2 (SD 2.1), day 26 = 0.9 (SD 1.8); (3) sham – day 7: 1.5 (SD 2.2), day 14: 1.4 (SD 2.1), day 21: 1.0 (SD 1.7), day 26 = 1.0 (SD 1.6); (4) C – day 7: 1.5 (SD 2.1), day 14: 1.6 (SD 2.2), day 21: 1.1 (SD 2.1), day 26 = 1.4 (SD 2.0). (<i>p</i>-values – all non-significant)</p>
Acupuncture vs. Chinese herbal medicine vs. Western medicine									
Mao 2009 ⁸⁷	Jiuquan Hospital of Traditional Chinese Medicine, Jiuquan, China	Effectiveness of acupuncture vs. Chinese herbal medicine vs. Western medicine, three-arm RCT	n = 90 (1 n = 30; comparator 1 n = 30; comparator 2 n = 30) (1) 1.8.30 (SD 1.60); (2) comparator 1 8.33 (SD 1.38); (3) comparator 2 8.57 (SD 1.66)	Women attending hospital clinic with vomiting and unable to tolerate oral intake	I mean number of vomiting episodes per 24 hours = 18.20 (8.54); comparator 1 mean number of vomiting episodes per 24 hours = 17.57 (7.06); comparator 2 mean number of vomiting episodes per 24 hours = 17.27 (8.50) (MODERATE-SEVERE)	All women given i.v. fluids to rehydrate and correct electrolyte imbalance. Traditional Chinese acupuncture twice daily for 7 days	Comparator 1, i.v. fluids given to rehydrate, phenobarbital given (30 mg) three times a day for 7 days Comparator 2, i.v. fluids given to rehydrate, traditional herbal remedy made daily and given in small doses through the day. Aim for three doses but if difficult to swallow then split into smaller more frequent doses	Author-defined scale (change in symptoms)	Symptom severity on day 4: (1) 1 complete recovery 12 (40%), obvious improvement 8 (26.7%), slight improvement 9 (30.0%), no effect 1 (3.3%); (2) comparator 1 complete recovery 2 (6.7%), obvious improvement 10 (10.0%), slight improvement 7 (23.3%), no effect 18 (60.0%); (3) comparator 2 complete recovery 3 (10.0%), obvious improvement 1 (3.3%), slight improvement 8 (26.7%), no effect 18 (60.0%)

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
									Day 4, proportion of women with symptom relief: (1) 1 96.7%; (2) comparator 1 40.0%; (3) comparator 2 40.0%
									Symptom severity on day 8: (1) 1 complete recovery 27 (90.0%), obvious improvement 2 (6.7%), slight improvement 0 (0%), no effect 1 (3.3%); (2) comparator 1 complete recovery 1 (3.3%), obvious improvement 6 (20.0%), slight improvement 4 (13.3%), no effect 16 (53.4%), (3) comparator 2 complete recovery 3 (10.0%), obvious improvement 10 (33.0%), slight improvement 5 (16.7%), no effect 12 (40%)
									Proportion of women gaining some benefit at day 8: (1) 1 96.7%; (2) comparator 1 46.6%; (3) comparator 2 60.0. %
									Acupuncture significantly better than other two groups, no significant differences between other two groups

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Zhang 2005 ¹⁵	Dongguan City, Zhangmutou Hospital, Guangdong, China	Effectiveness of Chinese acupuncture plus moxibustion vs. Chinese herbal medicine vs. Western medicine, three-arm RCT	n = 150 (I n = 50; comparator 1 n = 50; comparator 2 n = 50) At study entry: (1) I 6–8 weeks n = 30, 8–12 weeks n = 17, > 12 weeks n = 3; (2) comparator 1 6–8 weeks n = 28, 8–12 weeks n = 20, > 12 weeks n = 2; (3) comparator 3 6–8 weeks n = 25, 8–12 weeks n = 22, > 12 weeks n = 3	Women attending hospital with abnormal vomiting and unable to tolerate oral intake	Number of women per severity of vomiting: I < 5 times/day n = 15, 6–10 times/day n = 30, > 10 times/day n = 5; comparator 1 < 5 times/day n = 11, 6–10 times/day n = 27, > 10 times/day n = 12; comparator 2 < 5 times/day n = 13, 6–10 times/day n = 32, > 10 times/day n = 5 (MODERATE)	Traditional Chinese acupuncture at specific points plus gentle warming moxibustion for 10–15 minutes twice daily for 7 days. If the first round of treatment is ineffective a 3-day rest period was given before repeating the 7 days	Comparator 1: specific traditional formula of herbs boiled and made into drink. Full daily dose given in two parts over 7 days (repeated if not effective for further 7 days) Comparator 2: 2500–3000 mls i.v. fluids given daily to correct dehydration and electrolyte imbalance. Phenobarbital (30 mg) given orally three times daily for 7 days. If not effective treatment repeated for another 7 days after a 3-day rest period	Author-defined scale (change in symptoms)	(1) I acupuncture plus moxibustion complete recovery 21 (42%), obvious improvement 13 (26%), slight improvement 9 (18%), no effect 7 (14%); (2) comparator 1 complete recovery 9 (18%), obvious improvement 7 (14%), slight improvement 5 (10%), no effect 29 (58%); (3) comparator 2 complete recovery 5 (10%), obvious improvement 8 (16%), slight improvement 6 (12%), no effect 31 (62%)

continued

TABLE 9 Results for acupressure, nerve stimulation and acupuncture interventions NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Acupressure vs. no comparator									
Markose 2004 ^{1,24}	Christian Medical College Hospital, Vellore, India	To evaluate the effect of P6 acupressure in women who were given reassurance about the natural regression of nausea and who were not under any medication, case series	n = 35 recruited, n = 17 completed	Women experiencing nausea with or without vomiting	Percentage of women experiencing symptoms and distress: frequency of nausea 52.9%; frequency of vomiting 88.2%; frequency of dry retches 58.8%; distress due to nausea 58.8%; distress due to vomiting 82.3%; distress due to dry retches 70.5% (MILD-MODERATE)	Acupressure on point P6 from the fourth day (the first 3 days being the control phase), four times a day for 10 minutes on each hand	None, days 1-3 classed as a control period	RINVR	Frequency of nausea: day 1 52.9%, day 3 58.8%, day 7 11.7%; <i>p</i> = 0.008 Frequency of vomiting: day 1 88.2%, day 3 100%, day 7 17.6%; <i>p</i> < 0.001 Frequency of dry retches: day 1 58.8%, day 3 58.8%, day 7 5.8%; <i>p</i> = 0.004 Distress due to dry retches: day 1 70.5%, day 3 64.7%, day 7 23.5%; <i>p</i> = 0.016 Distress due to vomiting: day 1 82.3%, day 3 76.4%, day 7 29.4%; <i>p</i> = 0.008 Distress due to nausea: day 1 58.8%, day 3 70.5%, day 7 11.7%; <i>p</i> = 0.002

C, control; CV, Central Venter; I, intervention; IQR, interquartile range; MR, mean rank; NR, not reported; ns, not significant; SAS, sensory afferent stimulation; ST36, Stomach 36.

McGill Nausea Questionnaire

Only one trial used the McGill Nausea Questionnaire (a combined measure of various nausea severity indices) to assess symptom relief. Hsu and colleagues⁷⁹ reported no difference between intervention and sham comparator at baseline, 30 or 60 minutes for any of the indices ($p = 0.2$ for all).

Nausea outcomes

The trial of Bayreuther and colleagues⁶¹ used the 10-point VAS to assess the difference in mean levels of nausea severity between groups. They reported that the mean level of nausea was significantly lower for the intervention group (3.23 points on the VAS) than for the placebo group (4.92 points on the VAS) ($p = 0.019$).

Can Gurkan and Arslan⁴³ assessed the frequency, severity and discomfort from nausea using the VAS. They compared the difference in mean VAS scores between (1) the first 3 days and the second 3 days of the study; and (2) the second 3 days and the third 3 days for the each study group (intervention, placebo and a third no-treatment control arm). For frequency of nausea, a significant difference was reported between the first and second 3 days in both intervention ($z = -3.35$; $p < 0.001$) and placebo groups ($z = -0.28$; $p < 0.05$), but not in the control group ($z = -0.92$; $p > 0.05$). For frequency of nausea between the second and third 3 days of the study, no significant difference was found in any group (intervention $z = -1.28$, placebo $z = 10.18$, control $z = -1.40$; $p > 0.05$). For severity of nausea, the difference between the first and second 3 days was significant in the intervention and placebo group ($z = -0.04$ and $z = -2.96$, respectively; $p < 0.05$), but not in the control group ($z = -1.02$; $p > 0.05$). No significant difference was found in any group between the second and third 3 days (intervention $z = -1.48$; placebo $z = -1.26$, control $z = 0$; $p > 0.05$). The difference in the intensity of discomfort felt from nausea was significant for both the intervention group ($z = -3.7$; $p < 0.001$) and placebo group ($z = -2.4$; $p < 0.05$) but not control ($z = -1.2$; $p > 0.05$) between the first and second 3 days. However, the difference for any group for the second to third 3 was not significant (intervention $z = -0.22$; $p > 0.05$; placebo $z = -1.4$, $p > 0.05$; control $z = -0.44$; $p > 0.05$).

Naeimi Rad and colleagues⁹¹ also used the 10-point VAS to assess nausea severity between intervention (acupressure at point KID21) and placebo ('false' point) groups. Median VAS scores showed no significant difference between groups at day 1 [intervention 7 (IQR 8–6), comparator 7 (IQR 8–6); $p = 0.473$]. However, the difference was significant at day 2 [intervention 6 (IQR 7.75–4), comparator 7 (IQR 8–6); $p = 0.012$] and highly significant at days 3 [intervention 5 (IQR 5–3), comparator 7 (IQR 8–5); $p < 0.001$] and 4 [intervention 4 (IQR 5–2), comparator 7 (IQR 8–5); $p < 0.001$].

Werntoft and Dykes¹¹¹ assessed the difference in nausea severity using the 10-point VAS between an intervention group receiving acupressure at the P6 acupoint compared with either a placebo treatment group or a third control group receiving no treatment at all. Although significant differences were observed between both intervention and placebo group compared with control at days 1 and 3 ($p = 0.005$ and $p = 0.038$, respectively), no differences in favour of the intervention group compared with the placebo and control groups were found at days 6 and 14 of treatment ($p = 0.17$ and $p = 0.11$ respectively).

Belluomini and colleagues⁶² measured nausea via the RINVR between the pre-treatment period (mean of days 1–3) and the post-treatment period (mean of days 5–7). Both intervention and comparator groups reported a significant reduction in nausea scores [intervention from 8.38 ± 2.2 to 5.80 ± 2.9 ($p \leq 0.001$); comparator from 7.99 ± 2.5 to 7.04 ± 2.6 ($p \leq 0.001$)].

An author-defined assessment scale composed of number of episodes and self-reported severity of nausea was used in Steele and colleagues¹⁰⁴ to determine impact of acupressure at point P6 against placebo. During the 4 days of treatment, significant differences in favour of the intervention group were detected on both measures (mean rank for nausea frequency: intervention 40.30, comparator 72.82, $p < 0.001$; nausea severity: intervention 40.13, comparator 73.09, $p < 0.001$). However, when comparing the mean differences between the treatment period (days 1–4) and the post-treatment period (days 5–7), only the frequency of nausea showed a significant reduction ($p < 0.05$).

Vomiting outcomes

Emesis was measured with the RINVR in the trial of Belluomini and colleagues⁶² [change between the pre-treatment period (mean of days 1–3) and the post-treatment period (mean of days 5–7)]. A significant reduction was reported for the intervention group (from 2.09 ± 2.5 to 1.28 ± 1.9 ; $p = 0.03$). However, although the comparator group showed a reduction (from 1.83 ± 2.7 to 1.63 ± 2.3), it was not reported if this change was significant.

Naeimi Rad and colleagues⁹¹ used the 10-point VAS to compare vomiting frequency⁹¹ between intervention (acupressure at point KID21) and placebo ('false' point) groups. Increasingly significant differences between intervention and placebo groups were recorded on all days of treatment [day 1, intervention 1 (IQR 2–0), comparator 1 (IQR 2–1), $p = 0.012$; day 2, intervention 0 (IQR 1–0), comparator 1 (IQR 2–0.25), $p = 0.003$; day 3, intervention 0 (IQR 1–0), comparator 1 (IQR 2–0), $p = 0.001$; day 4, intervention 0 (IQR 0.75–0), comparator 1 (IQR 2–0), $p < 0.001$].

Can Gurkan and Arslan⁴³ assessed the difference in number of vomiting episodes between (1) the first and second 3 days of the study; and (2) the second and third 3 days for the each study group (intervention, placebo and a third no-treatment control arm).⁴³ No significant difference was observed at either time point in any of the three study groups [first 3 days compared with second 3 days: intervention $z = -1.38$, placebo $z = -1.85$ and control $z = -0.37$ ($p > 0.05$); second 3 days compared with third 3 days: intervention $z = -1.3$, placebo $z = -0.7$ and control $z = -0.57$ ($p > 0.05$)]. Bayreuther and colleagues⁶¹ recorded episodes of vomiting in both intervention and comparator groups but did not report those findings in their paper.

As above, Steele and colleagues¹⁰⁴ employed an author-defined assessment scale composed of number of episodes and self-reported severity of vomiting to measure the effectiveness of acupressure at point P6 against placebo. Significant differences favouring intervention were reported for both the frequency (intervention mean rank 41.51, comparator mean rank 70.94; $p < 0.001$) and severity (intervention mean rank 39.28, comparator mean rank 73.65; $p < 0.001$) of vomiting during the 4 intervention days. However, only the difference between groups for the severity of vomiting was significant when comparing days 1–4, with days 5–7 ($p < 0.05$).

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or adverse events were reported for any of the included trials. There were no relevant data on this intervention from the UKTIS.

Acupressure versus vitamin B6

One trial (Jamigorn and Phupong⁸⁰), judged as having a low risk of bias, compared the use of acupressure administered via a pressure band at acupoint P6 against vitamin B6 taken in tablet form.

Rhodes Index of Nausea, Vomiting and Retching

Jamigorn and Phupong⁸⁰ used the RINVR to measure changes in combined nausea, vomiting and retching scores. They reported significant improvements in overall scores in both the acupressure ($p < 0.001$) and vitamin B6 groups ($p < 0.001$) (see *Figure 2*). However, there were no statistically significant differences in the reduction of RINVR scores between baseline and end of treatment score between groups ($p > 0.05$).

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or adverse events were reported in any of the included trials. There were no relevant data on this intervention from the UKTIS.

Acupressure (case series)

Markose and colleagues¹²⁴ was a case series study that evaluated the effect of P6 acupressure on women with mild to moderate nausea following an initial control phase of 3 days without treatment (categorised as weak in terms of quality).

Rhodes Index of Nausea, Vomiting and Retching

The RINVR was used to assess symptoms, but the actual RINVR score was not presented. Results for individual components of score (nausea, vomiting and retching) are presented below.

Nausea outcomes

The RINVR was used to measure nausea outcomes. A significant reduction in the frequency of nausea (defined as percentage of women experiencing three or more adverse events) (day 1 52.9%, day 3 58.8%, day 7 11.7%; $p = 0.008$) and distress due to nausea (day 1 58.8%, day 3 70.5%, day 7 11.7%; $p = 0.002$) was reported.

Vomiting outcomes

Emesis outcomes were measured using the RINVR in terms of both the frequency of vomiting and the vomiting-related distress reported by the women themselves.¹²⁴ As with vomiting, the authors report a significant trend of reduced vomiting frequency by day 7 (defined as percentage of women reporting three or more events) (day 1 88.2%, day 3 100%, day 7 17.6%; $p < 0.001$). A similar positive trend was reported for distress experienced as a result of vomiting (defined as percentage of women self-reporting moderate to severe distress) (day 1 82.3%, day 3 76.4%, day 7 29.4%; $p = 0.008$).

Retching outcomes

Finally, Markose and colleagues¹²⁴ also measured the frequency of, and distress caused by, dry retches using the RINVR. A significant reduction in both retching outcomes (defined as percentage of women reporting three or more events, and distress from moderate to severe) was reported (retching frequency: day 1 58.8%, day 3 58.8%, day 7 5.8%, $p = 0.004$; distress due to dry retches: day 1 70.5%, day 3 64.7%, day 7 23.5%, $p = 0.016$).

Safety outcomes

No pregnancy outcomes or adverse events were reported in any of the included trials. There were no relevant data available on these interventions from the UKTIS.

Nerve stimulation versus placebo

Three studies compared the effect of a nerve stimulation device administered at pressure point P6 (as described in *Acupressure versus placebo*) against an identical inactive unit.^{73,98,109} Of these trials, two provided insufficient data to permit a clear judgement of bias,^{73,109} and the other trial was judged as having a low risk of bias.⁹⁸ Given the differences between trials in patient populations, settings, interventions and

in particular the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses and have outlined individual trial results in a narrative.

Rhodes Index of Nausea, Vomiting and Retching

Rosen and colleagues⁹⁸ used the RINVR to assess overall change in symptom severity. They determined that the time-averaged change in the RINVR score for total experience was significantly better in the study group than in the control group [6.48 (95% CI 5.31 to 7.66) vs. 4.65 (95% CI 3.67 to 5.63); $p = 0.02$]. Veciana and colleagues¹⁰⁹ also used the RINVR to detect the difference symptom severity scores between the intervention and control groups. However, they found no significant difference in RINVR score between groups (p -value not reported).

Author-defined scale

Evans and colleagues⁷³ measured subjective change in nausea and vomiting between those wearing active versus inactive nerve stimulation units via an author-defined scale (improved, worsened, no change). A significant change favouring the intervention group was detected in terms of both average nausea score (active unit 2.4, placebo 2.7; $p < 0.05$) and improvement in nausea score (active unit 15, placebo 10; $p < 0.05$).

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

The trial of Rosen and colleagues⁹⁸ did not report pregnancy outcomes, but three dehydration events were reported in the study group compared with 12 events in the control group ($p = 0.013$).⁹⁸ Neither of the other two trials reported pregnancy outcomes or side effects,^{73,109} and there were no UKTIS data available on this intervention.

Acupuncture versus placebo

The trials of Carlsson and colleagues⁶⁶ and Knight and colleagues⁸³ both compared acupuncture at various treatment points with placebo (sham) treatment. Smith and colleagues¹⁰¹ was a four-arm RCT comparing a combination of various treatment points (group 1); with comparators including single point acupuncture to P6 (group 2); sham acupuncture to ineffective points (group 3); and a control comprising a standard information sheet and telephone support (group 4). Smith and colleagues¹⁰¹ and Knight and colleagues⁸³ were both judged as carrying a low risk of bias; however, Carlsson and colleagues⁶⁶ was judged as high due to lack of blinding.

Combined severity score

No combined score reported.

Nausea outcomes

Carlsson and colleagues⁶⁶ assessed nausea severity between day 0 and day 1, and day 4 and day 5, in their crossover trial with the 10-point VAS. Focussing on 'speed' of change, the VAS reductions were calculated as the difference between the VAS estimates the day before acupuncture and those the day following the 2 acupuncture days for each patient. This intergroup crossover analysis showed a significantly faster reduction of nausea when giving the patients active acupuncture than when giving them placebo

acupuncture ($p = 0.032$). However, they found no time effect ($p = 0.138$). The VAS was also used by Knight and colleagues⁸³ to measure changes in nausea outcomes. They found evidence of a time effect ($p < 0.001$) in all groups [intervention scores reduced from 85.50 (IQR 71.25–89.75) on day 1 to 47.5 (IQR 29.25–69.5) at 3 days after session four; comparator scores reduced from 87.0 (IQR 73.0–90.0) on day 1 to 48.0 (IQR 14.0–80.0) at 3 days after session four. However, the authors reported that they found no evidence of a group effect ($p = 0.9$) or a group–time interaction ($p = 0.8$).

Smith and colleagues¹⁰¹ measured nausea severity with the RINVR. They found that women in the traditional acupuncture group were more likely to be free from nausea compared with women in the no acupuncture control group (relative risk 0.93, 95% CI 0.88 to 0.99) at the end of their first week of treatment. During the second week of the trial, women who received traditional acupuncture ($p < 0.001$), and P6 acupuncture ($p < 0.05$) reported lower nausea scores compared with women in the no acupuncture control group. This improvement in nausea continued for women receiving traditional acupuncture ($p < 0.001$) and P6 acupuncture ($p < 0.01$) into the third week compared with women in the no acupuncture control group. From the third week, women in the sham acupuncture group also reported lower nausea scores compared with women in the no acupuncture control group ($p < 0.01$). In the final week of the study, improvements in nausea continued for women in the traditional acupuncture (reported in the paper as $p < 0.001$, but actual value is not clear), P6 acupuncture ($p < 0.05$) and sham acupuncture ($p < 0.01$) groups compared with women in the no acupuncture group.

Vomiting outcomes

In the crossover trial of Carlsson and colleagues,⁶⁶ occurrence of vomiting (i.e. emesis vs. no emesis) was compared between study groups. For active acupuncture treatment, they reported that 7 out of 17 women were still vomiting after 2 acupuncture days. For the placebo group, 12 out of 16 women were still vomiting at the same time point. They did not report if this difference was significant. Knight and colleagues⁸³ measured number of vomiting episodes, but did not report these data.

Smith and colleagues¹⁰¹ measured vomiting severity with the RINVR. Lowered vomiting scores were recorded in all groups [intervention: day 7, mean 1.4 (SD 2.0) to day 26, mean 0.9 (SD 1.5); single point acupuncture: day 7, mean 1.2 (SD 2.0) to day 26, mean 0.9 (SD 1.8); placebo: day 7, mean 1.5 (SD 2.2) to day 26, mean 1.0 (SD 1.6)], including a small reduction in the control arm [day 7, mean 1.5 (SD 2.1) to day 26, mean 1.4 (SD 2.0)]. However, the differences between groups were not significant.

Retching outcomes

Smith and colleagues¹⁰¹ measured retching severity with the RINVR. They found that women in the traditional acupuncture ($p < 0.001$), P6 acupuncture ($p < 0.01$) and sham acupuncture ($p < 0.001$) groups all experienced fewer periods of dry retching compared with women in the no acupuncture control group at the end of the study. Sixty-eight (56%) women in the traditional acupuncture group were free from dry retching compared with 46 (39%) women in the no acupuncture control group by the end of the third week (relative risk 0.72, 95% CI 0.56 to 0.93; $p < 0.01$; number needed to treat 6, 95% CI 3 to 22). In the sham acupuncture group, 72 (59%) women were free from dry retching compared with 46 (39%) women in the no acupuncture control group (relative risk 0.68, 95% CI 0.52 to 0.87; $p < 0.001$; number needed to treat 6, 95% CI 3 to 13). These improvements continued to the end of the trial. In the P6 acupuncture group, no difference occurred in women free from dry retching compared with women in the no acupuncture control group.

Safety outcomes

No pregnancy outcomes or adverse events were reported in any of the included trials. There was no relevant data on this intervention from the UKTIS.

Acupuncture versus metoclopramide

The trial of Neri and colleagues⁹⁴ compared acupuncture to acupoints PC6, Central Venter 12 and Stomach 36 plus the wearing of an acupressure device at the PC6 point against the administration of a metoclopramide infusion at hospital plus oral supplementation with a vitamin B12 complex. The trial was judged as carrying a high risk of bias due to selective outcome reporting.

Combined severity score

No combined score reported.

Nausea outcomes

Neri and colleagues⁹⁴ measured the number of cases self-reporting improvements in the intensity of nausea (scored as 0 = no nausea; 1 = low intensity, no discomfort; 2 = high intensity, discomfort). Improvements were reported for both interventions [*n* (%) per session] [acupuncture: first session 1 (2.3%); second session 11 (25.5%); third session 19 (44.1%); metoclopramide: first session 1 (2.3%); second session 9 (23.6%); third session 12 (31.5%)], and the difference between groups was not reported as being non-significant (first session $p = 0.3$; second session $p = 0.6$; third session $p = 0.2$).

Vomiting outcomes

Neri and colleagues⁹⁴ measured improvements in the number of cases reporting improvements in vomiting episodes (grouped as 0 = no episodes, 1 = 1–3/day, 2 = > 3/day). Both groups showed improvements [*n* (%) per session] [acupuncture: 7 (16.2%) for the first session; 15 (34.8%) for the second session; and 24 (55.8%) for the third session; metoclopramide: 4 (10.5%) for the first session; 12 (31.5%) for the second session; 14 (36.8%) for the third session]. However, the difference between groups was not found to be significant ($p = 0.4$ for first session and $p = 0.5$ for the second), although it approached significance after the third treatment session, in favour of acupuncture ($p = 0.07$).

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or adverse events were reported in by Neri and colleagues.⁹⁴ There were no relevant data on this intervention from the UKTIS.

Acupuncture versus Chinese herbal medicine versus barbiturates

Two studies compared acupuncture against both Chinese herbal medicine and Western medicine. In the Mao and Liang trial,⁸⁷ all women were given i.v. fluids for rehydration purposes and to correct electrolyte imbalance. The following interventions were then compared: (1) traditional Chinese acupuncture at acupoints BL11, ST37, PC6 and SP4; (2) a traditional herbal remedy; and (3) phenobarbital. In the Zhang 2005 trial,¹¹⁵ traditional Chinese acupuncture was compared with a traditional herbal drink and i.v. fluids plus phenobarbital. Both trials were found to have a high risk of bias due to inadequate blinding and/or selective outcome reporting.

Author-defined scale

Mao and Liang⁸⁷ assessed symptom severity using an author-defined scale (complete recovery, obvious improvement, slight improvement, no effect) on days 4 and 8, alongside the proportion of women with symptom relief in each group. They reported that the acupuncture performed significantly better than comparator intervention groups (with no significant difference between those other two groups). However, they did not report *p*-values. In the trial of Zhang,¹¹⁵ a similar author-defined scale was also used to assess symptom relief. It reported the highest proportion of complete recovery (21 patients, 42%) among the

acupuncture group [compared with herbal medicine 9 (18%) or Western medicine 5 (10%)], but did not report if this difference was significant. The lowest numbers of patients reporting no effect were also found in the acupuncture group compared with either the herbal remedy or Western medicine groups [7 (14%) vs. 29 (58%) vs. 31 (62%) respectively]. However, again, it was not reported if this difference was significant.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or adverse events were reported in the included trials. There were no relevant data on this intervention from the UKTIS.

Summary

- The quality of evidence available for this group of interventions varied considerably, from low (eight studies^{61,62,66,80,83,91,98,101}) to high (five studies^{87,94,111,115,124}) risk of bias.
- Comparisons with placebo were equivocal: two studies (both where participants had mild symptoms and at low risk of bias)^{61,62} found no evidence of a difference or did not report the outcome for nausea and vomiting symptoms respectively.
- Acupressure at the KID21 acupoint reduced both nausea and vomiting compared with placebo.
- No evidence of a difference between acupressure and vitamin B6.⁸⁰
- Two studies^{73,98} of nerve stimulation at the P6 acupressure point reported better outcomes versus a non-stimulating device [one study = participants with mild–moderate severity with low risk of bias,⁹⁸ one study = unclear severity and risk of bias⁷³]. A third study found no evidence of a difference (unclear severity and risk of bias).¹⁰⁹
- Three studies^{66,83,101} compared acupuncture with placebo. There was no evidence of an effect on vomiting but one study (mild–moderate symptoms, low risk of bias)¹⁰¹ reported reduced nausea and retching in both traditional and P6 acupuncture groups.
- No evidence of a difference between acupuncture compared with metoclopramide, but symptoms improved from baseline in both groups (one study: high risk of bias, severity assessed as moderate–severe⁹⁴).
- Comparisons of traditional Chinese acupuncture and herbal medicine with Western medicine were at high risk of bias and impossible to emulate within the UK NHS setting.
- Overall conclusions are limited by the quality of included studies. Many participants appear to have improved symptoms from baseline in all treatment arms.
- There is some suggestion that acupressure may have some beneficial effect in women with mild symptoms of NVP.
- Acupressure at the KID21 point appears to improve symptoms of nausea and vomiting compared with acupressure to a false point, but data are limited.
- The evidence for nerve stimulation is mixed.
- Acupuncture may reduce symptoms of nausea and retching in women with mild–moderate symptoms, but data are limited and inconclusive.
- More larger, better-quality studies are required for these interventions.

Chapter 6 Clinical effectiveness: aromatherapy

Introduction

Aromatherapy was used as an intervention to treat nausea and/or vomiting in two pilot RCTs.^{76,97} Heterogeneity was observed in relation to the clinical setting and patient populations in which the studies were conducted, as well as the interventions, comparators and outcomes reported in each trial. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*) given these, we did not attempt to perform meta-analyses, and have thus reported a narrative summary only for each intervention and comparator set.

One trial compared aromatherapy with placebo in pre-natal wards⁹⁷ whereas the other trial compared aromatherapy with routine antenatal care.⁷⁶ Both trials were at unclear risk of bias: mainly due to poor reporting in the trial of Pasha and colleagues,⁹⁷ and for uncertainty surrounding whether the trial was truly randomised or was a quasi-RCT in Ghani and Ibrahim.⁷⁶ The women were described as experiencing symptoms at the mild to moderate end of the severity spectrum at baseline in both trials (*Table 10*).

Aromatherapy versus no aromatherapy

One trial⁹⁷ compared mint oil aromatherapy against placebo for the treatment of mild NVP.⁹⁷ Another trial compared mixed essential oils (lavender and peppermint) and routine antenatal care, where no treatment oils were given, in women with mild to moderate nausea and/or vomiting (i.e. requiring antiemetics but not hospitalisation).

Rhodes Index of Nausea, Vomiting and Retching

In the trial of Ghani and Ibrahim,⁷⁶ the observed difference in the overall change from baseline RINVR score in the aromatherapy group was reduced to 17.60 (SD 6.08) from 23.06 (SD 6.37) at the end of the 3-day trial ($p < 0.001$). The change score in the no aromatherapy control group was not reported so it is unclear if the aromatherapy was effective in reducing nausea and vomiting or if it was due to the time lag.

Nausea outcomes

Pasha and colleagues⁹⁷ examined the effectiveness of mint aromatherapy on pregnancy-induced nausea intensity using a VAS. There was no significant difference between the two groups. However, the authors reported a trend towards improvement by day 4 in the mint group, where the score had reduced to 3.50 (SD 1.95) from 4.78 (SD 1.62) at baseline, and an increase in nausea score at the end of day 4 from baseline score in the placebo group [mean 4.38 (SD 2.18) at day 4 compared with mean 3.00 (SD 2.19) at baseline].

Vomiting outcomes

The trial of Pasha and colleagues⁹⁷ also examined the effect of mint aromatherapy on the number of vomiting episodes.⁹⁷ There was no significant difference between the two groups. However, there was a reported trend towards improvement by day 4 in the mint group where the mean number of episodes had reduced to 2.23 (SD 1.88) from 4.85 (SD 1.82) at baseline. In the placebo group there was no significant difference between the number of vomiting episodes at the end of day 4 and baseline [mean 2.55 (SD 2.55) at day 4 compared with mean 2.52 (SD 2.4) at baseline].

Retching outcomes

No independent retching outcomes reported.

TABLE 10 Results for aromatherapy interventions for NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Pasha 2012 ⁹⁷	Pre-natal wards of seven health clinics, Iran	To evaluate the effect of mint oil on nausea and vomiting during pregnancy, pilot RCT	n = 67; I = 9.07 (SD 1.31), C = 9.73 (SD 2.2)	Women complaining of nausea and vomiting	Mean of nausea and vomiting intensity was: I = 4.78 ± 1.62, C = 3.00 ± 2.19 and I = 4.85 ± 1.82, C = 2.52 ± 2.4, 7 days before treatment (MILD)	Assigned to use a bowl of water with four drops of pure mint oil placed on the floor near their beds for 4 consecutive nights before sleeping. Dietary advice also given (n = 33)	As I except given saline instead of mint oil (n = 34)	VAS for nausea Episodes of vomiting	Trend towards improvement by day 4 in mint group Mint group, mean scores over the 4 days: nausea 3.50 (SD 1.95), vomiting 2.23 (SD 1.88) Placebo, mean scores over the 4 days: nausea 4.38 (SD 2.18), vomiting 2.55 (SD 2.55) Not significant
Ghani, 2013 ⁷⁶	Outpatient department, maternity teaching hospital plus three maternal and child health centres in Abha Kingdom, Saudi Arabia	To estimate the effect of mixed essential oils inhalation on nausea and vomiting in early pregnancy, pilot RCT	n = 101; I = 10 (SD 2.61), C = 10.2 (SD 2.29)	Women who had nausea and/or vomiting and required antiemetics but were not hospitalised	RINVR score: I = 23.06 (SD 6.37), C = 21.74 (SD 7.30) (MILD-MODERATE)	Four drops of lavender oils plus one drop peppermint oil in one spoon of water (ratio 4 : 1 : 1) heated using an oil burner. Perform twice a day for 3 days before sleep. Women were instructed to breathe deeply for 20 minutes. No other medication taken, diet/lifestyle advice given (n = 50)	Routine care, no treatment/oils given, no medication taken, diet/lifestyle advice given (n = 51)	RINVR score	Treatment group: change in score from 23.06 (SD 6.37) to 17.60 (SD 6.08); p < 0.001 Results from the C group not reported

C, control; I, intervention.

Safety outcomes

No pregnancy outcomes or adverse events were reported in either of the included trials. There were no relevant data on this intervention from the UKTIS.

Summary

- The evidence from the two trials^{76,97} available for aromatherapy was predominantly at an unclear risk of bias.
- The identified studies reported evidence of an improvement in symptoms over time, but there was no evidence of a difference compared with placebo or routine antenatal care.
- Overall, few data are available with no evidence of an effect.
- More larger, better-quality studies are required for to make any conclusions about this intervention.

Chapter 7 Clinical effectiveness: vitamin B6 (pyridoxine)

Introduction

Five trials compared the effectiveness of pyridoxine (vitamin B6) for the treatment of NVP.^{41,59,100,107,112} Two trials compared doses of vitamin B6 against placebo tablets,^{41,100} and they were at low⁴¹ and unclear¹⁰⁰ risk of bias, respectively. Tan and colleagues¹⁰⁷ examined the effect of vitamin B6 and metoclopramide combination versus metoclopramide alone and was at an unclear risk of bias, mainly due to the use of a dubious placebo (Tic Tac®, Ferrero UK Ltd, Greenford, UK). The trial of Wibowo and colleagues¹¹² compared high and low doses of vitamin B6, and was at low risk of bias. Babaei and Foghaha⁵⁹ compared the effectiveness of vitamin B6 against dimenhydrinate in the treatment of NVP and was at an unclear risk of bias.

In all five studies,^{41,59,100,107,112} women were described as experiencing mild to moderate symptoms at baseline (Table 11). However, as previously described (see Chapter 3, *Meta-analysis of included randomised controlled trials*), given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses. Thus we reported a narrative summary only for each intervention and comparator set.

Vitamin B6 versus placebo

Pregnancy-Unique Quantification of Emesis and Nausea scale

Wibowo and colleagues¹¹² measured the change in overall symptom severity using the PUQE scale for lower versus higher doses of vitamin B6. They found a higher mean change for the high dosage compared with lower, and there was a statistically significant improvement in PUQE score with the higher dose.

Nausea outcomes

The trial of Vutyavanich and colleagues⁴¹ used the 10-point VAS to assess mean change in nausea symptoms.⁴¹ The study determined a higher mean change in the pyridoxine group [2.9 (SD 2.2) vs. placebo 2.0 (SD 2.7)], and this difference was significant ($p < 0.001$). The Sahakian and colleagues¹⁰⁰ trial also employed the VAS to assess mean change in nausea. Overall, the study did not detect a significant difference between groups. However, for patients experiencing severe nausea, a significant mean change was observed in favour of the pyridoxine group ($p < 0.001$).

Vomiting outcomes

The change in the number of vomiting episodes was measured by Vutyavanich and colleagues.⁴¹ However, although a greater reduction was observed in the pyridoxine group, this was not significant ($p = 0.055$). Sahakian and colleagues¹⁰⁰ also assessed vomiting outcomes via number of vomiting episodes.¹⁰⁰ They observed a significant improvement both in the pyridoxine group as a whole ($p < 0.05$), and for the subgroup of women experiencing severe symptoms ($p < 0.055$).

Retching outcomes

No independent retching outcomes reported.

TABLE 11 Results for vitamin B6 interventions for NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Vitamin B6 vs. placebo									
Vutyavanich 1995 ⁴¹	Antenatal clinic, Maharaj Nakorn Chiang Mai Hospital, Chiang Mai University, Thailand	To determine the effectiveness of pyridoxine for NVP, double-blind RCT	n = 342; I 10.9 (± 2.7), C 10.9 (± 2.8)	Women with nausea, with or without vomiting	Nausea and vomiting pregnancy instrument: I 5.2 (± 5.3), C 4.9 (± 2.4) Episodes of vomiting in past 24 hours: I 1.8 ± 2.3, C 1.6 ± 2.0 (MILD-MODERATE)	Vitamin B6: 10-mg tablets of pyridoxine hydrochloride 8-hourly for 5 days. Dietary advice given, advised not to take any other medication (n = 173)	Placebo: identical tablets 8-hourly for 5 days. Dietary advice given, advised not to take any other medication (n = 169)	VAS for nausea Episodes of vomiting	Nausea mean change in score: pyridoxine 2.9 (SD 2.2) vs. placebo 2.0 (SD 2.7) (p < 0.001) Vomiting episodes change: pyridoxine 1.22 (SD 2.0) vs. placebo 0.65 (SD 2.4) (p = 0.055)
Sahakian 1991 ¹⁰⁰	General obstetric clinic, Iowa, IA, USA	To determine the efficacy of pyridoxine for the treatment of NVP, double-blind RCT	n = 74; I 9.3 (range 6–15.5), C 9.7 (range 6–19)	Women suffering from nausea and vomiting but not requiring hospitalisation	VAS for nausea: I 6.4 (SD 1.8), C 6.6 (SD 1.9) (MILD-MODERATE)	Vitamin B6 25-mg tablets of pyridoxine 8-hourly for 3 days. Dietary advice given (n = 31)	Placebo: identical tablets 8-hourly for 3 days. Dietary advice given (n = 28)	VAS for nausea Episodes of vomiting	Nausea mean change: pyridoxine 2.9 (SD 2.4), placebo 1.9 (SD 2.0), p-value non-significant Severe nausea mean change: pyridoxine 4.3 (SD 2.1), placebo 1.8 (SD 2.2); p < 0.01 Vomiting: improvement in total pyridoxine group (p < 0.05) and severe symptoms (p < 0.05). Crude OR for vomiting vs. no vomiting in pyridoxine vs. placebo = 0.3014 (95% CI 0.102 to 0.893); p < 0.05

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Wibowo 2012 ^{11,12}	Cipto Mangunkusumo National Hospital, University of Indonesia, Indonesia	To determine if supplementation with vitamin B6 improves NVP (and to compare circulating plasma vitamin B6 between sufferers and non-sufferers), RCT	60 sufferers (+60 non-sufferers who did not receive any treatment but acted as comparison for plasma B6 concentrations) C: < 8 weeks' gestation (n = 12); 8–12 weeks' gestation (n = 18) I: < 8 weeks' gestation (n = 8); 8–12 weeks' gestation (n = 22)	Women suffering nausea and vomiting	PUQE: 1.7 (range 4–15), C 6 (range 4–11) (MILD)	Pyridoxine (5 mg) mixed with 40 g of powdered milk twice a day for 2 weeks. Dose of pyridoxine was equivalent to 10 mg per day. Women were asked not to take other medications	As I but equivalent dose of pyridoxine (0.64 mg) twice daily for 2 weeks (= 1.28 mg per day). Women were asked not to take any other medications (n = 30)	PUQE score	Change in score: high-dose vitamin B6 3.86 ± 2.12 vs. low-dose vitamin B6 2.80 ± 1.78; p < 0.05
Tan 2009 ⁰⁷	University of Malaya Medical Centre, Malaysia	To evaluate the oral use of pyridoxine in conjunction with standard therapy in women hospitalised for HG, RCT	n = 94; I 10.5 ± 3.1, C 9.6 ± 2.8	Women with severe nausea and vomiting during pregnancy with clinical features	Nausea score via VAS: I median 7 (IQR 5), C median 7 (IQR 4) (MODERATE)	i.v. rehydration with saline (± potassium chloride), pyridoxine two 10 mg-tablets given plus i.v. metoclopramide (10 mg) given three times daily plus oral thiamine (10 mg) daily. Discharged with pyridoxine, two tablets three times daily plus oral metoclopramide and thiamine (n = 48)	As I group, except instead of pyridoxine two mint Tic Tac® (Ferreo UK Ltd, Greenford, UK) given (n = 46)	VAS for nausea Episodes of vomiting	Nausea score after 2 weeks: I median 2 (IQR 3), usual care median 2.5 (IQR 4); p = 0.69 Vomiting after 2 weeks: I mean 1.4 (SD 1.3), usual care mean 1.4 (SD 1.6); p = 0.98

continued

TABLE 11 Results for vitamin B6 interventions for NVP (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Vitamin B6 vs. serotonin antagonists									
Babaei 2014 ⁵⁹	Two antenatal clinics of referral hospitals affiliated to Shiraz University of Medical Sciences, Shiraz, Southern Iran	To compare effectiveness of vitamin B6 and dimenhydrinate in the treatment of NVP, double-blind RCT	140 participants, gestation < 16 weeks	Not reported (exclusion criteria included hospitalisation for severe vomiting)	RINVR nausea vomiting scores mean (SD) at baseline were 8.6 (SD 2.9) for vitamin B6 (group A) and 8.3 (SD 7.4) for dimenhydrinate (group B) (MILD)	Vitamin B6 tablet (50 mg) orally every morning for 1 week (n = 70). Both vitamin B6 and dimenhydrinate tablets were identical in size, colour and odour	Dimenhydrinate tablet (50 mg) orally every morning for 1 week (n = 70)	RINVR	Both groups decreased nausea and vomiting scores from baseline values Average score change in the vitamin B6 group was less than that in the dimenhydrinate group [mean 4.4 (SD 1.6) vs. mean 5.7 (SD 5.5); $p < 0.05$] Each day, there were significant differences in score change of vitamin B6 group vs. dimenhydrinate group

C, control; I, intervention.

Safety outcomes

None of the included trials reported on pregnancy outcomes or adverse events. UKTIS data on vitamin B6 enquiries are provided in *Appendix 7*.

Vitamin B6 and metoclopramide combination versus metoclopramide alone

The trial of Tan and colleagues¹⁰⁷ used the 10-point VAS to assess median nausea scores in women with severe nausea and vomiting during pregnancy with clinical features. The trial also reported the mean number of daily vomiting episodes.¹⁰⁷

Combined severity score

No combined score reported.

Nausea outcomes

Tan and colleagues¹⁰⁷ reported no significant difference in nausea score after 2 weeks between the pyridoxine (vitamin B6) and metoclopramide as a combination treatment and metoclopramide alone [combination median 2 (IQR 3), metoclopramide alone median 2.5 (IQR 4); $p = 0.69$].

Vomiting outcomes

Tan and colleagues¹⁰⁷ reported no significant difference in the mean number of daily vomiting episodes after 2 weeks between the vitamin B6 and metoclopramide as a combination treatment and metoclopramide alone [combination mean 1.4 (SD 1.3), metoclopramide alone mean 1.4 (SD 1.6); $p = 0.98$].

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

Tan and colleagues¹⁰⁷ did not report on pregnancy outcomes or adverse events. UKTIS data on vitamin B6 enquiries are provided in *Appendix 7*.

Vitamin B6 versus serotonin antagonist

The trial of Babaei and Foghaha⁵⁹ compared effectiveness of vitamin B6 against dimenhydrinate in the treatment of NVP in a double-blind RCT which was adjudged as being at unclear risk of bias.

Rhodes Index of Nausea, Vomiting and Retching

Babaei and Foghaha⁵⁹ compared nausea and vomiting scores at baseline and post treatment using the RINVR. Results showed that both vitamin B6 and dimenhydrinate groups decreased nausea and vomiting scores from baseline. However, the average score change in the vitamin B6 group was less than that in the dimenhydrinate group [mean 4.4 (SD 1.6) vs. mean 5.7 (SD 5.5); $p < 0.05$].

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

Babaei and Foghaha⁵⁹ did not report on pregnancy outcomes. Occurrence of drowsiness was significantly lower in the vitamin B6 group compared with the dimenhydrinate group [5 (4.5%) as opposed to 36 (53%); $p < 0.01$]. No other adverse effect was observed in either group during the 1-week follow-up. UKTIS data on vitamin B6 enquiries are provided in *Appendix 7*.

Summary

- The evidence available for vitamin B6 was predominantly at low risk of bias or the risk of bias was unclear.
- Participants in the five studies^{41,59,100,107,112} had symptoms categorised as mild to moderate at baseline.
- Comparisons of vitamin B6 preparations with placebo generally reported evidence of a reduction in symptoms of nausea, especially in women with more severe symptoms, and vomiting.
- Higher doses of vitamin B6 preparations resulted in a greater improvement in NVP symptoms.
- There was no evidence to suggest that vitamin B6 and metoclopramide as a combination treatment had an advantage over metoclopramide alone.
- Overall, there is a suggestion that vitamin B6 might be better than placebo in reducing the severity of symptoms especially at higher doses, but more studies are required using a range of comparators.

Chapter 8 Clinical effectiveness: pyridoxine/doxylamine combination

Introduction

Four trials^{65,84,95,117} assessed the effectiveness of vitamin B6 in combination with doxylamine for the treatment of NVP in comparison with a placebo (Koren and colleagues⁸⁴), ondansetron (Capp and colleagues⁶⁵ and Oliveira and colleagues⁹⁵) or when administered pre-emptively versus following symptom onset (Maltepe and Koren¹¹⁷). In the Koren and colleagues⁸⁴ and Maltepe and Koren¹¹⁷ studies this took the form of Diclectin (delayed-release doxylamine/pyridoxine combination), while Oliveira and colleagues⁹⁵ and Capp and colleagues⁶⁵ gave pyridoxine and doxylamine as separate preparations.

One of the trials⁹⁵ was only available in abstract form and risk of bias was unclear, whereas the trial of Koren and colleagues⁸⁴ was at a low risk of bias. Another trial¹¹⁷ examined pre-emptive treatment with Diclectin and this trial appeared to be at low risk of bias but some items were unclear. Capp and colleagues⁶⁵ was at unclear risk of bias due to lack of provided details across a number of measures.⁶⁵ The reporting of severity of symptoms was not possible for the Maltepe and Koren¹¹⁷ pre-emptive study, poor for the Oliveira and colleagues⁹⁵ study and unclear for the Capp and colleagues⁶⁵ study, but in the Koren and colleagues⁸⁴ study, severity appeared to range from mild to moderate. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*), given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses, and have thus reported a narrative summary only for each intervention and comparator set.

We also identified a non-randomised prospective observational study (Ashkenazi-Hoffnung and colleagues³⁶) which compared a combination regimen of doxylamine and pyridoxine versus metoclopramide only (control group). This study is prone to extreme selection bias and there was a noticeable difference at baseline: moderate to severe symptoms were present in 97% of the intervention group women versus 69% of control group women ($p < 0.01$). A summary of study conduct and results is depicted in *Table 12* for completeness.

Doxylamine/pyridoxine versus placebo

One double-blind, multicentre placebo-controlled trial⁸⁴ compared treatment with Diclectin, a combination preparation of doxylamine succinate (10 mg) and pyridoxine hydrochloride (10 mg), to placebo in women with symptoms of NVP and a PUQE score ≥ 7 . Women in the trial had not previously responded to conservative management.

Pregnancy-Unique Quantification of Emesis and Nausea scale

Diclectin led to significantly greater improvement in NVP symptoms as compared with placebo (PUQE score: -4.8 ± 2.7 vs. 3.9 ± 2.6 ; $p = 0.006$). The mean area under the curve of the change in PUQE from baseline as measured day-by-day was significantly larger with Diclectin compared with placebo (61.5 ± 36.9 in the Diclectin group vs. 53.5 ± 37.5 in the placebo group; $p < 0.001$).

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

TABLE 12 Results for pyridoxine-doxylamine interventions for NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Pyridoxine/doxylamine vs. placebo									
Koren 2010 ³⁴	Three university medical centres: the University of Texas, Texas, TX, USA; University of Pittsburgh, Pittsburgh, PA, USA; and Georgetown University, Washington, DC, USA	The effectiveness of Diclectin [doxylamine succinate (10 mg), pyridoxine hydrochloride (10 mg), delayed-release preparation] vs. placebo for NVP, double-blind RCT	n = 280; I 9.3 ± 2.0, C 9.3 ± 1.8	Women suffering from NVP with a PUQE score of ≥ 6 and had not responded to conservative management (dietary/lifestyle advice)	Mean PUQE: I 9.0 ± 2.1, C 8.8 ± 2.1 Median PUQE: I 9.0, C 8.0 (MODERATE)	Diclectin, two tablets at night, increasing up to four tablets daily as needed. Follow-up clinic visits and telephone calls (n = 140)	Identical placebo tablet: two tablets at night, increasing up to four tablets daily as needed. Follow-up clinic visits and telephone calls (n = 140)	PUQE score	Greater improvement in mean score change in Diclectin group vs. placebo (PUQE score: -4.8 ± 2.7 vs. 3.9 ± 2.6; p = 0.006) The mean area under the curve of the change in PUQE score from baseline was significantly larger with Diclectin as compared with placebo (61.5 ± 36.9 vs. 53.5 ± 37.5; p < 0.001)
Maltepe 2013 ¹⁷	Women recruited who were calling the Motherisk NVP Helpline for counselling, Canada	Does pre-emptive treatment before the symptoms begin vs. when the symptoms begin improve symptoms in patients at a high risk for recurrence of severe NVP, RCT	n = 60, NR	N/A (recruited onto study before symptoms stated)	Not applicable – see Severity Inclusion criteria	Diclectin, two tablets at bedtime following pregnancy confirmation, gradual increase if symptoms escalate. Frequent additional counselling provided. Motherisk algorithm of treatment followed (n = 31)	Diclectin, two tablets at bedtime on first day of NVP symptoms with gradual increase if symptoms escalated. Frequent additional counselling provided. Motherisk algorithm of treatment followed (n = 29)	PUQE score	Pre-emptive group, 43.3% reduction in HG between the previous pregnancy (19/30) and the present one (6/30), vs. 17.2% reduction (from 11/29 to 6/29) in the C group (p = 0.047) Pre-emptive group: 70% fewer cases of moderate–severe NVP (PUQE ≥ 11) [4/26 (15.4%) vs. 9/23 (39.13%)] in the C group during the first 3 weeks of NVP (p < 0.04)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Pyridoxine/doxylamine vs. metoclopramide									
Ashkenazi-Hoffnung 2013 ³⁶	Recruitment via BELTIS, a free call-in centre for queries regarding drug use during pregnancy and lactation, CA, USA	To evaluate the efficacy and safety of pyridoxine (50 mg twice daily) and doxylamine (25–50 mg) as an alternative treatment for NVP, described as a prospective case–controlled observational study	58, NR	Women who contacted the BELTIS regarding treatment of NVP were eligible for inclusion	Categorised as having moderate–severe NVP: I = 28/29 (97%), C = 18/26 (69%) (NOT CLEAR)	Pyridoxine (50 mg) twice daily. If vomiting persisted plus doxylamine (25 mg) at night, with two additional doses of 12.5 mg if required (n = 29)	Metoclopramide (10 mg) 8-hourly as needed (n = 29)	Maternal report on the severity of NVP (mild, moderate, severe) and efficacy of treatment (no or mild, moderate, high)	Moderate/severe symptoms in 28/29 of treatment group (97%) vs. 18/26 in the C group (69%) (p < 0.01) at baseline Comparison of the treatment vs. C following treatment = similar efficacy in 20/29 (69%) vs. 18/25 (72%) of women (p = 0.65)
Pyridoxine/doxylamine vs. ondansetron									
Capp 2014 ³⁵	Naval Medical Centre, San Diego, CA, USA	To determine whether ondansetron or the combination of doxylamine plus pyridoxine was superior for treatment of NVP, double-blind RCT	Trial was only available in abstract form so reporting lacks detail. Severity was NR so likely to have included all ranges from mild to severe	NR	Thirty-six women were enrolled in the trial with 30 fully completing the study. Presumably 18 were randomised in each group initially, NR – authors stated first trimester of pregnancy only (UNCLEAR)	4-mg ondansetron plus a placebo tablet every 8 hours for 5 days	25-mg pyridoxine plus 12.5-mg doxylamine every 8 hours for 5 days	VAS for nausea and vomiting	Patients randomised to ondansetron demonstrated a greater reduction in nausea as compared with those taking pyridoxine and doxylamine (p < 0.05) Furthermore, women taking ondansetron reported less vomiting (p < 0.05)

continued

TABLE 12 Results for pyridoxine-doxylamine interventions for NVP (*continued*)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Oliveira 2013 (abstract) ⁹⁵	Naval Medical Centre, San Diego, CA, USA	Is ondansetron or the combination of doxylamine plus pyridoxine superior for treating NVP, double-blind RCT	17, first trimester	Women with nausea and vomiting during first trimester and present at the emergency department	NR (MILD-MODERATE)	One tablet of pyridoxine (25 mg) plus one tablet of doxylamine (12.5 mg) (P + D group) every 8 hours for 5 days (n = 9)	One tablet of ondansetron (4 mg) plus a second (placebo) tablet (O group) (n = 8)	VAS for nausea, and vomiting	Greater mean reduction in nausea with ondansetron (56–15 mm) vs. pyridoxine plus doxylamine (27–29 mm); p = 0.02 No difference in vomiting between groups (28–25 mm for ondansetron vs. 10–31 mm for pyridoxine plus doxylamine; p = 0.38)

BELTS, Beilinson Teratology Information Service; C, control; I, intervention; N/A, not applicable; NR, not reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

Koren and colleagues⁸⁴ did not report on pregnancy outcomes or adverse events. UKTIS data on doxylamine/pyridoxine combination treatment are provided in *Appendix 7*.

Doxylamine/pyridoxine versus ondansetron

The double-blind RCT of Oliveira and colleagues⁹⁵ was in abstract form only and randomised women in the first trimester of pregnancy requesting treatment for NVP to either one tablet of ondansetron (4 mg) plus a second (placebo) tablet or one tablet of pyridoxine (25 mg) plus one tablet of doxylamine (12.5 mg) administered every 8 hours for 5 days.⁹⁵ All study medications were identical in appearance. Capp and colleagues⁶⁵ also compared 4-mg ondansetron plus a placebo tablet administered every 8 hours for 5 days against 25-mg pyridoxine plus 12.5-mg doxylamine.⁶⁵

Combined severity score

No combined score reported.

Nausea outcomes

Oliveira and colleagues⁹⁵ found a significantly greater mean reduction in nausea ($p = 0.02$) using the VAS in patients using ondansetron (56 ± 15 mm) compared with those taking pyridoxine plus doxylamine (27 ± 29 mm). A significant difference in favour of ondansetron was also reported by Capp and colleagues.⁶⁵ However, no detailed scores were provided as the paper was in abstract form only ($p < 0.05$).

Vomiting outcomes

There was no significant difference in vomiting between the two groups reported by Oliveira and colleagues⁹⁵ (28 ± 25 mm for ondansetron vs. 10 ± 31 mm for pyridoxine plus doxylamine; $p = 0.38$). However, Capp and colleagues⁶⁵ found a significant improvement in the ondansetron groups in terms of reduction in vomiting using the VAS ($p < 0.05$), although, as with nausea scores, no exact values were reported in the abstract.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

Oliveira and colleagues⁹⁵ did not report on pregnancy outcomes or adverse events. The trial of Capp and colleagues⁶⁵ did not report on pregnancy outcomes either but found no statistically significant difference between groups with respect to sedation or constipation ($p > 0.05$) (see *Appendix 8*). UKTIS data on doxylamine/pyridoxine combination treatment are provided in *Appendix 7*.

Pre-emptive doxylamine/pyridoxine

One RCT¹¹⁷ compared pre-emptive treatment with Diclectin before the onset of symptoms of NVP, versus when the symptoms first began in patients at high risk for recurrence of severe NVP.

Pregnancy-Unique Quantification of Emesis and Nausea scale

In the pre-emptive group there were 70% fewer cases of moderate–severe NVP (PUQE score of ≥ 11) compared with the control group [4/26 (15.4%) vs. 9/23 (39.13%)] during the first 3 weeks of NVP ($p = 0.05$). In addition, in the pre-emptive group there was a 43% reduction in HG between the previous

pregnancy (19/30) and the present one (6/30) compared with a 17% reduction (from 11/29 to 6/29) in the control group ($p = 0.047$).

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

The trial of Maltepe and Koren¹¹⁷ did not report either pregnancy outcomes or adverse events. UKTIS data on doxylamine/pyridoxine combination treatment are provided in *Appendix 7*.

Summary

- The evidence available for pyridoxine/doxylamine combinations is varied, but two trials appeared to be at low risk of bias^{84,117} with the other two having an unclear risk of bias profile.^{65,95}
- The quality of the evidence is low and was downgraded due to clinical heterogeneity and sparseness of data in most comparisons. Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimates.
- Diclectin appears to be more effective at relieving symptoms of NVP than placebo.
- Ondansetron appears more effective at reducing nausea than pyridoxine plus doxylamine but there was no difference for vomiting.
- Limited data from a single small study³⁶ showed no difference in efficacy between pyridoxine plus vitamin B6 versus metoclopramide but was subject to selection bias.
- Pre-emptive treatment with Diclectin appears to result in a reduced risk of moderate/severe NVP compared with treatment initiation once symptoms begin.
- Further larger, well-conducted trials are required to test the effectiveness of Diclectin or pyridoxine/doxylamine in combination compared with other treatment options.

Chapter 9 Clinical effectiveness: antihistamines

Introduction

Antihistamines were used as an intervention to treat nausea and/or vomiting in three RCTs.^{68,71,90} Heterogeneity was observed in relation to the clinical setting, and the patient populations in which the studies were conducted, but most notably there were differences in interventions, comparators and outcomes reported in each trial. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*), given these differences, we did not attempt to perform meta-analyses and have thus reported a narrative summary only for each intervention and comparator set.

The three trials had varying risk of bias profiles, with just one of the trials appearing to be at low risk of bias,⁷¹ and the remaining two at high risk of bias due to concerns with regard to blinding and incomplete outcome reporting,⁶⁸ and selective outcome reporting.⁹⁰ The women were described as experiencing mild symptoms at baseline in all three trials (*Table 13*).

Antihistamines versus placebo

Two trials^{71,90} compared antihistamines and placebo: one trial⁷¹ in women with mild recurrent nausea and who had vomited at least three times per week over the previous 2 weeks; and the other trial⁹⁰ in women complaining of nausea and/or vomiting. Hydroxyzine hydrochloride (25 mg) capsules were given twice daily for 3 weeks in the trial of Erez and colleagues,⁷¹ and cyclizine pyridoxine (dose not reported) tablets twice daily for 2 weeks in the Monias⁹⁰ trial.

Author-defined symptom severity/relief scale

Hydroxyzine significantly relieved symptoms of nausea and vomiting compared with placebo in the trial of Erez and colleagues,⁷¹ resulting in partial or initial or complete relief in 82% of women, whereas the placebo produced some effect in only 22% of women ($p < 0.01$). This trial was at low risk of bias.

In the trial of Monias,⁹⁰ 78, 5 and 17 out of 100 women experienced complete, partial and no relief in the intervention group, respectively. In the placebo group (also $n = 100$), only 13 women experienced complete relief, five partial relief, and in 82 women no relief of symptoms was observed. The intervention relieved symptoms in a higher percentage of women than the placebo group; however, no formal statistical analysis was undertaken.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting scores reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

The trial of Erez and colleagues⁷¹ found no statistically significant differences in terms of miscarriage, perinatal outcomes and fetal outcomes between groups ($p > 0.05$), and only minor side effects (slight drowsiness) were reported by 7% of the intervention group. No pregnancy outcomes or adverse events were reported by Monias.⁹⁰ See *Appendix 8* for details, with UKTIS data on hydroxyzine reported in *Appendix 7*.

TABLE 13 Results for antihistamine interventions for NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Antihistamines vs. placebo									
Erez 1971 ⁷¹	Prenatal clinics, Kasimpasa Naval Hospital and Golcuk Naval Hospital, Ismit, Turkey	To investigate the effectiveness of hydroxyzine hydrochloride as an antiemetic in pregnancy and its effect on fetal outcomes, RCT	n = 150 (I n = 100, C n = 50); states treatment in first 2 months of pregnancy only	Women reporting recurrent nausea and had vomited at least three times per week over the previous 2 weeks	NR (MILD)	Hydroxyzine hydrochloride capsules (25 mg) two times a day (a.m. and 2 p.m. for 3 weeks) (n = 100)	Identical placebo capsules two times a day (a.m. and 2 p.m. for 3 weeks) (n = 50)	Patient assessment: (1) complete relief; (2) partial relief; and (3) no relief	Hydroxyzine, partial or complete relief in 82% of the patients; placebo, some effect in 22% of the patients (p < 0.01)
Antihistamines ± vitamin B6									
Diggory 1962 ⁶⁸	Antenatal clinic, Queen Charlotte's and Chelsea Hospital, London, UK	To compare meclozine hydrochloride ± pyridoxine with placebo and simple dietary advice, four-arm RCT	n = 139 (group 1 n = 29, group 2 n = 34, group 3 n = 41, group 4 n = 35); ≤ 14 weeks	Women attending antenatal clinics who were experiencing nausea or vomiting	NR (MILD)	Group 3 = dietary info sheet plus antihistamines (25 mg a.m., 50 mg p.m.). Group 4 = dietary info sheet plus antihistamines (25 mg a.m., 50 mg a.m. plus pyridoxine 50 mg a.m., 100 mg p.m.)	Group 1 = dietary info sheet only. Group 2 = dietary info sheet plus placebo	Author-defined severity scale based on disruption to life (good, fair, poor)	Group 1 good (n = 6), fair (n = 4); poor (n = 19); group 2 good (n = 5), fair (n = 11), poor (n = 18); group 3 good (n = 28), fair (n = 12), poor (n = 1); group 4 good (n = 22), fair (n = 11), poor (n = 2)
Monias 1957 ⁶⁰	Military hospital, Massachusetts, MA, USA	To evaluate the effectiveness of cyclizine plus pyridoxine in treating NVP, RCT	n = 200 (n = 100 each arm); 6–20 weeks	Women complaining of nausea and/or vomiting	NR (MILD)	Instructed to take two cyclizine plus pyridoxine tablets half an hour before breakfast and an additional tablet before lunch if required for 10 days. Dose not reported (n = 100)	Instructed to take two placebo tablets half an hour before breakfast and an additional tablet before lunch if required for 10 days (n = 100)	Author defined (complete, partial or no relief)	I group: complete relief n = 78; partial relief n = 5; no relief n = 17 Comparator: complete relief n = 13; partial relief n = 5; no relief n = 82

C, control; I, intervention; NR, not reported.

Antihistamines alone or in combination with Vitamin B6 versus control

One four-arm trial⁶⁸ compared antihistamines (meclizine hydrochloride) with and without vitamin B6 (pyridoxine) with a placebo group and a no treatment group in women experiencing mild nausea or vomiting. In the intervention arms, meclizine hydrochloride was given at doses of 25 mg and 50 mg in the morning and evening, respectively. In the combination group, pyridoxine was given at 50 mg in the morning and 100 mg in the evening.

Author-defined scale

The trial used an author-defined severity scale based on disruption to life whereby women were asked to judge symptoms on the basis of whether the impact of the treatment on restoring disrupted routine could be categorised as good, fair or poor. No differences between the control groups or the two intervention groups were observed ($p > 0.05$). However, based on subjective assessments of restoration of disrupted routine, both the antihistamine alone and antihistamine with vitamin B6 groups significantly improved ($p < 0.001$) in comparison to the two control groups (no treatment group: good $n = 6$, fair $n = 4$, poor $n = 19$; placebo group: good $n = 5$, fair $n = 11$, poor $n = 18$; antihistamine alone: good $n = 28$, fair $n = 12$, poor $n = 1$; antihistamine with vitamin B6: good $n = 22$, fair $n = 11$, poor $n = 2$).

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting scores reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or adverse events were reported by Diggory and Tomkinson⁶⁸ and there were no UKTIS data relating to meclizine hydrochloride.

Summary

- Of the three studies available for antihistamines, two were at high risk of bias^{68,90} while one was at low risk.⁷¹ The quality of the evidence was very low and was downgraded due to the reporting of author-defined scales to measure symptom severity, and other concerns about the overall risk of bias in included RCTs. We are very uncertain about the effectiveness estimates in terms of validation and estimates may change in future trials.
- Participants in all three studies^{68,71,90} had mild symptoms so generalisability is restricted.
- Use of antihistamines resulted in an improvement over a range of symptoms.
- The addition of vitamin B6 does not appear to improve effectiveness.
- Evidence is limited in both quantity and quality.
- Antihistamines appear to be better than placebo in reducing the severity of symptoms, but better-quality, large studies are required for this and all other comparators.

Chapter 10 Clinical effectiveness: dopamine antagonists

Introduction

Dopamine antagonists were used as an intervention to treat HG in the trial of Tan and colleagues.¹⁰⁶ The trial was a head-to-head comparison of two interventions (promethazine vs. metoclopramide) and was at low risk of bias. We will focus on the results from this trial here. We also identified a non-randomised prospective study¹²² which examined two study groups (67 patients treated with i.v. droperidol 1 mg/hour plus diphenhydramine 25–50 mg every 6 hours; and 34 patients treated with i.v. droperidol 0.5 mg/hour plus diphenhydramine 50 mg every 6 hours; and a historical control of 54 patients receiving conventional antiemetic treatment). The study groups were then gradually weaned onto oral hydroxyzine and metoclopramide. This study is prone to extreme selection bias and other biases, and classified as ‘weak’ according to the EPHPP quality assessment tool. A summary of study conduct and results is depicted in *Table 14*, with safety data reported in *Appendix 8*.

Promethazine versus metoclopramide

The trial of Tan and colleagues¹⁰⁶ randomised women in early pregnancy (gestation of ≤ 16 weeks) with clinical HG to either 25-mg promethazine ($n = 76$) or 10-mg metoclopramide ($n = 73$), administered intravenously every 8 hours for 24 hours.

Combined severity score

No combined score reported.

Nausea outcomes

There were no significant differences in median nausea scores during all monitored time points (at baseline and at 8, 16 and 24 hours) between the promethazine and metoclopramide groups ($p = 0.95$). The median nausea score at 24 hours was 2 (IQR 1–4) in the promethazine group and 2 (IQR 1–5) in the metoclopramide group ($p = 0.99$).

Vomiting outcomes

There was no significant difference in the median number of vomiting episodes between the promethazine and metoclopramide groups [median episodes were 2 (range 0–3) and 1 (range 0–5) in the promethazine and metoclopramide groups respectively; $p = 0.8$].

Retching score

No independent retching score reported.

Safety outcomes

No pregnancy outcomes were reported by Tan and colleagues¹⁰⁶ and only minor side effects, with significantly more women in the promethazine group reporting drowsiness ($p = 0.001$) and dizziness ($p < 0.001$). Additional UKTIS data on promethazine are reported in *Appendix 7*.

TABLE 14 Results for dopamine antagonist interventions for NVP

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Dopamine antagonists vs. conventional treatment									
Ferreira 2003 ²²	Sainte-Justine Hospital, Montréal, Québec, Canada	To compare the efficacy of the droperidol/diphenhydramine combination with other conventional treatments used. Described as a case-control study but actually a cohort study	Group A n = 54, group B n = 67, group C n = 34 Group A = 11.1 ± 4.6, group B = 10.3 ± 3.9, group C = 10.4 ± 2.8	Women hospitalised for HG with at least one of the three following criteria: weight loss ≥ 5% of pre-pregnancy weight, ketonuria (objectified on urinary test strips), or hypokalaemia (< 3.5 mEq/l)	Mean score National Cancer Institute's Common Toxicity Criteria scale: nausea (scale 0-3) group A = 2.24 ± 0.8, group B = 1.33 ± 1.05, group C = 1.03 ± 1.00; vomiting (scale 0-4) group A = 1.06 ± 1.00, group B = 0.34 ± 0.66, group C = 0.41 ± 0.74 (MODERATE)	Two groups (groups B, C). Both groups received i.v. rehydration plus multivitamins. Group B: i.v. droperidol infusion at 1 mg/hour plus i.v. diphenhydramine (25-50 mg) 6-hourly Group C: i.v. droperidol at 0.5 mg/hour plus i.v. diphenhydramine (50 mg) 6-hourly. Gradually oral antiemetic treatment consisting of hydroxyzine and metoclopramide was started	A retrospective control group (A), treated with a variety of antiemetic treatments (i.m. chlorpromazine, i.v. dimenhydrinate, i.v. metoclopramide, or a combination of doxylamine-pyridoxine (Diclectin) taken orally according to the physicians' choice	National Cancer Institute's Common Toxicity Criteria scale: nausea analogue scale varies from 0 to 3, and the vomiting scale varies from 0 to 4	Mean score on day 1 (SD) to during hospitalisation: Nausea (n = 155): group A = 2.24 (SD 0.87) to 1.48 (SD 0.49); group B = 1.33 (SD 1.05) to 0.78 (SD 0.70); group C = 1.03 (SD 1.00) to 0.58 (SD 0.60); p < 0.001 Vomiting: group A = 1.06 (SD 1.00) to 0.62 (SD 0.54); group B = 0.34 (SD 0.66) to 0.25 (SD 0.36); group C = 0.41 (SD 0.74) to 0.24 (SD 0.39); p < 0.001 in favour of group B This trend continued through the hospital stay, although it was not clear whether or not analysis accounted for group differences at baseline

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Dopamine antagonists vs. metoclopramide									
Tan 2010 ¹⁰⁶	Gynaecology ward, university hospital in Kuala Lumpur, Malaysia	To compare the effects of promethazine with those of metoclopramide for HG, double-blind RCT	n = 159: metoclopramide 9.2 ± 2.3, promethazine 9.3 ± 2.6	Women hospitalised with presumed hyperemesis who were determined clinically to require i.v. antiemetic therapy	Nausea score via VAS: I = median 5 (IQR 2.75–7), C = median 5 (IQR 1.5–7) (MODERATE)	Metoclopramide (10 mg) given i.v. after randomisation, at 8, 16 and 24 hours (n = 79)	Promethazine (25 mg) given i.v. after randomisation and at 8, 16 and 24 hours (n = 80)	VAS for nausea Episodes of vomiting	Vomiting: metoclopramide = 1 (IQR 0–5), promethazine = 2 (IQR 0–3); p = 0.81 Nausea at 24 hours: metoclopramide = 2 (IQR 1–5), promethazine = 2 (IQR 1–4); p = 0.99 (Nausea scores at 8 and 16 hours also showed no significant difference) Repeated measures analysis of variance for nausea score: p = 0.95
C, control; I, intervention; i.m., intramuscular.									

Summary

- Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP.
- The quality of evidence was very low and was downgraded due to sparseness of data and imprecision, although the one trial¹⁰⁶ that reported a comparison on promethazine and metoclopramide was at low risk of bias.
- We are very uncertain about the estimates of effectiveness. Further, well-conducted studies are required to compare the effectiveness of metoclopramide against other comparators.

Chapter 11 Clinical effectiveness: serotonin antagonists (ondansetron)

Introduction

A total of five trials^{57,72,75,81,105} and one cohort study¹²¹ compared the serotonin antagonist (ondansetron) against a range of alternatives for the treatment of women experiencing various severities of nausea in pregnancy. Of these, one study focussed on the safety of ondansetron versus the usual treatment regimen,¹²¹ with symptom severity not specified. Three trials tested ondansetron against metoclopramide, with participants' symptoms classified as mild to moderate in two trials,^{75,81} and severe in one trial.⁵⁷ The remaining two trials compared ondansetron with antihistamines,^{72,105} with women classed as experiencing symptoms at the moderate to severe end of the spectrum.

Two of the identified trials were found to have a low risk of bias.^{57,81} The trial of Sullivan and colleagues¹⁰⁵ was classed as unclear due to insufficient information, with the remaining two found to carry a high risk of bias.^{72,75} The included cohort study of Einarson and colleagues¹²¹ was classed as 'weak' according to the EPHPP quality assessment tool. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*), given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses and have thus reported a narrative summary only for each intervention and comparator set. A summary of study conduct and results is depicted in *Table 15*, with safety data reported in *Appendix 8*.

Ondansetron versus usual treatment

Einarson and colleagues¹²¹ examined ondansetron versus the usual treatment regimen.¹²¹ As this involved a telephone survey of women already taking ondansetron, however, no direct measures of symptom relief were assessed. The outcomes of interest related to the incidence of fetal abnormalities in the surveyed patient population. These data are provided in *Appendix 8*, with additional UKTIS data on the intervention detailed in *Appendix 7*.

Ondansetron versus metoclopramide

Combined severity score

No combined score reported.

Nausea outcomes

All three trials comparing ondansetron against metoclopramide assessed nausea severity using the 10-point VAS.^{57,75,81}

The trial of Kashifard and colleagues⁸¹ (classed as low risk of bias), measured nausea in a group of women taking ondansetron versus a comparison group taking metoclopramide for pregnancy-related nausea and vomiting. They found that nausea scores for the ondansetron group were significantly less on the third and fourth days of treatment in comparison with those taking metoclopramide ($p = 0.024$ and $p = 0.023$ respectively). However, there was no significant difference between groups in the overall nausea trend over time.

TABLE 15 Results for serotonin antagonist interventions for NVP/HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ondansetron vs. usual treatment Abas 2014 ⁵⁷	Full-service, state-funded university hospital, Kuala Lumpur, Malaysia	To compare effectiveness of ondansetron with metoclopramide in the treatment of HG	n = 160; ≤ 16 weeks with clinical dehydration and ketonuria (of 2+ or greater)	Presence of nausea and intractable vomiting sufficient to cause dehydration and metabolic disturbance of a severity to require hospitalisation	Nausea score: ondansetron median 8 (IQR 7–9); metoclopramide median 8 (IQR 7–10) (SEVERE)	4-mg ondansetron was diluted in 100-ml normal saline and standard 100-ml normal saline packs were labelled as A or B (n = 80)	10-mg metoclopramide was diluted in 100-ml normal saline and the standard 100-ml normal saline packs were labelled as A or B (n = 80)	VAS for nausea Episodes of vomiting	Nausea visual numeric rating scale scores [median (IQR)] at 8 hours [4 (IQR 3–6) vs. 5 (IQR 4–6)], 16 hours [3 (IQR 1–4) vs. 3 (IQR 2–4; 75)], and 24 hours [1 (IQR 1–3) vs. 2 (IQR 1–3)] after randomisation were not significantly different when assessed separately at each time point Vomiting episodes in the first 24 hours were median 1 (IQR 0–2) vs. median 2 (IQR 0–2; 75) (p = 0.38) for ondansetron and metoclopramide respectively Repeated measures analysis of variance also showed no difference across the trial arms (p = 0.22) for nausea visual numeric rating scale scores, although generally across both arms, nausea score lessened significantly over at 24 hours (12.5% compared with 30%; p = 0.01; number needed to treat to benefit, 6) for the ondansetron arm

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Einarson 2004 ²¹	Women calling the Helpline or TIS at the MotherRisk Program in Toronto, Canada, or the MotherSafe Program in Sydney, Australia	To determine whether or not the use of ondansetron during pregnancy is associated with an increased risk for major malformations. Prospective, three-arm, comparative study	Group 1 n = 188 (ondansetron), group 2 n = 176 (antiemetics), group 3 n = 176 (non-teratogen)	Women who called either the MotherRisk and MotherSafe counselling services, who were taking ondansetron for NVP	NR (NOT CLEAR)	Group 1: women who called either service and were taking ondansetron at the time of call within a 2-year period were enrolled Group 2: women who called one of the helplines but were not exposed to ondansetron (used other antiemetics including Diclectin, metoclopramide, phenothiazines and ginger) Group 3: women exposed to other drugs considered safe to use in pregnancy or those who had not used any medication	Group 2: women who called one of the helplines but were not exposed to ondansetron (used other antiemetics including Diclectin, metoclopramide, phenothiazines and ginger) Group 3: women exposed to other drugs considered safe to use in pregnancy or those who had not used any medication	PUQE NR	NR
Ondansetron vs. metoclopramide									
Kashifard 2013 ³⁰	Ruhani Hospital of Babol University of Medical Sciences, North Iran	To compare the effectiveness of ondansetron vs. metoclopramide in the treatment of HG, double-blind RCT	n = 83, gestational age in weeks 8.7 (SD 2.6) both groups	Women vomiting three times a day with weight loss > 3 kg and presence of ketonuria	NR (MILD-MODERATE)	Ondansetron hydrochloride tablets (4 mg), three times a day for 1 week. Dose gradually reduced and discontinued: two times a day for 3 days; once a day for 4 days. Medication stopped after second week	Metoclopramide (10 mg), three times daily following same regime	VAS for nausea Episodes of vomiting	Nausea was significantly less in the ondansetron group on third and fourth days of treatment vs. metoclopramide group ($p = 0.024$ and $p = 0.023$, respectively) Vomiting episodes in the ondansetron group were fewer than the metoclopramide group from the second to the eighth days Trend of change for vomiting in the ondansetron group was lower ($p = 0.045$) No difference in nausea trend

continued

TABLE 15 Results for serotonin antagonist interventions for NVP/HG (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ghahiri 2011 ²⁶	Shahid Beheshti Hospital, Iran	To compare the effectiveness of ondansetron vs. metoclopramide in the treatment of NVP, RCT	n = 70; metoclopramide 12 weeks (SD 3.8), ondansetron 10.8 weeks (SD 3.3)	Women suffering from chronic NVP, requiring hospitalisation and treatment	Mean (SE): nausea VAS 1.3.14 (SE 0.55), C 3.09 (SE 0.66); vomiting VAS 1.2.29 (SE 0.71), C 2.09 (SE 0.78) (MILD-MODERATE)	All participants rehydrated with i.v. fluids plus 10-mg metoclopramide twice daily orally for 3 weeks (n = 35)	All participants rehydrated with i.v. fluids plus ondansetron (4 mg) orally twice daily for 3 weeks (n = 35)	VAS for nausea Episodes of vomiting	No difference in severity of NVP at 3 days or 1 week At 2 weeks nausea was less in ondansetron group, (p = 0.05) and vomiting was less in the metoclopramide group (p = 0.04) At 3 weeks vomiting was less in the ondansetron group (p = 0.02)
Ondansetron vs. antihistamines									
Sullivan 1996 ⁶⁵	Women admitted to the University of Mississippi Medical Centre, MS, USA	To investigate whether or not ondansetron may be an effective antiemetic in pregnancy, RCT	n = 30, (n = 15 in each arm); I 11.0 (SD 2.7), C 10.2 (SD 3.8)	Women admitted to hospital with severe HG; > 5 lb weight loss, ketonuria > 80 mg/dl, hypokalemia < 3.0 mEq/dl, hyponatremia < 134 mEq/dl requiring i.v. replacement, positive serum acetone, two or more visits to obstetric emergency department for HG which required i.v. hydration or promethazine suppositories, as an outpatient	NR (MODERATE-SEVERE)	i.v. hydration. Ondansetron (10 mg) given intravenously in 50 ml of compatible i.v. fluid over 30 minutes. First dose mandatory, then as needed 8-hourly. I.v. hydration continued until patient ingesting a bland diet. Discharged with promethazine suppositories plus dietary instructions. Seen in clinic on a weekly basis. If no change after 48 hours patient considered treatment failure and was excluded	As I except promethazine (50 mg) given intravenously in 50 ml of compatible i.v. fluid over 30 minutes for initial and subsequent inpatient doses	VAS for nausea	Daily VAS score presented graphically, no numerical values presented No significant difference between treatments over 5 days of treatment via p-values

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Eftekhari 2013 ²	University hospital, Kerman, Iran	To compare the effectiveness of treating HG with either ondansetron or promethazine, RCT	n = 60 (n = 30 in each group); 171.56 days (SD 15–125), C 80.06 days (SD 35–128)	Women with chronic vomiting, not able to take oral fluids, dehydration, weight loss, in need of hospitalisation	NR (MODERATE–SEVERE)	All participants rehydrated with i.v. fluids. The ondansetron group received 8 mg i.m. 8-hourly for 48 hours	All participants rehydrated with i.v. fluids. Promethazine group received 25 mg i.m. 6-hourly for 48 hours	Author-defined symptom and relief scales	Mean change severity: ondansetron = 6.4 (SD 2.02), promethazine = 5.34 (SD 3.1); p = 0.46 Relief: ondansetron = 12.16 (SD 3.7), promethazine = 11.65 (SD 3.4); p = 0.178

C, control; I, intervention; i.m., intramuscular; NR, not reported; TIS, Teratology Information Service.

Ghahiri and colleagues⁷⁵ (classified as unclear risk of bias), compared women taking metoclopramide versus a comparison group taking ondansetron for pregnancy-related nausea and vomiting. They found no difference in symptom severity at either 3 days or 1 week. However, nausea was significantly lower in the ondansetron group at 2 weeks ($p = 0.05$).

Finally, the trial of Abas and colleagues⁵⁷ compared the effectiveness of ondansetron with metoclopramide in the treatment of HG. No significant difference in the median (IQR) of well-being VAS scores were found between treatment arms [intervention = 9 (IQR 8–10) vs. comparator = 9 (IQR 7–10) ($p = 0.33$)].

Vomiting outcomes

Kashifard and colleagues⁸¹ measured vomiting outcomes using the number of episodes of vomiting recorded per group. They reported that both fewer episodes of vomiting were found in the ondansetron group compared with the metoclopramide group, and that the trend over time for vomiting in the ondansetron group was significantly lower compared with the comparator ($p = 0.045$). Ghahiri and colleagues⁷⁵ found no differences in episodes of vomiting between groups until 2 weeks, when episodes were lower in the metoclopramide group ($p = 0.04$). However, by 3 weeks episodes of vomiting were significantly lower in the ondansetron group ($p = 0.02$). Finally, no significant difference was reported by Abas and colleagues⁵⁷ in terms of median (IQR) number of vomiting episodes in the first 24 hours between treatment arms [intervention median 1 (IQR 0–2) compared with comparator median 2 (IQR 0–2.75); $p = 0.38$].

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No adverse events were reported in the trial of Kashifard and colleagues,⁸¹ in terms of either pregnancy outcomes or side effects. Ghahiri and colleagues⁷⁵ found no significant difference between groups in relation to minor side effects [headaches, dizziness, sedation or anxiety ($p > 0.05$)]. However, although Abas and colleagues⁵⁷ also reported some minor side effects (difficulty sleeping, dizziness, diarrhoea, headache, palpitations and skin rash) in similar proportions across the trial arms ($p > 0.5$), significant differences were found in self-reported drowsiness ($p = 0.011$) and dry mouth ($p = 0.003$) in favour of ondansetron. Full details are presented in *Appendix 8* (although as previously emphasised, given the anticipated rarity of these events small trials are likely to provide unreliable estimates). Additional UKTIS data on ondansetron and metoclopramide enquiries can be found in *Appendix 7*.

Ondansetron versus antihistamines

Author-defined relief scale

The study by Eftekhari and Mehralhasani⁷² used an author-defined scale to assess change in symptom severity and relief. No significant difference between treatment groups was found for either category (severity $p = 0.46$, relief $p = 0.178$). Furthermore, this trial was found to have a high risk of bias due to inadequate allocation concealment procedures.

Nausea outcomes

Sullivan and colleagues¹⁰⁵ (classified as unclear risk of bias due to lack of information), assessed daily scores using the VAS. No significant difference between ondansetron and antihistamine was reported during the 5-day treatment period (data presented graphically, therefore no specific p -value is available).

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or adverse event data were reported in the trial of Eftekhari and Mehralhasani.⁷² In the trial of Sullivan and colleagues,¹⁰⁵ eight women reported sedation versus none in the ondansetron group ($p = 0.002$). UKTIS data on ondansetron related inquiries are presented in *Appendix 7*.

Summary

- The evidence available for serotonin receptor antagonists, specifically ondansetron, was predominantly at high or unclear risk of bias with only one at low risk.⁸¹
- Evidence for the comparison of ondansetron with metoclopramide was mixed, with both drugs improving symptoms.
- Low risk of bias studies found ondansetron more effective at reducing symptoms of vomiting compared with metoclopramide after 4 days.
- Both ondansetron and antihistamine improved symptoms with no evidence of a difference between them.
- Overall, ondansetron does reduce the severity of symptoms, but more larger, better-quality studies are required to assess benefit over other comparators.

Chapter 12 Clinical effectiveness: intravenous fluids

Introduction

Two studies compared the effectiveness of i.v. fluids for the treatment of women with HG.^{69,108} One trial by Tan and colleagues¹⁰⁸ tested different compositions of i.v. solution (dextrose plus saline vs. saline only)¹⁰⁸ and was judged as low risk of bias. The other study compared the use of diazepam against i.v. fluids containing vitamins⁶⁹ but was judged as carrying an unclear risk of bias due to lack of sufficient information in a number of areas. Both papers described the trial participants as suffering from HG; however, we classified severity as either moderate,⁶⁹ or moderate to severe¹⁰⁸ based on the participant data provided. As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*), given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses and have thus reported a narrative summary only for each intervention and comparator set. A summary of study conduct and results is depicted in *Table 16*, with safety data reported in *Appendix 8*.

Dextrose saline versus saline only

Combined severity score

No combined score reported.

Nausea outcomes

The 10-point VAS was used by Tan and colleagues¹⁰⁸ to assess nausea in their trial comparing dextrose plus saline versus saline only i.v. fluids. Although the difference after 24 hours between groups was not found to be significant [dextrose plus saline, VAS score = 2 (SD 1–4), saline only, VAS score = 2 (SD 2–4); $p = 0.39$], repeated-measures of the analysis of variance for nausea scores detected a significantly greater reduction in favour of the dextrose plus saline preparation ($p = 0.046$).

Vomiting outcomes

Tan and colleagues¹⁰⁸ measured number of episodes of emesis to assess vomiting outcomes and did not detect a difference in the median change in episodes between groups [both groups = 0 (IQR 0–2); $p = 0.66$].

Retching outcomes

No independent retching scores reported.

Safety outcomes

Tan and colleagues¹⁰⁸ did not report any pregnancy outcomes or side effects. No specific UKTIS data were available on this intervention.

Intravenous fluids versus intravenous fluids plus diazepam

Combined severity score

No combined score presented.

TABLE 16 Results for i.v. fluid interventions for NVP and HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
<i>i.v. fluids D-Saline vs. N-Saline</i>									
Tan 2013 ¹⁰⁸	University hospital, Kuala Lumpur, Malaysia	To compare 5% dextrose, 0.9% saline against 0.9% saline solution in the i.v. rehydration of HG, double-blind RCT	n = 222 (n = 111 each group); I 9.8 ± 2.8, C 9.8 ± 2.5	Women at their first hospitalisation for HG (intractable nausea and vomiting of pregnancy with dehydration and starvation clinically judged to require hospitalisation for i.v. rehydration and antiemetic drug administration)	VAS median nausea score 9 (QR 7–10) for both groups (MODERATE–SEVERE)	5% dextrose, 0.9% saline by i.v. infusion, 125 ml/hour over 24 hours plus potassium chloride 24 hours plus potassium chloride (9.5 mmol) as required, plus multivitamin containing 250-mg thiamine given intravenously. i.v. antiemetic prescribed according to health-care providers' preference. If capillary glucose > 8 mmol/l, infusion fluid changed to open-label standard 0.9% saline	0.9% saline by i.v. infusion, 125 ml/hour over 24 hours plus potassium chloride (9.5 mmol) as required, plus multivitamin containing 250-mg thiamine given intravenously. i.v. antiemetic prescribed according to health-care providers' preference. If capillary glucose > 8 mmol/l, infusion fluid changed to open-label standard 0.9% saline	Numerical scale 0–10 for nausea Episodes of vomiting	Median vomiting episodes both groups 0 (IQR 0–2); p = 0.66 Nausea in D-Saline group at 24 hours = 2 (IQR 1–4), N-Saline group = 2 (IQR 2–4); p = 0.39 Repeated-measures analysis of variance of nausea score: p = 0.046 in favour of D-Saline
<i>i.v. fluids ± diazepam</i>									
Ditto 1999 ⁶⁹	Department of obstetrics and gynaecology, Siena University Hospital, Italy	The efficacy of parenteral fluids with vitamins with or without diazepam; sedation in cases of HG	n = 50; i.v. fluids plus diazepam group 11.2 weeks ± 3.17, i.v. fluids only 11.5 weeks ± 2.96	Women with HG	Numerical values for nausea and baseline not provided. Proportion of women who had lost > 5% of pre-pregnancy weight: I 52%, C 56% (MODERATE)	i.v. fluids plus multivitamins given plus diazepam (10 mg) i.v. twice daily. Discharged with oral diazepam tablets (5 mg), twice daily (n = 25)	i.v. fluids plus multivitamins only. Discharged with placebo tablets (n = 25)	VAS for nausea Episodes of vomiting	There was a significant decrease in nausea in both groups but for women on diazepam the reduction was significantly greater on day 2 (p < 0.002) and day 3 than for the comparator group

C, control; D-Saline, dextrose saline; I, intervention; N-Saline, normal saline.

Nausea outcomes

For patients with a nausea score of 4+ (classed by the authors as severe nausea), significant reductions for both groups were found on the second and third days of therapy using the VAS ($p < 0.05$) in the trial of Ditto and colleagues⁶⁹. They reported that on day 2, this reduction was significantly greater in the diazepam group ($p < 0.002$); however, post treatment, the difference between groups was not significant (no p -value reported).

Vomiting outcomes

Ditto and colleagues⁶⁹ assessed vomiting outcomes via number of vomiting episodes. No significant difference between groups was observed (exact p -value not provided).

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No side effects were reported in the trial of Ditto and colleagues,⁶⁹ and no statistically significant differences were reported in terms of gestation at delivery; preterm delivery; caesarean section rate; mean birthweight or neonatal abnormalities (p -value not reported). As above, no specific UKTIS data were available on this intervention.

Summary

- The evidence available for i.v. fluids was at low¹⁰⁸ or unclear⁶⁹ risk of bias.
- i.v. fluid improves reported symptoms. Dextrose saline may be more effective at improving nausea over time for those with moderate nausea.
- Diazepam appears to be more effective than i.v. fluids alone at reducing nausea on day 2 but there was no evidence post treatment for those with moderate/severe nausea.
- Overall, i.v. fluids help correct dehydration and improve symptoms, dextrose saline may be more effective at reducing nausea than normal saline. (The lower concentration of sodium in dextrose saline may exacerbate any pre-existing hyponatraemia. High doses/concentrations of dextrose solutions may increase the risk of Wernicke's encephalopathy, but concentrations in dextrose saline are unlikely to provoke this response.)
- Future studies are required which focus on interventions given alongside rehydration therapy.

Chapter 13 Clinical effectiveness: transdermal clonidine

Introduction

One trial by Maina and colleagues⁸⁵ studied the efficacy of a 5-mg transdermal clonidine patch in the treatment of HG, in a double-blind, placebo-controlled, crossover RCT. Patients were classified at the severe end of the HG spectrum, with the trial judged as having a low risk of bias. A summary of study conduct and results is depicted in *Table 17*, with safety data reported in *Appendix 8*.

Transdermal clonidine versus placebo patch

Pregnancy-Unique Quantification of Emesis and Nausea scale

Maina and colleagues⁸⁵ assessed overall improvements in symptom severity using the PUQE scale. They reported that the transdermal clonidine patch led to a greater improvement in PUQE score [mean 6.3 (95% 5.5 to 7.1) vs. mean 8.5 (95% CI 7.7 to 9.3); $p = 0.001$].

Nausea outcomes

The 10-point VAS was also used to assess nausea severity. As above, the study found an improvement in VAS scores favouring the intervention group [mean 22 (95% CI 19 to 26) vs. mean 29 (95% CI 25 to 32); $p = 0.009$].

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

A number of pregnancy outcomes were reported in the trial of Maina and colleagues.⁸⁵ These included two major pregnancy complications: a central venous catheter-related sepsis and a postpartum haemorrhage. In addition, one baby was small for gestational age. However, no adverse fetal outcomes were reported (defined as miscarriage, stillbirth, preterm delivery or low birthweight) and no major or minor birth defects were detected. In addition, there was no significant association to an increase of adverse effects such as lassitude, drowsiness, dry mouth, headache, dizziness, fainting or skin intolerance in comparison with the placebo group ($p = 0.2$). These data are reported in *Appendix 8* (with the proviso that this was a small trial in terms of the generalisability of the results). No UKTIS data were available on transdermal clonidine patches.

Summary

- Evidence from one study⁸⁵ suggests that the use of transdermal clonidine patches may be effective for the treatment of severe HG, but more and larger studies are required to compare effectiveness against comparators.

TABLE 17 Results for transdermal clonidine interventions for HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Maina 2014 ⁸⁵	The trial was carried out in a single hospital setting (Ospedale Sant'Anna, Italy) after admission of patients	To study the efficacy of transdermal clonidine in the treatment of severe refractory HG, randomised, double-blind, placebo-controlled, crossover design (RCT)	Major grade of HG clinical severity who were unresponsive to standard antiemetic treatment (e.g. pyridoxine, metoclopramide or ondansetron plus an antireflux medication such as ranitidine or omeprazole)	Clinical severity was defined by a PUQE score index ≥ 13	Twelve women of gestational age 6–12 weeks (SEVERE)	With and without transdermal clonidine (5-mg clonidine patch or sham patch) for two consecutive periods of 5 days, other antiemetic drugs and antireflux drugs (ranitidine, omeprazole) being administered on a scheduled or as-needed basis. All women received i.v. hydration and supplementation with thiamine during both periods. The use of steroids was allowed as a rescue medication in the case of further worsening of symptoms	Sham patch crossover trial, see column 7	PUQE VAS	Clonidine vs. placebo, mean (95% CI): PUQE score 6.3 (95% CI 5.5 to 7.1), 8.5 (95% CI 7.7 to 9.3); $p = 0.001$ Mann-Whitney <i>U</i> -test, VAS score 22 (95% CI 19 to 26), 29 (95% CI 25 to 32); $p = 0.009$

I, intervention.

Chapter 14 Clinical effectiveness: outpatient/day case management

Introduction

One study compared the effectiveness of either midwife-led outpatient management versus standard inpatient care,⁸⁸ while another examined the feasibility of day case management,¹²⁰ for the treatment of moderate to severe HG/NVP. The trial of McParlin and colleagues⁸⁸ was judged as having a low risk of bias, while a 2007 case series study by Alalade and colleagues¹²⁰ was classed as of weak quality. A summary of study conduct and results is depicted in *Table 18*, with safety data reported in *Appendix 8*.

Outpatient management versus standard inpatient care

Pregnancy-Unique Quantification of Emesis and Nausea scale

In the trial of McParlin and colleagues,⁸⁸ the effectiveness of the same treatments (i.v. cyclizine and i.v. fluids), administered via an outpatient management setting versus standard inpatient care was assessed via mean change in PUQE scores.⁸⁸ No significant difference was found between groups [intervention = 6.9 (SD 4.1), comparator = 6.2 (SD 2.3); $p > 0.05$] when followed up for 7 days from initiation of treatment.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No adverse events were reported in the trial of McParlin and colleagues⁸⁸ and no significant differences between groups were found in the rates of miscarriage, termination of pregnancy, gestation at delivery, birthweight, incidence of small for gestational age, or admissions to a special care baby unit (see *Appendix 8* for full data). There were no UKTIS data available for this intervention and, given the anticipated rarity of these events, small trials are unlikely to provide unreliable estimates.

Day case management

The feasibility and efficacy of day case management of HG was assessed by Alalade and colleagues¹²⁰ using rates of patient discharges within a 24-hour period. These case series data are reported in *Appendix 8*.

Summary

- The evidence available for day case management was at low risk⁸⁸ and high risk¹²⁰ of bias.
- The identified studies indicate that day case management of women with moderate to severe symptoms is feasible and acceptable to women.
- The results indicate that day case management is as effective at improving severity scores as inpatient management for some women.
- More, larger studies are required to provide definitive results and women's views.

TABLE 18 Results for day case/outpatient management for NVP and HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
McParlin 2008 ¹⁸	Maternity assessment unit, Newcastle upon Tyne Hospitals, UK	To compare the effectiveness of midwife-led outpatient management versus standard inpatient care, RCT	n = 53, (I n = 27, C n = 26); I = 9.3 (SD 2.8), C = 10.3 (SD 2.9)	Women attending the maternity assessment unit with severe NVP/PUQE score of ≥ 9	Mean PUQE score: I 12.6 (2.2), C 11.5 (2.2) (MODERATE–SEVERE)	Cyclizine (50 mg, i.v.), given followed by 3 l of Hartmann's solution over 6 hours (i.e. 1 hour, 2 hours and 3 hours). 50 mg of oral thiamine given, discharged home with prescription for oral cyclizine (50 mg), three times daily. Women telephoned on day 3 and day 7 to offer ongoing support and advice	Admission to antenatal ward for routine care, i.v. fluids, i.v. cyclizine, oral thiamine	PUQE score	Mean change in PUQE score: I 6.9 (SD 4.1), C 6.2 (SD 2.3); p > 0.05
Alalade 2007 ²⁰	Whipps Cross University Hospital, London, UK	Feasibility and clinical efficacy of day case management of HG and patients' satisfaction	27; 8.8 weeks	Women suffering from NVP with severe dehydration; inability to retain fluids orally; urine dipstick 4+ ketones and electrolyte imbalance	Not applicable (MODERATE–SEVERE)	Direct admission to the gynaecological day ward. 2 l of normal saline were infused over 4 hours with 20 mmol of potassium chloride in each bag. Regular i.m./i.v. antiemetics were given. Discharged with oral antiemetics, folic acid and thiamine	N/A	Not assessed	Not reported

C, control; I, intervention; i.m., intramuscular; N/A, not applicable.

Chapter 15 Clinical effectiveness: corticosteroids

Introduction

Corticosteroids were tested against a variety of alternative treatments in seven individual studies.^{58,64,93,99,114,116,125} Two trials compared the effectiveness of corticosteroids against placebo tablets in patient populations categorised as experiencing moderate to severe symptoms.^{93,114} Three trials compared corticosteroids with either promethazine^{99,116} or phenergan suppositories⁵⁸ in patients classed as experiencing moderate, unclear or moderate to severe level symptoms respectively. One trial tested the effectiveness of steroids against metoclopramide for women with moderate to severe HG.¹¹⁶ As previously described (see *Chapter 3, Meta-analysis of included randomised controlled trials*), given the differences between trials in patient populations, settings, interventions and, in particular, the heterogeneous nature of the reported outcomes across trials, we did not attempt to perform meta-analyses and have thus reported a narrative summary only for each intervention and comparator set.

Of these trials, three were classed as carrying a low risk of bias^{64,93,99} and two were unclear due to lack of information.^{58,116} One study was judged to have a high risk of bias due to lack of blinding and unclear outcome reporting.¹¹⁴ A final case series study examined the effect of corticosteroids in comparison with the usual treatment regimen in women with severe HG.¹²⁵ A summary of study conduct and results is depicted in *Table 19*, with safety data reported in *Appendix 8*.

Corticosteroids versus placebo

Combined severity score

No combined score reported.

Nausea outcomes

Nelson-Piercy and colleagues⁹³ assessed the effect of corticosteroids compared with placebo tablets on nausea symptoms using the VAS. However, although the observed change was greater in the intervention group [median 6.5 (range 2.0–10.0) compared with median 4.0 (range –5.0 to 9.0)], this difference was not significant (relative risk for proportion with nausea 0.10; CI not reported).

Vomiting outcomes

In the trial of Nelson-Piercy and colleagues,⁹³ changes in vomiting status were assessed using numbers of patients still vomiting at 1 week of treatment; numbers of patients vomiting more than five times a day at day 5 of treatment; and via an author-defined scale which coded severity from 0 to 4.⁹³ Overall, although reported scores favoured the steroid treatment group across all three measures, the difference in effect sizes was not found to be significant [number still vomiting at 1 week, relative risk 1.4 (range 0.6–3.2); number vomiting more than five times per day, relative risk 2.5 (range 0.6–10.5); median reduction in vomiting score intervention 2.0 (range –1.0 to 4.0) vs. comparator 1.5 (range –3.0 to 4.0)].

The trial of Yost and colleagues¹¹⁴ assessed the difference in the impact on readmission rates between groups. These data are presented in *Appendix 8*.

Retching outcomes

No independent retching outcomes reported.

TABLE 19 Results for corticosteroid interventions for NVP and HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Corticosteroids vs. placebo									
Nelson-Piery 2001 ⁹³	Inpatient gynaecology wards in eight collaborating centres in the UK	To test the hypothesis that corticosteroids would lead to rapid and complete remission of the symptoms of NVP in cases of severe HG, RCT	n = 25, I n = 13, C n = 12	Onset NVP < 12 weeks, dependent on i.v. fluids for at least 1 week (first admission for HG) or 24 hours (second or subsequent admission), receiving regular treatment with at least 1 antiemetic, ketonuria on admission, vomiting at least two times per day or nausea so severe that unable to eat or drink, receiving regular treatment with oral thiamine or a single dose of parenteral thiamine	I: number vomiting five or more times per day = 6; number requiring one or more antiemetics = 4 C: number vomiting five or more times per day = 6; number requiring one or more antiemetics = 2 (MODERATE-SEVERE)	A 1-week course of prednisolone, 20 mg (four 5-mg tablets) orally 12-hourly, if, following 72 hours, a woman was still vomiting or vomiting tablets, and still dependent on i.v. fluids and electrolyte replacement, the therapy was changed to an i.v. equivalent [i.e. hydrocortisone (100 mg) 12-hourly]	A 1-week course of placebo (four 5-mg tablets) orally 12-hourly, if, following 72 hours, a woman was still vomiting or vomiting tablets, and still dependent on i.v. fluids and electrolyte replacement, the therapy was changed to an i.v. equivalent (i.e. N-Saline)	VAS for nausea Episodes of vomiting	Number still vomiting at 1 week: I 5, C 7; relative risk 1.4 (95% CI 0.6 to 3.2) Number vomiting five or more times per day: I 2, C 5; relative risk 2.5 (95% CI 0.6 to 10.5) Reduction in vomiting score: I median 2.0 (range -1.0 to 4.0), C median 1.5 (range -3.0 to 4.0) Nausea score improvement I = 6.5 (range 2.0-10.0), C = 4.0 (range -5.0 to 9.0), relative risk 0.10 for proportion with nausea

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Yost 2003 ¹¹⁴	Parkland Memorial Hospital, Dallas, TX, USA	To estimate the effect of corticosteroids in reducing the number of women requiring rehospitalisation for HG, RCT	n = 126, I n = 64, C n = 62	Women who previously had not responded to outpatient therapy and had three or four dipstick urinary ketones as evidence of severe dehydration	NR (MODERATE-SEVERE)	All women: i.v. rehydration with crystalloid until ketonuria cleared [first litre included thiamine (100 mg)]. Conventional treatment: promethazine (25 mg) and metoclopramide (10 mg) intravenously every 6 hours for 24 hours, followed by the same regimen orally until discharge from the hospital. The women were also randomised, I = methylprednisolone (125 mg, i.v.). This was followed by a tapering regimen of oral prednisone (40 mg for 1 day, 20 mg for 3 days, 10 mg for 3 days, and 5 mg for 7 days)	All women: i.v. rehydration with crystalloid until ketonuria cleared [first litre included thiamine (100 mg)]. Conventional treatment: promethazine (25 mg) and metoclopramide (10 mg) intravenously every 6 hours for 24 hours, followed by the same regimen orally until discharge from the hospital. The women were also randomised, C = i.v. placebo. This was followed by an identical tapering regimen)	N/A	Not reported
Corticosteroids vs. phenothiazines/promethazine/Phenergan® (Sanofi-Aventis)									
Adamczak 2007 ³⁸	Unclear, obstetrics and gynaecology department, New Jersey, NJ, USA	To compare SDP with Phenergan suppositories in the treatment of symptomatic NVP, RCT	n = 110, I n = 55, C n = 55; states 8-14 weeks	Patients presenting with NVP	Diagnosed with NVP (NOT CLEAR)	SDP included 8-mg t.i.d. tapered over 6 days	Phenergan suppositories were dosed at 25-mg q.i.d.	Episodes of vomiting	I (SDP), number of vomiting episodes: day 1: 7.1 (SD 1.8), day 3: 3.0 (SD 1.9), day 7: 1.8 (SD 1.6), day 14: 0.6 (SD 0.8) C, number of vomiting episodes: day 1: 6.6 (SD 1.9), day 3: 4.7 (SD 1.8), day 7: 3.9 (SD 1.7), day 14: 2.5 (SD 1.4) p-values for differences: day 1 p = 0.2, days 3, 7 and 14 p < 0.05

continued

TABLE 19 Results for corticosteroid interventions for NVP and HG (*continued*)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Safari 1998 ⁹⁹	Women's and children's hospital, Los Angeles, CA, USA	To compare the efficacy of methylprednisolone with that of promethazine for the treatment of HG, RCT	n = 40, I n = 20, C n = 20; I gestational age in weeks = 9.8 (SD 2.1), C gestational age in weeks = 9.5 (SD 2.7)	Diagnosed with HG, given i.v. hydration but with no resolution of symptoms, or second admission for HG	NR. The duration of HG before admission was longer in the promethazine group than in the methylprednisolone group ($p = 0.03$) (MODERATE–SEVERE)	Methylprednisolone (16 mg) orally three times a day for 3 days, followed by a tapering regimen, halving of dose every 3 days, to none during the course of 2 weeks (10 a.m., 2 p.m., 8 p.m.). After 2 days women were discharged with their study medication. If no improvement study allocation unblinded and patients withdrawn from further data collection	Promethazine (25 mg) orally three times a day for 2 weeks (10 a.m., 2 p.m., 8 p.m.). After 2 days women were discharged with study medication. If no improvement study allocation unblinded and patients withdrawn from further data collection	NR	Not reported

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Ziaei 2004 ¹¹⁶	Najmieh Hospital, Iran	To determine whether or not low dosages of prednisolone are effective in the treatment of outpatients with HG, RCT	n = 80, I n = 40, C n = 40	Vomiting more than three times per day during the last 72 hours or ketonuria that did not respond to dietary manipulation and caused weight loss	Episodes of vomiting/day: I 3 (range 2–5), C 3 (range 2–6) (MODERATE)	Prednisolone, (5 mg) given once in the morning for 10 days	Promethazine was administered (25 mg) three times daily for 10 days	VAS for nausea Episodes of vomiting	Severity of nausea: Mild/moderate: first 48 hours, I = 20 (50%), C = 30 (75%); third to tenth day, I = 6 (65%), C = 25 (62.5%); seventeenth day, I = 17 (43.6%), C = 12 (30.8%) Severe: first 48 hours, I = 20 (50%), C = 10 (25%); third to tenth day, I = 14 (35%), C = 15 (37.5%); seventeenth day, I = 22 (56.4%), C = 27 (69.2%) For prednisolone group OR (95% CI) of nausea: during first 48 hours OR 0.33 (95% CI 0.13 to 0.86); between third and tenth days OR 1.11 (95% CI 0.14 to 2.6); seventeenth day OR 1.7 (95% CI 0.68 to 4.4) Episodes of vomiting: first 48 hours, I median 3 (range 1–7), C median 1 (range 0–4), p = 0.04; third to tenth day, I median 1.5 (range 1–5), C median 1 (range 0–5), p = 0.80; tenth to seventeenth day, I median 3 (range 0–6), C median 3 (range 0–5), p = 1.0

continued

TABLE 19 Results for corticosteroid interventions for NVP and HG (continued)

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Corticosteroids vs. metoclopramide									
Bondok 2006 ²⁴	Intensive care unit of Ain Shams University Maternity Hospital, Cairo, Egypt	To compare the efficacy of pulsed hydrocortisone therapy with that of metoclopramide for the management of intractable HG, prospective double-blind study	n = 40, I n = 20, C n = 20; I = 10 ± 2.68, C = 11 ± 2.44	Women admitted to the intensive care unit with intractable hyperemesis (defined as severe persistent vomiting, ketonuria and weight loss > 5% of pre-pregnancy weight)	NR (MODERATE-SEVERE)	300 mg of i.v. hydrocortisone for 3 days, followed by a tapering regimen of 200 mg for 2 days, then 100 mg for another 2 days. Patients received three syringes, each every 8 hours, 10 ml each, one containing the drug diluted in normal saline and the other two containing normal saline. Nursing services recorded the daily number of emetic episodes	10 mg of metoclopramide, in a 10-ml syringe diluted in normal saline, intravenously every 8 hours for the same 7-day period	Episodes of vomiting	Mean vomiting episodes: hydrocortisone group reduced by 40.9% on second day, 71.6% on third day, and 95.8% on seventh day. Metoclopramide group reduced by 16.5% on second day, 51.2% on third day, and 76.6% on seventh day (p < 0.001)
Corticosteroids vs. usual treatment									
Moran 2002 ²⁵	Inpatient antenatal ward, Newcastle upon Tyne, UK	To document the effect of prednisolone therapy in women with defined severity of HG, case series	Thirty pregnancies in 25 women in treatment group, matched with 25 women treated with conventional therapy (i.v. fluids and antiemetics), treatment group median age in weeks = 9.6 (range 8.6–11.1)	HG requiring hospital admission plus weight loss > 5 kg and/or evidence of muscle wasting; onset of nausea and vomiting before 6 weeks; ketonuria on admission; i.v. fluids for > 1 week or > 24 hours if a repeat admission; failure of traditional antiemetic treatment; vomiting at least twice per day or severe nausea precluding any oral intake; gestation over 8 weeks	Assessed by physician and deemed severe enough to meet inclusion criteria (SEVERE)	Oral prednisolone (10 mg) 8-hourly prescribed, replacing traditional antiemetics. If unable to tolerate tablets stabilisation achieved with i.v. hydrocortisone (50 mg) 8-hourly. Prednisolone dosage was reduced in a stepwise fashion. Typically dosage decreased to 15 mg daily within 5 weeks, remaining between 12.5 mg and 15 mg for a further 3–8 weeks	Retrospective case series of 25 consecutive women hospitalised for hyperemesis but judged not to require steroid therapy	VAS for nausea	Six women completed VAS for nausea over 1 week Data for intensity show a clear pattern of resolution in the active group Three comparator group women received steroids Median number of inpatient days pre-steroid treatment = 8 (range 4–14) and after commencement = 3 (range 1–6.5)

C, control; I, intervention; N/A, not applicable; NR, not reported; N-Saline, normal saline; q.i.d., four times a day; SDP, solumedrol dose pack; t.i.d., three times a day.

Safety outcomes

Nelson-Piercy and colleagues⁹³ did not report side effects, but found no difference in birthweight or gestational age at delivery (excluding the triplet pregnancy) between groups ($p > 0.05$), or in terms of the numbers of babies born with birthweights less than the 5th centile ($p > 0.05$). Yost and colleagues¹¹⁴ reported no significant differences between groups in terms of spontaneous abortion ($p = 0.6$); gestational diabetes ($p = 0.96$); hypertension ($p = 0.2$); preterm delivery ($p = 0.4$); or primary or repeat caesarean delivery ($p = 0.06$ and $p = 0.5$ respectively). Full data are reported in *Appendix 8*. No UKTIS data were available on this intervention and it should be noted that given the anticipated rarity of these events, small trials are unlikely to provide unreliable estimates.

Corticosteroids versus promethazine (Phenergan)

Combined severity score

No combined score reported.

Nausea outcomes

Ziaei and colleagues¹¹⁶ used the VAS to assess severity of nausea and found that women who received promethazine responded better in the first 48 hours of treatment ($p = 0.02$). However, with continuation of the treatment, the difference decreased, and 1 week after completion of the treatment, the subjects who had received corticosteroids had fewer symptoms, although this difference was not significant ($p = 0.23$).

Vomiting outcomes

Number of episodes of vomiting was assessed in the trials of Ziaei and colleagues¹¹⁶ and Adamczak and colleagues.⁵⁸ In the Ziaei and colleagues¹¹⁶ study, median episodes of vomiting in the first 4 hours were lower in the promethazine group.¹¹⁶ However, as with nausea severity, by the end of the study period, there was no significant difference between groups ($p = 1.0$). In contrast, Adamczak and colleagues⁵⁸ reported that episodes of emesis were significantly lower in the steroid intervention group compared with those receiving Phenergan suppositories at days 3, 7 and 14 ($p < 0.05$ at all time points).⁵⁸

The primary outcome in the Safari and colleagues⁹⁹ study related to numbers of patients for whom therapy had failed at different points in the treatment period. These are reported in detail in *Appendix 8*.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcomes or side effects were reported by Adamczak and colleagues.⁵⁸ Safari and colleagues⁹⁹ reported no significant difference between the two groups in relation to birthweight or American Pediatric Gross Assessment Record scores at 1 and 5 minutes ($p > 0.05$). One patient in the methylprednisolone group was delivered at 35 weeks' gestation of a male infant with Smith–Lemli–Opitz syndrome. No pregnancy outcomes were reported by Ziaei and colleagues,¹¹⁶ but significant differences were found in terms of drowsiness during the first 48 hours, and between the third and tenth days ($p = 0.026$ in both instances in favour of the intervention). These data are detailed in *Appendix 8*, but it should be noted that given the anticipated rarity of these events, small trials are unlikely to provide unreliable estimates. No UKTIS data were available on this intervention.

Corticosteroids versus metoclopramide

Combined severity score

No combined score reported.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

One trial by Bondok and colleagues⁶⁴ compared the effectiveness of corticosteroids against metoclopramide in terms of recorded episodes of emesis. A significant improvement was observed in favour of the corticosteroid group ($p < 0.001$).

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No pregnancy outcome or side effect data were reported in the trial of Bondok and colleagues,⁶⁴ and no UKTIS data were available.

Corticosteroids against usual treatment**Combined severity score**

No combined score reported.

Nausea outcomes

Moran and Taylor¹²⁵ assessed the change in nausea symptoms in 6 out of the 25 case series patients who were treated with oral or i.v. corticosteroids for severe HG using the VAS. Data presented graphically suggested that the intensity of nausea experienced showed a pattern of improvement in the intervention group. However, no precise numerical scores were provided.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

Moran and Taylor¹²⁵ did not report any side effects and there was no difference in mean gestation at delivery or birthweight for term infants. Full data are reported in *Appendix 8*.

Summary

- Evidence available for corticosteroids was predominantly at low risk of bias (three studies^{64,93,99}), or the risk of bias was unclear/high (three studies^{58,114,116}).
- Steroids versus placebo had a trend towards improved symptoms, but results were not statistically significant.
- For steroids versus promethazine, there was no evidence of a difference in improvement by 1 week.
- Steroids were more effective at reducing vomiting episodes than Phenergan suppositories or metoclopramide.
- Overall, treatment with corticosteroids reduces the severity of symptoms, but more and larger studies are required to compare effectiveness against comparators.

Chapter 16 Clinical effectiveness: nasogastric enteral/jejunostomy feeding

Introduction

Two case series of nasogastric¹²³ and jejunostomy¹²⁶ feeding for women experiencing HG at the severe end of the spectrum were found. Both studies were judged as weak in quality. A summary of study conduct and results is depicted in *Table 20*, with safety data reported in *Appendix 8*.

Nasogastric enteral feeding

Hsu and colleagues¹²³ reported their experience of treating HG with nasogastric enteral feeding in a case study with seven women conducted at the Genesee Hospital, Rochester, New York. Enteral feeding was administered via an 8-Fr Dobbhoff nasogastric tube (Ross Products Division, Abbott Laboratories, Columbus, OH), delivering Jevity or Osmolite incrementally to meet daily caloric requirements (initial rate of 25 ml/hour, increasing as tolerated by 25 ml an hour per day). Patients were discharged once stabilised. Six women continued enteral feedings at home with nasogastric feedings discontinuing when nutritional needs were being met orally. Key outcomes of interest were the period of time (in days) after which patients were discharged and the mean duration of feedings (see *Appendix 8*).

Combined severity score

No combined scores reported.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No UKTIS data were available on this intervention.

Jejunostomy feeding

A further case series study by Saha and colleagues¹²⁶ looked at the feasibility and efficacy of feeding via jejunostomy in five women with HG compared with standard therapy at the Women and Infants Hospital, Providence, Rhode Island. Participants all displayed persistent severe nausea and vomiting plus one of the following: weight loss of > 5% of pre-pregnancy weight; ketonuria; multiple emergency room visits for dehydration; and/or inability to tolerate oral intake, despite i.v. hydration, i.v. ondansetron, i.v. ranitidine or pantoprazole, and i.v. metoclopramide. The J-tubes were placed between 12 and 26 weeks' gestation (median 14 weeks) for a mean duration of 19 weeks (range 8–28 weeks). Four J-tubes remained in place until delivery. One tube was removed at 34 weeks at the patient's request because of emotional distress and one tube fell out at 30 weeks' gestation and was not replaced. They reported that all patients had continued nausea and vomiting over the study period which required continued standard therapy in

TABLE 20 Results for nasogastric enteral/jejunostomy feeding for HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Nasogastric feeding									
Hsu 1996 ^{1,23}	The Genesee Hospital, Rochester, NY, USA	To report experiences of treating HG with nasogastric enteral feeding, case series	Seven participants (reported as individual not as mean)	Hospitalised due to intractable nausea, vomiting and weight loss, symptoms present at 1–4 weeks and had failed to respond to dietary manipulation and antiemetics	NR (SEVERE)	Enteral feeding via an 8-Fr Dobhoff nasogastric tube, of Jevity or Osmolite incrementally to meet daily caloric requirements. Initial rate of 25 ml/hour, increasing as tolerated by 25 ml/hour/day. Patients were discharged once stabilised. Six women continued enteral feedings at home. Patients were instructed to eat when they felt able. Nasogastric feedings were discontinued when nutritional needs were being met orally	NA	NR	NR
Jejunostomy									
Saha 2009 ^{7,26}	Women and Infants Hospital, Providence, RI, USA	To assess the feasibility and efficacy of surgically placed feeding jejunostomy (J-tube) in women with HG refractory to standard therapy, case series	Five patients (covering 6 pregnancies), 16.3 weeks	Persistent severe nausea and vomiting and one of the following: weight loss of > 5% of pre-pregnancy weight, ketonuria, multiple ER visits for dehydration, and/or inability to tolerate oral intake, despite i.v. hydration/ondansetron/ranitidine or pantoprazole, and metoprolamide	NR (SEVERE)	The J-tubes were placed between 12 and 26 weeks' gestation (median 14 weeks). The J-tubes were in place for a mean duration of 19 weeks (range 8–28 weeks). Four J-tubes remained in place until delivery. One tube was removed at 34 weeks at the patient's request because of emotional distress (patient 1) and one tube fell out at 30 weeks' gestation and was not replaced (patient 2)	NA	NR	All patients had continued nausea and vomiting

ER, emergency room; I, intervention; NA, not applicable; NR, not reported.

addition to J-tube feedings. Outcomes measured by the study included use of additional medication and adverse events (see *Appendix 8*).

Combined severity score

No combined scores reported.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes reported.

Safety outcomes

No UKTIS data were available on this intervention.

Summary

- Enteral feeding is an effective but extreme method of supporting women suffering from very severe symptoms as a last resort.

Chapter 17 Clinical effectiveness: gabapentin

Introduction

No controlled trials of gabapentin use in the treatment of HG were identified. The only available data were derived from one uncontrolled pilot study by Guttuso and colleagues³⁹ which enrolled seven consecutive women of 20 weeks' gestation with HG to examine gabapentin therapy. This study was classed as 'weak' by the EPHPP quality assessment tool.

Author-defined scale

The authors concluded that gabapentin appeared to be well-tolerated and may be effective in the treatment of HG but that they did not recommend the use of gabapentin in the treatment of HG until larger controlled trials have properly assessed gabapentin's efficacy in HG. This is only a very small pilot study, but a summary of study conduct and results is depicted in *Table 21*.

Nausea outcomes

No independent nausea outcomes reported.

Vomiting outcomes

No independent vomiting outcomes reported.

Retching outcomes

No independent retching outcomes.

Safety outcomes

Guttuso and colleagues³⁹ reported two serious congenital defects among seven of the subjects' infants (tethered spinal cord and hydronephrosis). Four subjects experienced mild–moderate side effects of sleepiness or dizziness when first starting gabapentin. It is important to highlight that this was a very small trial so unlikely to provide reliable estimates; however, the authors did conclude that gabapentin's safety during pregnancy needed to be further assessed. No UKTIS data were available on this intervention but full details are reported in *Appendix 8, Secondary outcome data*.

Summary

- We identified only one very small pilot trial which examined gabapentin therapy in women with HG.
- More research is needed, including monitoring of safety, in light of the reported cases of congenital anomalies among the seven exposed infants.

TABLE 21 Results for gabapentin interventions for HG

Study	Setting, location	Research question, study design	Number of participants, gestation	Severity inclusion criteria	Severity scores (reviewers' assessment)	Intervention	Comparator	Outcome assessment scale	Symptom relief outcomes
Guttuso 2010 ³⁹	Combination of inpatients and outpatients included, Buffalo, NY, USA	To perform an open-label pilot study examining the safety, tolerability and effectiveness of gabapentin in the treatment of HG, case series	Seven participants; 8 weeks	Eligible subjects needed to have severe nausea and vomiting, refractory to at least one antiemetic, causing at least 3+ ketonuria or 5% weight loss compared with the pre-pregnancy weight	All seven subjects had at least 3+ ketonuria and 5% weight loss from their pre-pregnancy weight at baseline (MODERATE-SEVERE)	Gabapentin initiated at 300 mg orally, three times daily. Every day the total dose could be increased by 300 mg (maximum of 3600 mg/day) if the subject was still experiencing nausea or emesis and was not experiencing bothersome side effects. After 14 days of therapy, gabapentin was discontinued for 2 days and could then be resumed on day 17 and for the remainder of the pregnancy, if necessary	N/A	PUQE scale	Mean reductions in nausea and emesis from baseline to days 12–14 of 80% and 94%, respectively, and 84% and 98%, respectively, from baseline to days 19–21. There was a 3 times increase in mean nausea and a seven times increase in mean emesis scores associated with discontinuing gabapentin during days 15–16

I, intervention; N/A, not applicable.

Chapter 18 Economic analysis

Introduction

This chapter has two aims: (i) to review systematically the published evidence relating to the cost-effectiveness of interventions used in the treatment of NVP/HG; and (ii) to assess the relative cost-effectiveness of such interventions from a NHS and Personal Social Services perspective. To assess cost-effectiveness, an economic model was developed in the first instance. However, due to the lack of evidence required to populate the economic model, an alternative method of economic evaluation was used. The methods and findings of the systematic review are presented first, followed by those of the economic evaluation.

Systematic review of economic evaluations

Methods

The broad methods of this systematic review were similar to those presented in *Chapter 2* and thus, only key details relevant to the economic review are given here.

Search strategy

Searches for economic studies were run on MEDLINE, EMBASE, Cumulative Index to Nursing and Allied Health Literature, PsycInfo, Allied and Complementary Medicine Database, British Nursing Index, Cochrane Central Register of Controlled Trials, Scopus, Web of Science, NHS Economic Evaluation Database and Health Economic Evaluations Database. The same terms as used for the main review were used, with the addition of health economic-related terms (included in *Appendix 9*).

Study selection

As part of the main review, titles and abstracts were screened for relevance and any potentially relevant articles were to be obtained for scrutiny against the full selection criteria by the health economics lead. The criteria were:

Study design Cost–consequence analysis, cost-effectiveness analysis, cost–benefit analysis, cost–utility analysis, cost studies.

Population Women experiencing severe nausea, vomiting and/or retching in pregnancy where recruitment to a trial took place up to 20 weeks' gestation. Owing to the difficulty in differentiating between HG and severe or intractable NVP, specific approaches were used to identify relevant populations of women (described in *Chapter 3*). Studies with mixed populations were to be included as long as data for relevant patients were extractable.

Intervention All pharmacological and non-pharmacological interventions relevant to the NHS in the community and in hospital as either an inpatient or an outpatient. The list of eligible interventions has been outlined previously in *Chapter 2*.

Comparator A no treatment group, a treatment as usual group or an alternative intervention group.

Outcome Cost-effectiveness, cost estimates, utilisation estimates, QoL estimates.

Data extraction and quality assessment strategy

Data on the following, where available, were to be extracted from included studies by one health economic reviewer and checked by another:

- study characteristics, such as study question, form of economic analysis, populations, interventions, comparators, perspective, time horizon and form of modelling used
- clinical effectiveness and cost parameters, such as effectiveness data, health state valuations (utilities), resource use data, unit cost data, price year, discounting and key assumptions; and
- results and sensitivity analyses.

Studies were to be quality assessed using the following tools as part of the data extraction process: the Consensus on Health Economic Criteria list¹³⁰ for economic evaluations and the checklist by Philips and colleagues¹³¹ for model-based analyses.

Results

From the main systematic review, 11 papers (no duplicates) were identified. None of the records were deemed relevant to this economic review and, as such, no hard copies were obtained for scrutiny against the inclusion criteria for the review. A flow diagram presenting the process of selecting studies can be found in *Figure 5*.

Discussion

No economic evaluations were found during the search for literature on the cost and cost-effectiveness of interventions used in the treatment of NVP/HG.

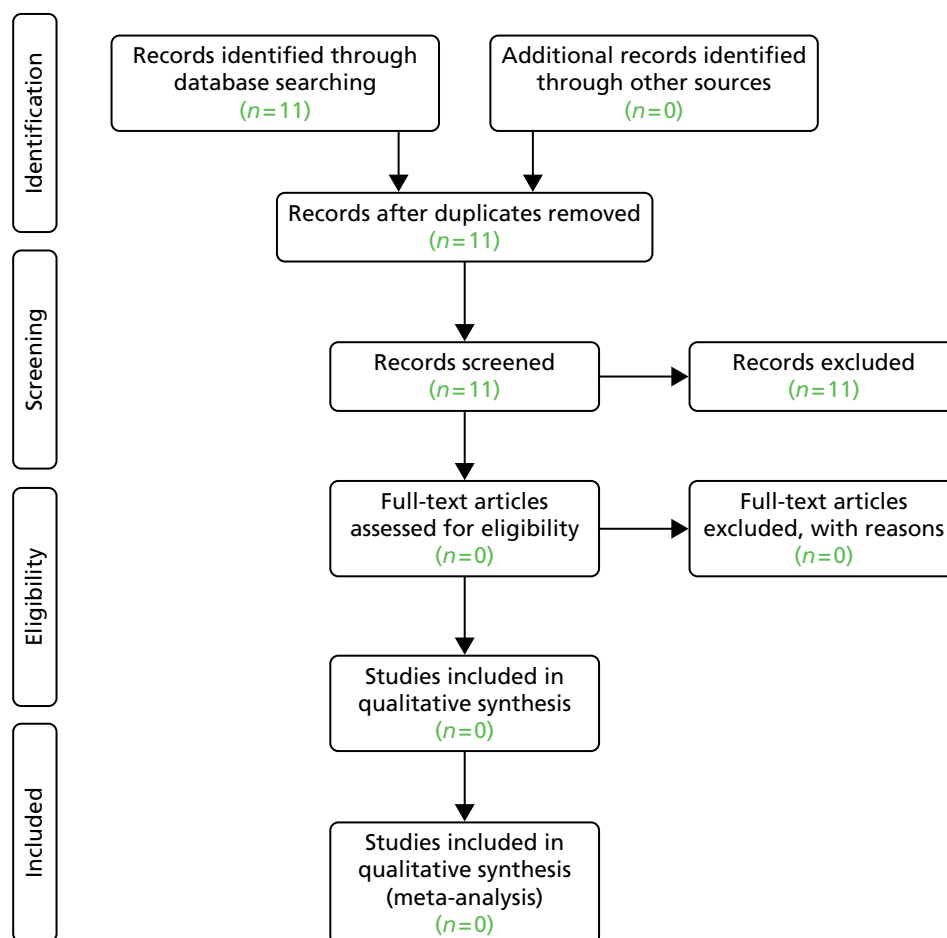


FIGURE 5 Flow diagram showing study selection for economic evaluations review.

Economic modelling

Introduction

This section provides a detailed description of the economic model developed to estimate the relative cost-effectiveness of interventions used to treat pregnant women suffering from NVP/HG. The model describes the potential care pathways experienced by patients at different levels of severity (mild, moderate and severe). A separate, identical model was created for each group, with the plan to adjust probabilities and outcomes in each model to reflect reality. Unfortunately, due to the lack of available clinical evidence it was not possible to populate the economic model. However, a model now exists for analysis should additional information become available. An overview of the key characteristics of the proposed cost-effectiveness analysis is presented in *Box 1*.

Methods

Outline of model

We initially proposed to develop an economic model to estimate costs, long-term effects and relative cost-effectiveness of the alternative interventions for NVP and HG from the perspective of the UK NHS and Personal Social Services. The model was to describe the pathways of individuals who have different severities of symptoms and have treatment initiated in different sectors of the health service (primary care, hospital outpatients and hospital inpatients). It was to cover the period of initial intervention and the costs and consequences of any subsequent outcomes including further interventions. Some of the effects of NVP and HG are short term; however, there may be some persisting impact on the mother and longer-term effects on the child. It was the intention to model the cumulative costs and quality-adjusted life-years for the mother and longer-term effects on the child (reported either in natural units or, if data allowed, quality-adjusted life-years). An outline structure of the core pathways for this proposed model is presented in *Figure 6*. The full model is available from the authors.

BOX 1 Characteristics of the cost-effectiveness analysis

Intervention: All clinically relevant interventions used to treat pregnant women suffering from NVP and HG.

Comparator: Alternative interventions used in the treatment of NVP/HG.

Population: Cohort of patients suffering from mild, moderate or severe NVP/HG and who are receiving a clinically relevant intervention in response.

Time frame: Lifetime time horizon, 3-month time cycle.

Perspective: NHS/Personal Social Services.

Effects: Any adverse events and necessary hospital treatments related to NVP/HG.

Costs: Pharmacological costs associated with medical treatment and changing medical treatment and the non-pharmacological costs associated with treating patients who have experienced a progression of symptoms or an adverse event.

Outcomes: Mortality, QoL, QALYs.

Assessment of cost-effectiveness: Cost per additional QALY gained.

QALY, quality-adjusted life-year.

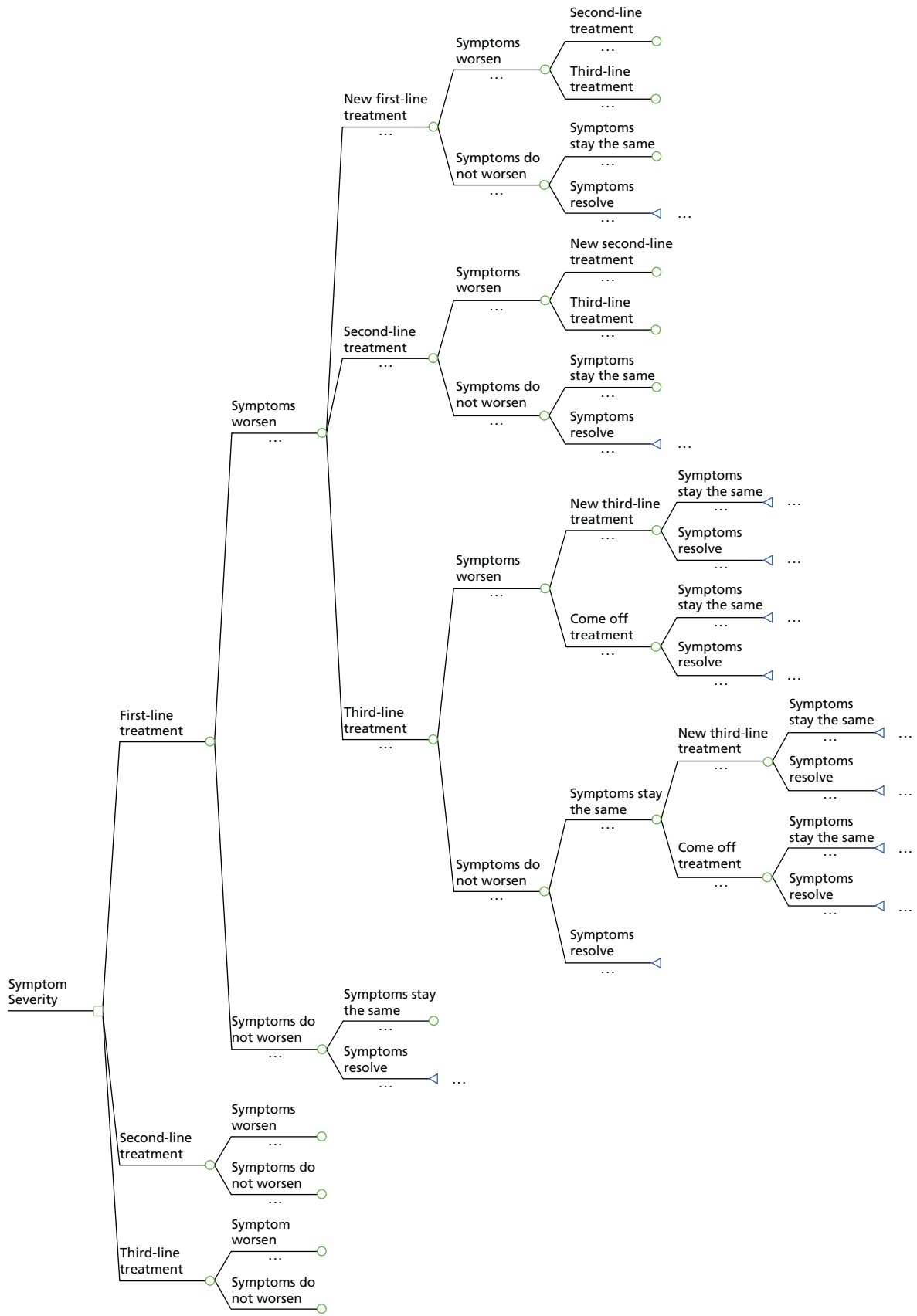


FIGURE 6 Structure of the decision model.

The economic model was built in TreeAge Pro® 2014 (TreeAge Software, Inc., Williamstown, MA, USA) to estimate the relative cost-effectiveness of the pharmacological and non-pharmacological interventions used to treat NVP/HG. The populations considered were women experiencing mild, moderate and severe nausea, vomiting and/or retching in pregnancy.

The decision tree structure can be described as follows. Depending on the severity of symptoms, patients began treatment on first-, second- or third-line treatment. The categorisation of interventions is described in detail in *Chapter 2* but can be distinguished loosely as patient-initiated treatments (first line), clinician-prescribed antiemetics (second line) and clinician-prescribed corticosteroids (third line). The categorisation of these treatments as first line, second line and third line is not medically recognised and there may be discrepancies in how clinicians describe particular interventions. However, such a hierarchy allows us to compare the relative cost-effectiveness of categories of treatments rather than the multiple individual treatments that exist, while allowing us to think through the sequence of treatments that are likely to be relevant to patients at the different levels of severity of symptoms.

Beginning at 'first-line treatment', symptoms may worsen or not worsen. If symptoms worsened, patients moved to a 'new first-line treatment', 'second-line treatment' or 'third-line treatment'. From each of these points, symptoms may again worsen or not worsen. If symptoms worsened, patients moved to the next hierarchical treatment strategy. This sequence was followed until patients reached 'third-line treatment', at which point symptoms may worsen or not worsen. If symptoms worsened, patients moved to a 'new third-line treatment' or 'came off treatment', at which point symptoms could either 'stay the same' or 'resolve' (terminal nodes). Beginning at the same starting point ('first-line treatment'), if symptoms did not worsen, symptoms could 'stay the same' or 'resolve' (terminal node). If 'symptoms stayed the same', the pathways described at the same point as symptoms worsening were replicated.

For those patients beginning at 'second-line' and third-line' treatment, the subsequent care pathways were broadly similar to those previously described. However, the model assumes that those patients starting at second- and third-line treatments are unable to progress to less severe treatment strategies. Therefore, the care pathways are progressively shorter as we move through the hierarchy of treatments. The decision model allows one to assess costs and effects across the three alternative treatment strategies and to assess the relative cost-effectiveness of interventions used to treat NVP/HG.

Economic evaluation

Introduction

Unfortunately, as described in previous chapters, the evidence base supporting alternative treatments was not sufficiently robust to support an informative model. Therefore, the focus of the economic component was to estimate the cost of delivering each of the interventions and then use these data in a disaggregated form of economic evaluation. The differences between interventions, in terms of resource use (costs) and natural and clinical measures of effectiveness are presented. Such an approach serves to highlight the choices and trade-offs between the various treatments. Nonetheless, any decision based on this approach is made using an implicit (rather than explicit) synthesis of the available data. These differences between interventions are presented for each of the comparisons presented in *Chapters 4–17*.

The cost data are used to estimate the implied value for the benefits of treatment should a decision be made to adopt one treatment over another. The approach relies on the conditions required for an efficient allocation of resources. When resources are efficiently allocated the ratio of marginal costs to marginal benefits for all interventions a to n must be equal, that is:

$$\frac{MC_a}{MB_a} = \frac{MC_b}{MB_b} = \frac{MC_c}{MB_c} = \dots = \frac{MC_n}{MB_n} \quad (1)$$

Rearranging the above and simplifying shows that when there is an efficient allocation of resources the ratio of costs is equal to the ratio of benefits, such that:

$$\frac{MC_a}{MC_b} = \frac{MB_a}{MB_b} \quad (2)$$

Although benefits may not be clearly known, it is possible to use information about the costs of interventions a and b to imply how much more effective a needs to be compared with b to be considered efficient. For example, if the ratio of the costs of a to b is 1.2 : 1 then this means that a is 20% more costly than b and for a to be considered a worthwhile use of resources compared with b it should provide at least 20% more benefits. This implied value is then compared with the evidence on effectiveness reported in Chapters 4–17 to inform judgements made on the relative cost-effectiveness of treatments. This section provides a detailed description of the costs of interventions used in the treatment of NVP/HG.

Estimation of costs of interventions

In order to inform the economic evaluation, a detailed cost analysis of interventions used in the treatment of NVP/HG was performed. The interventions considered were categorised as follows: (1) patient-initiated first-line interventions; (2) clinician-prescribed second-line interventions; and (3) clinician-prescribed third-line interventions. The cost of interventions includes both the pharmacological (medication) and non-pharmacological (GP visits, outpatient care, inpatient care, medical tests and management) costs of treatment. The weekly pharmacological costs of clinically relevant interventions (as advised by clinical experts) are listed in *Table 22*, whereas non-pharmacological costs are listed in *Table 23*. A full table of interventions included in the analysis can be seen in *Appendix 10*. These additional treatments were included for completeness but are rarely prescribed in the NHS.

The costs of medication were sourced from the *British National Formulary 2014*.¹³² For all medications, the non-proprietary costs were sought in the first instance. However, if these were unavailable the patented drug costs were used. Where available, the costs of medications which have been incorporated into a tablet or capsule were obtained. Clinical experts in the study team were able to advise on which medications are also commonly administered by i.v. or intramuscular (i.m.) injection and through suppositories, and the relevant costs were included. The overall cost of medication and the average daily dose were used to calculate a low- and high-estimate weekly cost of medication, unless otherwise stated. Where average daily dose was presented as a minimum and maximum, the minimum was used to calculate the low-estimate cost and the maximum was used to calculate the high-estimate cost. Where average daily dose was not presented over a range, costs have been included under high-estimate weekly cost. These drug costs and the recommended daily doses are included in *Appendix 10*.

Non-pharmacological costs included the unit costs of health and social care services, as well as any tests and medication administered, in a primary and secondary care setting. Clinical experts advised on the tests and treatments which are commonly given to women suffering from NV/HG in each setting. Standard sources such as the NHS Reference Healthcare Research Group tariffs 2012–13,¹³³ the *British National Formulary 2014*¹³² for medications and the Personal Social Services Research Unit *Unit Costs of Health and Social Care 2013*¹³⁴ were used to obtain these costs.

Estimation of total cost of care

In order to estimate the total cost of interventions, the pharmacological and non-pharmacological costs were combined together in informative packages of care. All likely scenarios experienced by patients suffering from NVP/HG, based on the advice of clinical experts, were included. These scenarios ranged from treatments initiated before consultation with a medical professional to treatments prescribed in primary and secondary care. The flow chart presented in *Figure 1* shows the treatments that patients are likely to receive at each level of care and the scenarios presented in this section are based on these care pathways. The costs of managing patients in each setting, as well as the costs of treatment, were used to estimate the overall cost of care. For patient-initiated interventions and medication prescribed in a

TABLE 22 Weekly costs of pharmacological interventions

Preparation	Patient-initiated first-line intervention	Low-estimate weekly cost	High-estimate weekly cost
Tablets	Vitamin B6: pyridoxine hydrochloride (non-proprietary)	£0.12	£2.59
Tablets	Vitamin B12: cyanocobalamin (non-proprietary)	£0.87	£2.62
Solution	Ginger (FortiCare)	NE	£2.21 (assumes 125 ml per week) ^a
Physical therapy	Acupressure/acupuncture	NE	First appointment: £50–70 Subsequent appointments: £35–50 ^a
Hypnosis	Hypnotherapy	NE	£50–90 for a private hypnotherapy session ^a
Preparation	Clinician-prescribed second-line interventions	Low-estimate weekly cost	High-estimate weekly cost
Antihistamines			
Tablets	Cyclizine (non-proprietary)	£0.74	£2.22
i.v./i.m. injection	Cyclizine: Valoid® (Amdipharm Mercury Company Ltd)	NE	£13.65
Tablets	Chlorpromazine (non-proprietary)	NE	£1.55
i.v./i.m. injection	Chlorpromazine (non-proprietary)	£12.60	£33.60
Dopamine antagonists			
Tablets	Promethazine: Phenergan	£0.74	£2.22
i.v./i.m. injection	Promethazine (non-proprietary)	£0.68 (daily) ^a	£1.20 (daily) ^a
Tablets	Prochlorperazine (non-proprietary)	£0.47	£1.42
i.v./i.m. injection	Prochlorperazine (non-proprietary)	NE	£0.52 (daily) ^a
Tablets	Domperidone (non-proprietary)	£0.39	£1.17
Tablets	Metoclopramide (non-proprietary)	£0.22	£0.66
i.v./i.m. injection	Metoclopramide (non-proprietary)	£2.24	£6.72
Serotonin antagonists			
Tablets	Ondansetron (non-proprietary)	NE	£8.69 (daily) ^a
i.v./i.m. injection	Ondansetron (non-proprietary)	NE	£1.00 (daily) ^a
Preparation	Clinician-prescribed third-line interventions	Low-estimate weekly cost	High-estimate weekly cost
Corticosteroids			
i.v./i.m. injection	Methylprednisolone: Solu-Medrone® (Pharmacia)	NE	£17.30 (daily) ^a
Tablets	Prednisone: Lodotra® (Napp Pharmaceuticals Limited)	£12.46	£24.92
Tablets	Prednisolone (non-proprietary)	£0.67	£1.33
i.v./i.m. injection	Hydrocortisone: Efcortisol® (Amdipharm Mercury Company Ltd)	£3.24 (daily) ^a	£19.56 (daily) ^a
NE, not estimated.			
^a All costs are weekly, unless otherwise stated.			

TABLE 23 Non-pharmacological costs

Primary care	Costs
GP consultation	
Per surgical consultation lasting 11.7 minutes	£45
Ketone Reagent Strips: Ketosis (Bayer Diabetes Care) (50 pack)	£2.50
Secondary care	
Outpatient	
Obstetrics	£122
Ultrasound scan in obstetrics (less than 20 minutes)	£38
Urinary test (urine culture)	£8.51
Ketone Reagent Strips: Ketosis (Bayer Diabetes Care) (50 pack)	£2.50
Liver function test	£6.80
Thyroid function test	£13.55
Glucose	£2.96
Urea and electrolytes	£5.84
Full blood count	£4.94
Sodium chloride i.v. infusion	£1.29 per litre
Potassium chloride concentrate, sterile (non-proprietary) (10-ml ampoule)	£0.48
Cost of administering the fluid	£50.42 per day
Thiamine supplements	£0.05 per day/£0.35 per week (low estimate) £0.08 per day/£0.56 per week (high estimate)
Inpatient	
Non-elective inpatient (long stay) excess bed-days	£265
Urinary test (urine culture)	£8.51
Ketone Reagent Strips: Ketosis (Bayer Diabetes Care) (50 pack)	£2.50
Liver function test	£6.80
Thyroid function test	£13.55
Glucose	£2.96
Urea and electrolytes	£5.84
Full blood count	£4.94
Sodium chloride i.v. infusion	£1.29 per litre
Potassium chloride concentrate, sterile (non-proprietary) (10-ml ampoule)	£0.48
Cost of administering the fluid	£50.42 per day
Pabrinex® (Archimedes Pharma) (10 ml)	£2.25 per week
Treatment for thromboembolism:	
Clexane® (Sanofi-Aventis) (40 mg)	£3.03 per day
Tinzaparin (Innohep®, Leo Laboratories) (4500 units)	£3.56 per day
Fragmin® (Pfizer) (5000 units)	£2.56 per day
	£9.15 per day

Bold text represents the point at which individual drug costs are added to calculate the total daily cost of treatment for thromboembolism.

primary care setting, costs were estimated on a weekly basis. For all other interventions administered in a secondary care setting, costs were estimated according to length of stay. The packages of care are described in scenarios 1–7 below, and total costs for all clinically relevant interventions, as advised by clinical experts, are displayed in *Tables 24–30*. The estimation of total costs for all interventions is included in *Appendix 10*.

Packages of care

Scenario 1

Initially, we assumed that the patient does not visit her GP. Rather, she initiates treatment herself. The patient may seek advice from a community midwife or GP at this stage. However, costs are for the medication or treatment costs alone. These costs are outlined in *Table 24*.

Scenario 2

We assumed that the patient visits her GP and receives a urinary test. The GP decides to reassure the patient and advises her to remain on one of the patient-initiated first-line interventions. It was judged unlikely that the GP would advise on acupressure/acupuncture or hypnosis at this stage as these therapies are not routinely available on the NHS. Costs are displayed in *Table 25*.

TABLE 24 Cost of patient-initiated first-line interventions

Pharmacological preparation	Patient-initiated first-line intervention	Total low-estimate weekly cost	Total high-estimate weekly cost
Tablets	Vitamin B6: pyridoxine hydrochloride (non-proprietary)	£0.12	£2.59
Tablets	Vitamin B12: cyanocobalamin (non-proprietary)	£0.87	£2.62
Solution	Ginger (FortiCare)	NE	£2.21 (assuming 125 ml per week)
Physical therapy	Acupressure/acupuncture	£35	£50
Hypnosis	Hypnotherapy	£50	£90
NE, not estimated.			

TABLE 25 Cost of patient-initiated first-line interventions following a GP visit

Pharmacological preparation	Patient-initiated first-line intervention	Total low-estimate weekly cost	Total high-estimate weekly cost	GP clinic consultation	Urine ketones strip	Total
Tablets	Vitamin B6: pyridoxine hydrochloride (non-proprietary)	£0.12	£2.59	£45	£0.05	£45.17–47.64
Tablets	Vitamin B12: cyanocobalamin (non-proprietary)	£0.87	£2.62	£45	£0.05	£45.92–47.67
Solution	Ginger (FortiCare)	NE	£2.21 (assuming 125 ml per week)	£45	£0.05	£47.26
NE, not estimated.						

Scenario 3

We assumed that the patient visits her GP and receives a urinary test. Symptoms are severe enough that the GP prescribes one of the clinician-prescribed second-line interventions (oral antiemetic only).

Costs are displayed in *Table 26*.

Scenario 4

If the patient is in a serious enough condition that they cannot be managed in primary care, the GP may refer the woman to secondary care (or alternatively, women may refer themselves to hospital maternity services). The following costs reflect those costs that might be incurred in a secondary care setting.

In *Table 27* we include the costs of the woman attending hospital as a 'day case', receiving the relevant outpatient tests, receiving an antiemetic and then being discharged. Included also are the daily cost of receiving a thiamine supplement (high estimate) and the cost of receiving 3 l (in severe cases) of sodium chloride i.v. infusion along with the appropriate amount of potassium chloride. All costs are calculated on a daily basis.

Scenario 5

In very severe cases, or where the patient refuses outpatient treatment, or if outpatient treatment fails (i.e. the patient feels as unwell at the end of outpatient treatment), the woman may be admitted as an inpatient. In *Table 28* the assumption is that the woman is admitted as an inpatient for 2 days (based on expert opinion). In this scenario, it is assumed that the woman would have received a one-off ultrasound scan prior to being admitted as an inpatient and thus, this cost is not included. Based on expert clinical advice, we have assumed that the patient receives all relevant tests on day of admission and day of discharge, medication used in the prevention of venous thromboembolism, i.v. infusion of 3 l of sodium chloride solution every day along with the appropriate amount of potassium chloride, i.v. thiamine (Pabrinex, 10 ml) once a week, and appropriate antiemetics (as advised by clinical experts) (no steroids) over the course of the 2-day admission before being discharged.

TABLE 26 Cost of clinician-prescribed second-line interventions following a GP visit

Pharmacological preparation	Clinician-prescribed second-line intervention	Total low-estimate weekly cost	Total high-estimate weekly cost	GP clinic consultation	Urine ketones strip	Total
Antihistamines						
Tablets	Cyclizine (non-proprietary)	£0.74	£2.22	£45	£0.05	£45.79–47.27
Tablets	Chlorpromazine (non-proprietary)	NE	£1.55	£45	£0.05	£46.60
Dopamine antagonists						
Tablets	Domperidone (non-proprietary)	£0.39	£1.17	£45	£0.05	£45.44–46.22
Tablets	Metoclopramide (non-proprietary)	£0.22	£0.66	£45	£0.05	£45.27–45.71
Tablets	Prochlorperazine (non-proprietary)	£0.47	£1.42	£45	£0.05	£45.52–46.47
Tablets	Promethazine: Phenergan	£0.74	£2.22	£45	£0.05	£45.79–47.27
Serotonin antagonists						
Tablets	Ondansetron (non-proprietary)	NE	£60.83	£45	£0.05	£105.88

NE, not estimated.

Scenario 6

In this scenario, it is assumed that the patient has not responded to the single antiemetic prescribed while admitted as an inpatient and so has been admitted as an inpatient for 2 days (based on expert opinion) with two antiemetics prescribed. Based on expert clinical advice, we have assumed that the patient receives all relevant tests on day of admission and day of discharge, i.v. infusion of 3 l of sodium chloride solution every day along with the appropriate amount of potassium chloride, medication used in the prevention of venous thromboembolism, i.v. thiamine (Pabrinex, 10 ml) once a week, and a combination of two antiemetics (no steroids) over the course of the 2-day admission before being discharged. Owing to the minimal difference in cost between antiemetics, the low- and high-estimate costs represent combinations of the two cheapest and two most expensive medications (based on high-estimate daily cost). These costs are displayed in *Table 29*.

Scenario 7

In this scenario, it is assumed that the patient has so far not responded to any antiemetics prescribed while admitted as an inpatient. In the following table of costs (see *Table 30*), the assumption is that the woman has been admitted as an inpatient for 5 days (based on expert opinion) to follow a regime of multiple antiemetics and steroids. Up until this point the woman has experienced (a) weight loss, (b) failed second-line interventions, or (c) difficulty coping. As the condition of the patient is so severe, it is assumed that she receives all relevant tests on each day of admission (following the guidance of clinical experts). It is assumed that i.v. infusion of 3 l of sodium chloride solution is given every day along with the appropriate amount of potassium chloride, medication used in the prevention of venous thromboembolism is given for each night of stay, i.v. thiamine (Pabrinex, 10 ml) is given once over the course of the admission, and a combination of three antiemetics is given to the patient for the first 3 days before a steroid is provided in days 4 and 5, before discharge. Owing to the minimal difference in cost between antiemetics, the low- and high-estimate costs represent combinations of the three cheapest and three most expensive medications (based on high-estimate daily cost). Similarly, there is little difference in cost between steroids and so, the low- and high-estimate costs are representative of packages including the cheapest and most expensive steroid (based on high-estimate daily cost). These costs are displayed in *Table 30*.

The packages of care presented above are representative scenarios that combine the pharmacological and non-pharmacological costs of interventions used in the treatment of patients suffering from NVP and HG. If symptoms have not resolved following scenario 7, extreme measures such as enteral or parenteral nutrition may sometimes be used. However, this is quite rare and the costs can be significant. Kilonzo estimated the mean cost of treatment for standard parenteral nutrition to be £337 among a group of patients in intensive care units and high-dependency units for ≥ 48 hours, with gastrointestinal failure and requiring parenteral nutrition (Kilonzo M, University of Aberdeen, 2014, personal communication). Enteral and parenteral feeding is a last-resort therapy among women suffering from NVP/HG and would usually only ever be used as a last resort.

The packages of care indicate that the costs of treatment increase rapidly as women move through the hierarchy of treatments. In scenario 1, we see that in less severe cases, where women initiate treatment themselves, costs may be as low as £0.12 per week. Alternative patient-initiated treatment strategies such as acupressure/acupuncture or hypnosis are significantly more expensive, but may only be required on a one-off basis. If symptoms are persistent, it is likely that the woman would seek the advice of a clinician and begin to take oral prescribed medication. In scenarios 2 and 3, we see the cost of attending a GP and taking oral medication for 1 week. Following this first week, costs would revert back to the cost of medication alone, which is small in comparison with the cost of attending the GP, provided no further consultations are required. In scenario 4, we see the costs that would be incurred if the woman was to be referred to hospital as a 'day case'. This is the woman's first point of contact with secondary care services, and a number of tests, including an ultrasound scan and relevant bloods, are likely to be done at this stage. When we factor in the costs of hospital care and medication that is likely to be administered at this stage, costs range from £286 in the least costly scenario to £295 in the most costly (based on

TABLE 27 Cost of clinician-prescribed second-line interventions if attending hospital as a 'day case'

Pharmacological preparation	Clinician-prescribed second-line intervention	Total low-estimate daily cost (£)	Total high-estimate daily cost (£)	Obstetrics unit (£)	Ultrasound scan (< 20 minutes) (£)	Urinary test (urine culture) (£)
Antihistamines						
Tablets	Cyclizine (non-proprietary)	0.11	0.32	122	38	8.51
i.v./i.m. injection	Cyclizine: Valoid	NE	1.95	122	38	8.51
Tablets	Chlorpromazine (non-proprietary)	NE	0.22	122	38	8.51
i.v./i.m. injection	Chlorpromazine (non-proprietary)	1.80	4.80	122	38	8.51
Dopamine antagonists						
Tablets	Promethazine: Phenergan	0.11	0.32	122	38	8.51
i.v./i.m. injection	Promethazine (non-proprietary)	0.68	1.20	122	38	8.51
Tablets	Prochlorperazine (non-proprietary)	0.07	0.20	122	38	8.51
i.v./i.m. injection	Prochlorperazine (non-proprietary)	NE	0.52	122	38	8.51
Tablets	Domperidone (non-proprietary)	0.06	0.17	122	38	8.51
Tablets	Metoclopramide (non-proprietary)	0.03	0.09	122	38	8.51
i.v./i.m. injection	Metoclopramide (non-proprietary)	0.32	0.96	122	38	8.51
Serotonin antagonists						
Tablets	Ondansetron (non-proprietary)	NE	8.69	122	38	8.51
i.v./i.m. injection	Ondansetron (non-proprietary)	NE	1.00	122	38	8.51
NE, not estimated.						

Urine ketones strip (£)	Liver function test (£)	Thyroid function test (£)	Glucose (£)	Urea and electrolytes (£)	Full blood count (£)	Thiamine supplement (£)	Normal saline + a proportional amount of potassium chloride + cost of administering the fluid (£)	Total (£)
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.93–286.14
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	287.77
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.04
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	287.62–290.62
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.93–286.14
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.50–287.02
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.89–286.02
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.34
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.88–285.99
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.85–285.91
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.14–286.78
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	294.51
0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.82

TABLE 28 Cost of clinician-prescribed second-line interventions if admitted as an inpatient

Pharmacological preparation	Clinician-prescribed second-line interventions	Total low-estimate daily cost × 2 (£)	Total high-estimate daily cost × 2 (£)	Inpatient excess bed-days × 2 (£)	Urinary test (urine culture) × 2 (£)	Urine ketones strip × 2 (£)	Liver function test × 2 (£)
Antihistamines							
i.v./i.m. injection	Cyclizine: Valoid	NE	3.90	530	17.02	0.10	13.60
Tablets	Chlorpromazine (non-proprietary)	NE	0.44	530	17.02	0.10	13.60
i.v./i.m. injection	Chlorpromazine (non-proprietary)	3.60	9.60	530	17.02	0.10	13.60
Dopamine antagonists							
i.v./i.m. injection	Metoclopramide (non-proprietary)	0.64	1.92	530	17.02	0.10	13.60
Tablets	Prochlorperazine (non-proprietary)	0.13	0.41	530	17.02	0.10	13.60
i.v./i.m. injection	Prochlorperazine (non-proprietary)	NE	1.04	530	17.02	0.10	13.60
Tablets	Domperidone (non-proprietary)	0.11	0.33	530	17.02	0.10	13.60
Serotonin antagonists							
i.v./i.m. injection	Ondansetron (non-proprietary)	NE	2.00	530	17.02	0.10	13.60
NE, not estimated.							

TABLE 29 Cost of clinician-prescribed second-line interventions × 2 if admitted as an inpatient

Clinician-prescribed second-line intervention	Total daily cost (two antiemetics) × 2 (£)	Inpatient excess bed-days × 2 (£)	Urinary test (urine culture) × 2 (£)	Urine ketones strip × 2 (£)	Liver function test × 2 (£)	Thyroid function test × 2 (£)
Least costly	0.50	530	17.02	0.10	13.60	27.10
Most costly combination	50.70	530	17.02	0.10	13.60	27.10

TABLE 30 Cost of clinician-prescribed third-line interventions if admitted as an inpatient

Clinician-prescribed third-line intervention	Total daily cost (three antiemetics) × 3 (£)	Total daily cost (corticosteroid) × 2 (£)	Inpatient excess bed-days × 5 (£)	Urinary test (urine culture) × 5 (£)	Urine ketones strip × 5 (£)	Liver function test × 5 (£)	Thyroid function test × 5 (£)
Least costly combination	1.26	0.38 (prednisolone tablets)	1325	42.55	0.25	34.00	67.75
Most costly combination	82.86	39.12 (hydrocortisone injection)	1325	42.55	0.25	34.00	67.75

Thyroid function test × 2 (£)	Glucose × 2 (£)	Urea and electrolytes × 2 (£)	Full blood count × 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) × 1 (£)	Normal saline + a proportional amount of potassium chloride × 2 + cost of administering the fluid over 2 days (£)	Total (£)
27.10	5.92	11.68	9.88	9.15	2.25	166.18	796.78
27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.32
27.10	5.92	11.68	9.88	9.15	2.25	166.18	796.48–802.48
27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.52–794.80
27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.01–793.29
27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.92
27.10	5.92	11.68	9.88	9.15	2.25	166.18	792.99–793.21
27.10	5.92	11.68	9.88	9.15	2.25	166.18	794.88

Glucose × 2 (£)	Urea and electrolytes × 2 (£)	Full blood count × 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) × 1 (£)	Normal saline + a proportional amount of potassium chloride × 2 + cost of administering the fluid over 2 days (£)	Total (£)
5.92	11.68	9.88	9.15	2.25	166.18	793.38
5.92	11.68	9.88	9.15	2.25	166.18	843.58

Glucose × 5 (£)	Urea and electrolytes × 5 (£)	Full blood count × 5 (£)	Treatment for thromboembolism (4-night stay) (£)	Thiamine supplement (Pabrinex) × 1 (£)	Normal saline + a proportional amount of potassium chloride × 5 + cost of administering the fluid over 5 days (£)	Total (£)
14.80	29.20	24.70	36.60	2.25	415.45	1994.19
14.80	29.20	24.70	36.60	2.25	415.45	2114.53

high-estimate costs). Again, the cost of managing the woman in this setting is the main cost driver, with medication costs becoming less significant in comparison. Finally, scenarios 5–7 are representative of scenarios where the woman has been admitted to hospital as an inpatient due to the extremity of her symptoms. This is the point at which costs escalate rapidly, largely driven by length of stay. This is reflected in the fact that there is very little difference in the total cost of care between scenarios 5 and 6, despite antiemetics being doubled in the latter scenario. Similarly, in the most severe cases, where women are admitted for a 5-day period, < 6% of the total cost of care is accounted for by the medication costs.

As we have seen, the cost of medication used in the treatment of NVP/HG is less important when compared with the cost of managing women with the condition in a primary and secondary care setting. One issue which may be of importance to decision-makers is whether it is more economical to manage women with NVP/HG as day cases or as inpatients. In *Table 31*, the cost of first-, second- and third-line interventions is omitted and the focus is placed solely on the hospital management costs. The cost of managing women as day cases on 2 separate days is presented alongside the cost of managing women as inpatients over a 2-day period. All treatment costs relevant to each scenario are presented.

The difference in cost between inpatient management and day case management is largely driven by the fact that inpatient bed-days are substantially more expensive than the obstetrics unit day case costs. The only other major cost differentials are that an ultrasound scan is likely to initially be carried out on day case patients, whereas it is assumed that patients would have received a scan prior to being admitted as an inpatient. Additionally, clinical experts have advised that inpatients would require prophylactic treatment to prevent deep-vein thrombosis and pulmonary embolism, whereas this cost is not incurred when treating day case patients. The total cost difference is £259, with inpatient treatment significantly more expensive than day case treatment in the 2-day scenario presented in *Table 31*.

Methods

The total cost data were used to estimate the implied value for the benefits of treatment should a decision be made to adopt one treatment over another. Within each package of care, the ratio of the cost of one treatment to another was calculated, with the results informing us as to the increase in benefits that the more expensive option would need to provide in order to be considered a worthwhile use of resources. Benefits of treatment are not clearly known, but these results were used to imply how much more effective one treatment needs to be compared with another to be considered efficient. The implied value was then used to compare with the evidence on effectiveness reported in *Chapters 4–17*, in a disaggregated form of economic evaluation. Each comparison presented in *Chapters 4–17* is included. The implied value for the benefits of treatments is presented in the subsequent section.

Results

Implied value for the benefits of treatment

Within each package of care, the ratio of the cost of one treatment to another was calculated. The size of the ratio was indicative of the increase in benefits that the more costly option would need to provide in order to be considered efficient compared with the less costly option. Effect sizes were categorised according to Grimshaw and colleagues'¹³⁵ review of guideline dissemination and implementation strategies and, in particular, the description of the size of effect for process dichotomous measures. The effects sizes were categorised as follows:

- 'small' to describe effect sizes $\leq 5\%$
- 'modest' to describe effect sizes $> 5\%$ and $\leq 10\%$
- 'moderate' to describe effect sizes $> 10\%$ and $\leq 20\%$
- 'large' to describe effect sizes $> 20\%$.

TABLE 31 Cost of day case management compared with inpatient management

Day case	244	38	17.02	0.10	13.60	27.10	5.92	11.68	9.88	0.16	166.18	533.64
	Obstetrics unit x 2 (£)	Ultrasound scan (<20 minutes) x 1 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Thiamine supplement x 2 (£)	Normal saline and a proportional amount of potassium chloride and cost of administering the fluid on 2 days (£)	Total (£)
Inpatient	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	792.88
	Inpatient excess bed-days x 2 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) (£)	Normal saline and a proportional amount of potassium chloride and cost of administering the fluid over 2 days (£)	Total (£)

Cost comparisons for all clinically relevant interventions in the UK, as advised by clinical experts, were based on the high-estimate costs and are presented in *Tables 32–39*. Unfortunately, in a large number of cases, while a comparison of costs could be made between interventions within each package of care, evidence on effect was unavailable. Cost comparisons for all interventions included in the analysis can be seen in *Appendix 10*.

For patient-initiated first-line interventions, the ratio of the cost of vitamin B6 to ginger was 1.2 : 1. Therefore, in order to be considered a worthwhile use of resources, vitamin B6 would need to provide at least 20% more in benefits than ginger. It is feasible that such a difference in effect could exist. However, there was no evidence to support a difference of this magnitude. The ratio of the cost of acupressure/acupuncture to both vitamin B6 and ginger is extremely large and it is inconceivable that an equally large difference in effectiveness could exist to justify this. The evidence on effect showed no significant difference between the interventions, although the evidence was limited. Finally, ginger, acupressure/acupuncture and vitamin B6 were all compared with placebo. Although an implied valuation was not assessable, evidence on effect showed that all three treatments looked promising in reducing symptoms compared with placebo, but the findings were not conclusive. With respect to ginger and vitamin B6 the cost of these therapies might be considered modest and were quite similar. From an economics perspective, this suggests that we are indifferent about which is used as a first treatment.

For patient-initiated first-line interventions following a GP visit, the ratio of the cost of vitamin B6 to ginger was 1.008 : 1. Therefore, vitamin B6 would only need to provide 0.8% more in benefits than ginger in

TABLE 32 Cost comparisons of patient-initiated first-line interventions

Comparison	Implied valuation	Effect size	Evidence on effect
Vitamin B6 : ginger	1.2 : 1	Moderate	No evidence of a difference between groups
Vitamin B12 : vitamin B6	1.01 : 1	Small	Unknown
Vitamin B12 : ginger	1.2 : 1	Moderate	Unknown
Acupressure/acupuncture: vitamin B12	19.1 : 1	Large	Unknown
Acupressure/acupuncture : vitamin B6	19.3 : 1	Large	No evidence of a difference between groups
Acupressure/acupuncture : ginger	22.6 : 1	Large	Ginger looks promising in reducing symptoms when compared with acupressure but findings are not conclusive
Hypnotherapy : acupressure/acupuncture	1.8 : 1	Large	Unknown
Hypnotherapy : vitamin B12	34.4 : 1	Large	Unknown
Hypnotherapy : vitamin B6	34.7 : 1	Large	Unknown
Hypnotherapy : ginger	40.7 : 1	Large	Unknown
Ginger : placebo	Not assessable	Not assessable	Ginger looks more effective when compared with placebo. Additional cost of ginger is small
Acupressure/acupuncture : placebo	Not assessable	Not assessable	Acupressure looks promising in reducing symptoms when compared with placebo in a small number of studies, while the rest show no difference between the groups
Vitamin B6 : placebo	Not assessable	Not assessable	Vitamin B6 looks more effective when compared with placebo. Additional cost of vitamin B6 is small

TABLE 33 Cost comparisons of patient-initiated first-line interventions following a GP visit

Comparison	Implied valuation	Effect size	Evidence on effect
Vitamin B6 : ginger	1.008 : 1	Small	No evidence of a difference in effect
Vitamin B12 : vitamin B6	1.0006 : 1	Small	Unknown
Vitamin B12 : ginger	1.009 : 1	Small	Unknown
Ginger : placebo	Not assessable	Not assessable	Ginger looks promising in reducing symptoms when compared with placebo, but findings are not conclusive
Vitamin B6 : placebo	Not assessable	Not assessable	Vitamin B6 looks promising in reducing symptoms when compared with placebo, but findings are not conclusive

TABLE 34 Cost comparisons of clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit

Comparison	Implied valuation	Effect size	Evidence on effect
Domperidone : metoclopramide	1.01 : 1	Small	Unknown
Prochlorperazine : domperidone	1.005 : 1	Small	Unknown
Prochlorperazine : metoclopramide	1.02 : 1	Small	Unknown
Chlorpromazine : prochlorperazine	1.003 : 1	Small	Unknown
Chlorpromazine : domperidone	1.008 : 1	Small	Unknown
Chlorpromazine : metoclopramide	1.02 : 1	Small	Unknown
Cyclizine : chlorpromazine	1.01 : 1	Small	Unknown
Cyclizine : prochlorperazine	1.02 : 1	Small	Unknown
Cyclizine : domperidone	1.02 : 1	Small	Unknown
Cyclizine : metoclopramide	1.03 : 1	Small	Unknown
Promethazine : cyclizine	1 : 1	Small	Unknown
Promethazine : chlorpromazine	1.01 : 1	Small	Unknown
Promethazine : prochlorperazine	1.02 : 1	Small	Unknown
Promethazine : domperidone	1.02 : 1	Small	Unknown
Promethazine : metoclopramide	1.03 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Ondansetron : promethazine	2.2 : 1	Large	Unknown
Ondansetron : cyclizine	2.2 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no evidence of significant difference in effects
Ondansetron : chlorpromazine	2.3 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no evidence of significant difference in effects
Ondansetron : prochlorperazine	2.3 : 1	Large	Unknown
Ondansetron : domperidone	2.3 : 1	Large	Unknown
Ondansetron : metoclopramide	2.3 : 1	Large	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Antihistamines : placebo	Not assessable	Not assessable	Antihistamines appear to be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required

TABLE 35 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case'

Comparison	Implied valuation	Effect size	Evidence on effect
Domperidone (tablets) : metoclopramide (tablets)	1.0003 : 1	Small	Unknown
Prochlorperazine (tablets) : domperidone (tablets)	1.0001 : 1	Small	Unknown
Prochlorperazine (tablets) : metoclopramide (tablets)	1.0004 : 1	Small	Unknown
Chlorpromazine (tablets) : prochlorperazine (tablets)	1.00007 : 1	Small	Unknown
Chlorpromazine (tablets) : domperidone (tablets)	1.0002 : 1	Small	Unknown
Chlorpromazine (tablets) : metoclopramide (tablets)	1.0005 : 1	Small	Unknown
Cyclizine (tablets) : chlorpromazine (tablets)	1.0003 : 1	Small	Unknown
Cyclizine (tablets) : prochlorperazine (tablets)	1.0004 : 1	Small	Unknown
Cyclizine (tablets) : domperidone (tablets)	1.0005 : 1	Small	Unknown
Cyclizine (tablets) : metoclopramide (tablets)	1.0008 : 1	Small	Unknown
Promethazine (tablets) : cyclizine (tablets)	1 : 1	Small	Unknown
Promethazine (tablets) : chlorpromazine (tablets)	1.0003 : 1	Small	Unknown
Promethazine (tablets) : prochlorperazine (tablets)	1.0004 : 1	Small	Unknown
Promethazine (tablets) : domperidone (tablets)	1.0005 : 1	Small	Unknown
Promethazine (tablets) : metoclopramide (tablets)	1.0008 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Prochlorperazine (injection) : promethazine (tablets)	1.0007 : 1	Small	Unknown
Prochlorperazine (injection) : cyclizine (tablets)	1.0007 : 1	Small	Unknown
Prochlorperazine (injection) : chlorpromazine (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : prochlorperazine (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : domperidone (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : metoclopramide (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (injection)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : promethazine (tablets)	1.002 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Metoclopramide (injection) : cyclizine (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : chlorpromazine (tablets)	1.003 : 1	Small	Unknown

TABLE 35 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (continued)

Comparison	Implied valuation	Effect size	Evidence on effect
Metoclopramide (injection) : prochlorperazine (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : domperidone (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : metoclopramide (tablets)	1.003 : 1	Small	Unknown
Ondansetron (injection) : metoclopramide (injection)	1.0001 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (injection) : prochlorperazine (injection)	1.002 : 1	Small	Unknown
Ondansetron (injection) : promethazine (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : cyclizine (tablets)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : chlorpromazine (tablets)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : prochlorperazine (tablets)	1.003 : 1	Small	Unknown
Ondansetron (injection) : domperidone (tablets)	1.003 : 1	Small	Unknown
Ondansetron (injection) : metoclopramide (tablets)	1.003 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Promethazine (injection) : ondansetron (injection)	1.0007 : 1	Small	Unknown
Promethazine (injection) : metoclopramide (injection)	1.0008 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Promethazine (injection) : prochlorperazine (injection)	1.002 : 1	Small	Unknown
Promethazine (injection) : promethazine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : cyclizine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : chlorpromazine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : prochlorperazine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : domperidone (tablets)	1.004 : 1	Small	Unknown

continued

TABLE 35 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (continued)

Comparison	Implied valuation	Effect size	Evidence on effect
Promethazine (injection) : metoclopramide (tablets)	1.004 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Cyclizine (injection) : promethazine (injection)	1.003 : 1	Small	Unknown
Cyclizine (injection) : ondansetron (injection)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Cyclizine (injection) : metoclopramide (injection)	1.003 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (injection)	1.005 : 1	Small	Unknown
Cyclizine (injection) : promethazine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : cyclizine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : chlorpromazine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : domperidone (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : metoclopramide (tablets)	1.007 : 1	Small	Unknown
Chlorpromazine (injection) : cyclizine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : promethazine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : ondansetron (injection)	1.01 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Chlorpromazine (injection) : metoclopramide (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : promethazine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : cyclizine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : chlorpromazine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : domperidone (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : metoclopramide (tablets)	1.02 : 1	Small	Unknown
Ondansetron (tablets) : chlorpromazine (injection)	1.01 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects

TABLE 35 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (continued)

Comparison	Implied valuation	Effect size	Evidence on effect
Ondansetron (tablets) : cyclizine (injection)	1.02 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : promethazine (injection)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : ondansetron (injection)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : metoclopramide (injection)	1.03 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (tablets) : prochlorperazine (injection)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : promethazine (tablets)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : cyclizine (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : chlorpromazine (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : prochlorperazine (tablets)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : domperidone (tablets)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : metoclopramide (tablets)	1.03 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Antihistamines : placebo	Not assessable	Not assessable	Antihistamines appear to be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required

TABLE 36 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient

Comparison	Implied valuation	Effect size	Evidence on effect
Prochlorperazine (tablets) : domperidone (tablets)	1.0001 : 1	Small	Unknown
Chlorpromazine (tablets) : prochlorperazine (tablets)	1.00004 : 1	Small	Unknown
Chlorpromazine (tablets) : domperidone (tablets)	1.0001 : 1	Small	Unknown
Prochlorperazine (injection) : chlorpromazine (tablets)	1.0008 : 1	Small	Unknown

continued

TABLE 36 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient (continued)

Comparison	Implied valuation	Effect size	Evidence on effect
Prochlorperazine (injection) : prochlorperazine (tablets)	1.0008 : 1	Small	Unknown
Prochlorperazine (injection) : domperidone (tablets)	1.0009 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (injection)	1.001 : 1	Small	Unknown
Metoclopramide (injection) : chlorpromazine (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : domperidone (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : metoclopramide (injection)	1.0001 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (injection) : prochlorperazine (injection)	1.001 : 1	Small	Unknown
Ondansetron (injection) : chlorpromazine (tablets)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : prochlorperazine (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : domperidone (tablets)	1.002 : 1	Small	Unknown
Cyclizine (injection) : ondansetron (injection)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Cyclizine (injection) : metoclopramide (injection)	1.002 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (injection)	1.004 : 1	Small	Unknown
Cyclizine (injection) : chlorpromazine (tablets)	1.004 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (tablets)	1.004 : 1	Small	Unknown
Cyclizine (injection) : domperidone (tablets)	1.005 : 1	Small	Unknown
Chlorpromazine (injection) : cyclizine (injection)	1.007 : 1	Small	Unknown
Chlorpromazine (injection) : ondansetron (injection)	1.01 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Chlorpromazine (injection) : metoclopramide (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : chlorpromazine (tablets)	1.01 : 1	Small	Unknown

TABLE 36 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient (continued)

Comparison	Implied valuation	Effect size	Evidence on effect
Chlorpromazine (injection) : prochlorperazine (tablets)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : domperidone (tablets)	1.01 : 1	Small	Unknown
Antihistamines : placebo	Not assessable	Not assessable	Antihistamines appear to be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required

TABLE 37 Cost comparison of clinician-prescribed second-line interventions × 2 if admitted as an inpatient

Comparison	Implied valuation	Effect size	Evidence on effect
Most expensive : least expensive	1.06 : 1	Modest	Unknown

TABLE 38 Cost comparison of clinician-prescribed third-line interventions if admitted as an inpatient

Comparison	Implied valuation	Effect size	Evidence on effect
Most expensive : least expensive	1.06 : 1	Modest	Unknown

TABLE 39 Cost comparison of 2-day day case management with 2-day inpatient management

Comparison	Implied valuation	Effect size	Evidence on effect
Inpatient : day case	1.5 : 1	Large	Results indicate that day case management is as effective at improving severity scores as inpatient management for some women. However, more, larger studies are required to provide definitive results

order to be considered a worthwhile use of resources within this package of care. Unfortunately, there was little evidence on effect available and a conclusion could not be drawn. Both ginger and vitamin B6 look promising in reducing symptoms compared with placebo, but the findings from the effectiveness review were not conclusive. The main difference in cost between the use of vitamin B6 and ginger was the difference in medications. The cost of these therapies might be considered modest and were quite similar. From an economics perspective this suggests that we are indifferent about which is used.

Although this may not be considered acceptable to women or clinically, the results from *Tables 32* and *33* allow comparisons to be drawn whether or not women are encouraged to self-medicate before seeking medical treatment from primary care. For example, as an initial response to NVP, seeking advice from a GP who goes on to recommend vitamin B6 as opposed to self-medicating with vitamin B6 implies that the benefits of advice and treatment from the GP are at least 17 times more than the benefits of self-medication. The question for decision-makers is whether or not the benefits from the advice and reassurance gained from contacting a GP are of sufficient value to be worth the additional cost.

Cost comparisons were carried out for all clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit. In the comparison between promethazine and metoclopramide the implied valuation was 1.03 : 1, meaning that in order to be considered efficient, promethazine would need to provide at least 3% more benefits than metoclopramide. No difference in effectiveness was reported from the review; however, it is feasible that such a difference in effect could exist. Ondansetron was compared with both antihistamines (cyclizine and chlorpromazine) within this package of care. Although ondansetron is more expensive (2.2 : 1 and 2.3 : 1, respectively), no difference in effects was shown in the review. In the cost comparison between ondansetron and metoclopramide, the ratio of the cost of ondansetron to the cost of metoclopramide was 2.3 : 1. While the evidence on effect showed that ondansetron was more effective at reducing the symptoms of vomiting than metoclopramide after 4 days, it would need to provide at least 1.3 times more in benefits than metoclopramide in order to be considered a worthwhile use of resources. Finally, antihistamines were compared with placebo and, while an implied valuation was not assessable, the limited information available from the effectiveness review showed that antihistamines appear to be better than placebo in reducing the severity of symptoms.

Within this package of care, it is the patient management costs which are the main cost drivers. Therefore, the difference in cost between all interventions in this package of care was extremely small. The cost comparison ratios ranged from 1 : 1 to 1.03 : 1. From the evidence on effect, ondansetron was more effective at reducing the symptoms of vomiting than metoclopramide after 4 days. Antihistamines also appear to be better than placebo in reducing the severity of symptoms. Other than this, there was no significant difference in effect between any of the comparators included in the effectiveness review within this package of care. Owing to the small difference in cost between all comparators, it is conceivable that any of the more costly options could provide sufficient benefit in order to be worthwhile. However, the limited data available on clinical effectiveness mean that a definitive conclusion cannot be drawn.

As before, it is the patient management costs which are the main cost drivers within this package of care. Cost comparison ratios ranged from 1.00004 : 1 in the smallest instance to 1.01 : 1 in the largest. From the evidence on effect, ondansetron was more effective at reducing the symptoms of vomiting than metoclopramide after 4 days. Antihistamines also appear to be better than placebo in reducing the severity of symptoms. Other than this, there was no significant difference in effect between any of the comparators included in the effectiveness review within this package of care. Again, because of the small difference in cost between all comparators, it is conceivable that any of the more costly options could provide sufficient benefit in order to be considered worthwhile. However, the limited data available on clinical effectiveness mean that a definitive conclusion cannot be drawn.

In order to be considered a worthwhile use of resources, the more expensive option of second-line inpatient care would need to provide at least 6% more in benefits than the less expensive option. Evidence on effect is unknown but it is not implausible that such a difference could exist.

In order to be considered a worthwhile use of resources, the more expensive option would need to provide at least 6% more in benefits than the less expensive option. Evidence on effect is unknown.

The cost of inpatient management compared with day case management in the 2-day scenario presented is almost 50% greater. This means that the benefits of inpatient management would need to be at least 50% greater than those of day case management in order for it to be considered a worthwhile use of resources. Although the results are not definitive due to limited data, the evidence on effectiveness indicates that day case management is as effective at improving severity scores as inpatient management for some women. Therefore, the additional cost of inpatient management in the scenario presented may represent an inefficient use of resources.

For a large number of comparators included in this section, evidence on effect is unknown. All clinically relevant interventions in the UK were included in each package of care; however, clinical evidence is unavailable for many interventions. The ratio of the cost of one intervention to another was calculated

within packages of care to determine the increase in effectiveness that the more costly option would need to provide in order to be considered a worthwhile use of resources. However, as illustrated above, this technique could also be applied to treatments in separate packages of care. This would allow one to, for instance, compare a patient-initiated first-line intervention with a clinician-prescribed second-line intervention to calculate the implied valuation. This information may be of interest to decision-makers.

Summary

- Currently, there are no economic evaluations on the cost-effectiveness of interventions used in the treatment of NVP/HG.
- An economic model has been developed to estimate the relative cost-effectiveness of interventions used in the treatment of NVP/HG and may be used in a model-based economic evaluation once additional information becomes available.
- Pharmacological and non-pharmacological costs of treatment have been combined together to develop informative packages of care. The cost of treatment increases rapidly as patients move from patient-initiated first-line interventions to clinician-prescribed second-line interventions to clinician-prescribed third-line interventions. The cost of medication is small in comparison with the cost of care.
- An implied valuation form of economic evaluation has been used to assess cost-effectiveness. Within each package of care, cost comparisons have been carried out in order to determine the increase in benefits that the more costly option would need to provide in order to be considered efficient.
- For patient-initiated first-line interventions, the cost comparison ratios ranged from 1.01 : 1 in the comparison between vitamin B12 and vitamin B6 to 41 : 1 for the comparison between hypnotherapy and ginger.
- The final cost comparison between inpatient management and day case management indicated that inpatient management would need to provide at least 50% more benefits than day case management to be considered worthwhile. The results on clinical effectiveness indicate that day case management is equally as effective.
- The economic evaluation requires additional information on clinical effectiveness for a large number of interventions in order to make definitive conclusions on cost-effectiveness.

Chapter 19 Issues of importance to patients

Introduction

Nausea and vomiting of pregnancy is a frequently occurring, often debilitating condition which can result in physical, emotional and psychological distress, and a reduced QoL for sufferers. Management of the condition by health-care providers is inconsistent, with many women reporting a poor experience of the health-care system. It is therefore essential to consider the views and opinions of previous sufferers alongside research evidence for clinical effectiveness if the care and the treatment of this condition is to be improved.

Key to carrying out this review was the role played by practitioners and service user members of the research team. Women who had suffered from NVP and/or HG were members of our Project Steering Group along with representatives of key patient advocacy groups in the field. These members represented their own views, not necessarily the views of the advocacy groups. These representatives were consulted at the start of the study, when we informed them about the evidence review and invited their comments on how the project could be improved. Engagement continued throughout the project, so that they were able contribute to meetings in the latter stages of the review when we sought their input to help the research team generate ideas for dissemination.

Background

Previous research and an online survey carried out by the PSS Group (unpublished data, 2014, www.pregnancysicknesssupport.co.uk/), has highlighted the problems women face when seeking advice, support and care from health-care professionals (HCPs). *Tables 40* and *41* represent themes and examples drawn from comments made to women that were reported as part of this survey.

Relating what patients want to review findings

The remit of this systematic review was to evaluate the available research evidence surrounding interventions to treat NVP/HG. It is hoped that results from this review can be used to make suggestions as to how to address some of the problems women face.

Women want HCPs to acknowledge that NVP/HG is a chronic condition and that symptoms can vary dramatically between individuals, from mild to very severe, with no time limit on when symptoms will cease. The review has highlighted inconsistencies with assessing severity of symptoms in many of the included studies. This is likely to be the case within the wider health-care system. HCPs need to acknowledge and be able to assess the wide spectrum of symptoms severity that can be experienced. The introduction of a validated, standard method of assessment may help, so enabling HCPs to discuss the most appropriate treatment options with women. This decision-making process should take into consideration the individual woman's situation, experience, knowledge and past history. Pre-emptive therapy may be appropriate in some cases where women have a history of HG in a previous pregnancy.

The review results will inform clinical guidelines which in turn will support HCPs to advise and prescribe the most effective treatment depending on location, gestation and severity. Different pathways of care make it clear that not all treatments work for all women and that there are different options which should be tried.

TABLE 40 Examples of bad HG/NVP practice by HCPs

Theme	Explanation	Quotes
Knowledge and beliefs	<ul style="list-style-type: none"> • Lack of knowledge regarding NVP and HG • Unaware of appropriate medication or believing medication harmful 	<p><i>Biggest difficulty trying to find GP who understood or would prescribe help</i></p> <p><i>Medical care awful if I'd been diagnosed earlier and given medication earlier I wouldn't have terminated three babies</i></p> <p><i>I felt that they dismissed my symptoms, even tried to convince me I had gastroenteritis</i></p> <p><i>I was responsible for researching all options regarding drugs and asking for things</i></p> <p><i>The total lack of understanding and support from the medical profession both during and after my pregnancy</i></p> <p><i>Despair at lack of understanding or clear direction from GP and hospital staff</i></p> <p><i>Had to see 4 doctors before I was prescribed Buccastem. Previous doctors refused to prescribe me any medication</i></p>
Attitudes	<ul style="list-style-type: none"> • Trivialising the condition • Patronising • Judgemental 	<p><i>Told (by GP) that only 'stupid people' need admitted to hospital</i></p> <p><i>I was very much treated as weak and hysterical by hospital staff and when I asked for injections</i></p> <p><i>I was made to feel like I was wasting their [community midwife and consultant] time and that it was a normal part of pregnancy</i></p> <p><i>Despair at lack of understanding or clear direction from GP and hospital staff</i></p> <p><i>I was treated by medical professionals as though HG was a matter of my mind and not real</i></p> <p><i>His [GP] exact words were 'you can't expect pregnancy to be a walk in the park'</i></p>
Lack of sympathy/ understanding	<ul style="list-style-type: none"> • Disbelief that symptoms could be so bad • Psychosomatic • Comparing women who are suffering from severe symptoms with those suffering mild symptoms 	<p><i>Made to feel as if should just get on with it</i></p> <p><i>I regret the termination and feel if the medical profession had shown any empathy I might have been able to get through the HG that first time</i></p> <p><i>No empathy from GPs. Felt as though I had to beg for medication</i></p> <p><i>I was ignored by my GP, turned away from the hospital due to no ketones in my urine. Little to no support from healthcare professionals until I found a consultant who specialised in HG</i></p>

TABLE 41 Examples of good HG/NVP practice by HCPs

Theme	Explanation	Quotes
Communication and attitude	<ul style="list-style-type: none"> • Being listened too • Being believed • Feeling supported • No fuss made when asking for help 	<p><i>When my GP did see me, she was fantastic but had to rush me into hospital due to dehydration</i></p> <p><i>Midwife was awful and ended up complaining and changing at 30 weeks, wish I'd done it sooner as the next midwife was amazing!</i></p>
Knowledge and understanding	<ul style="list-style-type: none"> • Given appropriate advice • Medication prescribed without hesitation • Early recognition of the condition • Admission to hospital for treatment when needed 	<p><i>I was lucky to have an excellent GP who was quick to diagnose, quick to treat, and got me through</i></p>

The results from this review can also provide women with evidence-based information about different treatment options. Women experiencing NVP for the first time may want to try 'patient initiated' options before seeking medical support. For example, trying preparations of fresh root ginger or acupressure regularly for 3–4 days may help to reduce mild symptoms. Similarly, vitamin B6 taken regularly for at least 3 days may help to reduce mild to moderate symptoms in some women.

If women do seek medical support, prior knowledge relating to treatment effectiveness may then help to empower them, enabling a more satisfactory interaction and outcome.

Summary

Although it is beyond the scope of this review to address all the problems and issues sufferers of NVP/HG have had with their care, it will add to the body of evidence-based knowledge surrounding treatment. This in turn should inform clinical guidelines, so leading to improved care, attitudes and advice for women suffering from NVP/HG. Similarly, if women also have this information they will hopefully be less likely to be treated dismissively and inappropriately.

Dissemination of findings to appropriate audiences will be essential. As well as the published report, the findings will be presented at conferences (medical, midwifery, service users so that women have access to the information) and submitted to peer-reviewed journals.

Chapter 20 Issues of importance to practitioners

Introduction

When providing care and support to women suffering from moderate or severe NVP, HCPs need to consider the evidence on effectiveness and safety as well as the direct and indirect costs to the NHS and to the women. The effectiveness of any intervention is likely to be dependent on the severity of symptoms. Ideally, the severity of NVP should be assessed in a reliable and valid way (e.g. using the PUQE, RINVR or Nausea Questionnaire score). A mild, moderate or severe score, alongside an assessment of how the individual woman is coping, will help HCPs make appropriate recommendations.

The following findings include evidence of effectiveness from the results of the review and associated treatment costs from the economic evaluation presented in *Chapter 18, Economic evaluation*. No reliable data on fetal outcomes [fetal or neonatal death, congenital abnormalities, low birthweight (< 2.5 kg), preterm birth (before 37 weeks' gestation) or small for gestational age (birthweight < 10th centile)] were identified as part of the systematic review. All safety data presented below are derived from large population-based observational studies. This evidence is indirect in that it relates to pregnancy but not specifically to NVP.

The interventions that can be recommended to women fall into three categories:

1. Self-help, 'over-the-counter' interventions which can be recommended by GPs, midwives, practice nurses and pharmacists, in information leaflets and on NHS approved websites.
2. Primary care interventions which can be prescribed by GPs. These may be in addition to or instead of self-help interventions.
3. Secondary care interventions which are delivered in a hospital setting. Some women are referred to hospital by primary care/community HCPs after failure of self-help and/or primary care interventions. However, increasingly, women refer themselves to hospital maternity services (via day/maternity assessment units) often without any prior intervention. Depending on symptom severity and duration, sufferers may need to progress from one category to another and possibly try a number of different, or combinations of interventions.

Findings

First-line 'over-the-counter' interventions and alternative therapies

Ginger

Fresh root ginger may be effective in improving mild symptoms, especially nausea, but findings are not conclusive. When compared with acupuncture, ginger may be more effective, but the quality and quantity of evidence is limited. There are too few data to suggest ginger is any better or worse than vitamin B6 or pharmacological interventions.

Ginger appears to be safe for women during pregnancy. Data from two birth cohorts and several small RCTs suggest there is no evidence of an increased risk of congenital anomalies or adverse perinatal outcomes.^{136,137} In a birth cohort of 68,522 women, 1020 reported using ginger, with no associated increased risks of congenital malformations, stillbirth, perinatal death, preterm birth or low birthweight babies.¹³⁷

The estimated weekly cost of using a ginger preparation was £2.20.

Acupressure at pericardium point P6

Acupressure using correctly placed 'sea-bands' may be effective in improving symptoms of NVP in women with mild to moderate symptoms. Acupressure at KID21 may also be effective but data are limited. The quality of the evidence is low and it is difficult to ascertain whether acupressure is more or less effective than other interventions.

Acupressure is a non-invasive intervention and no safety data are available. The price of sea-bands is approximately £10.

Acupuncture

The evidence of effectiveness of acupuncture is inconclusive. The cost of acupuncture consultations and treatments range between £35 and £70. These data suggest that acupuncture should not be recommended.

Vitamin B6 (pyridoxine)

Vitamin B6 appears to have some effectiveness at improving NVP in women with mild to moderate symptoms, especially at higher doses (≥ 10 mg three times daily for at least 3–4 days).

There is little information regarding the safety of vitamin B6 in isolation. However, the safety of vitamin B6 in combination with antihistamines (doxylamine) has been studied extensively. The results of two meta-analyses, which involved over 200,000 women, found no evidence of increased risk of congenital malformations (relative risk 0.95, 95% CI 0.88 to 1.04¹³⁸ and OR 1.02, 95% CI 0.66 to 1.55).¹³⁹ These findings have since been confirmed by several epidemiological studies.

The weekly cost of vitamin B6 preparations has been estimated at between £0.12 and £2.60.

Second-line interventions prescribed by general practitioners in primary care settings

All quoted medication costs will require the addition of £45 for the GP consultation.

Doxylamine plus pyridoxine

A combination of high-dose pyridoxine and doxylamine may be as effective as metoclopramide at relieving moderate symptoms of NVP. However, a combination of these medications has not been shown to be as effective as ondansetron. Given other data, it can be inferred that this therapy is more effective than no treatment (proxied by placebo in trials).

Evidence suggests that the combined therapy of doxylamine and pyridoxine is safe to use during pregnancy (see *Appendix 7*).

The cost of this combined therapy would be £0.12–2.60 plus cost of doxylamine [price currently not available as not routinely used in the UK NHS, price of alternative antihistamine (e.g. cyclizine) is £0.74–2.22].

Diclectin (delayed release doxylamine 10 mg plus pyridoxine 10 mg) appears to be more effective than placebo when given to treat moderate symptoms and when given pre-emptively to women with a history of moderate NVP in a previous pregnancy. However, Diclectin is currently not available in the UK.

Antihistamines (hydroxyzine, meclizine, cyclizine)

Evidence from the review suggests that these drugs appear to be more effective in treating mild symptoms of NVP compared with no treatment.

A large meta-analysis which involved over 200,000 women who took antihistamines during pregnancy found no evidence of an increased risk of teratogenicity (OR for major malformations was 0.76, 95% CI 0.60 to 0.94), with no other serious maternal or fetal outcomes.¹⁴⁰

Estimated weekly cost of treatment with oral cyclizine was £0.74–2.22.

Dopamine receptor antagonists

There was no evidence of a difference between metoclopramide and promethazine in improving symptoms in moderate cases of NVP. There was also very low-quality evidence from a non-randomised study that droperidol may be effective, especially at higher doses but higher-quality evidence is needed to confirm or refute this.

Dopamine antagonists such as the phenothiazines promethazine and prochlorperazine are regarded as safe: a meta-analysis of eight studies ($n = 2948$) identified no difference in the risk of major malformations (pooled relative risk 1.03, 95% CI 0.89 to 1.22).¹⁴¹ Other drugs in this class used to treat NVP/HG include domperidone, droperidol, trimethobenzamide and metoclopramide. Limited evidence suggests that trimethobenzamide is safe while a recent large study of metoclopramide used during the first trimester of pregnancy ($n = 3458$) found no evidence of an increased risk of major malformations (OR 1.04, 95% CI 0.89 to 1.14) or adverse obstetric outcome.¹⁴² These findings agreed with those of a large, Danish birth cohort (exposed cases $n = 28,486$), which reported no increased risk of congenital malformations (OR 0.95, 95% CI 0.88 to 1.03), miscarriages or stillbirths.¹⁴³ The estimated weekly costs of these drugs are metoclopramide £0.22–0.66, promethazine £0.74–2.22, prochlorperazine £0.47–1.42, domperidone £0.39–1.17.

Serotonin receptor antagonists (ondansetron)

Ondansetron appears to be effective at all levels of symptom severity. However, there is little evidence to say that it is more effective than metoclopramide or antihistamines.

Serotonin antagonists may be safe in pregnancy but experience is limited. One recent case-control study reported an increased risk of cleft palate (adjusted OR 2.37, 95% CI 1.18 to 4.76).¹⁴⁴ However, another Danish study, involving over 600,000 pregnancies, found no association with any adverse fetal outcomes.¹⁴⁵ Concerns have been raised regarding the risk of cardiac arrhythmias [time between start of the Q wave and of the T wave in the heart's electrical cycle (QT) prolongation] with large doses of i.v. ondansetron. A recent systematic review identified that out of all cases of QT prolongation, 67% of patients had a significant medical history or were using a concomitant QT prolonging medication. The study concluded that prior screening of electrocardiogram (ECG) and electrolytes should be limited to high risk patients receiving ondansetron intravenously.¹⁴⁶

The estimated weekly cost of prescribing oral ondansetron was £60.83.

Second- and third-line interventions delivered in secondary care settings

Outpatient/day case management

Limited evidence suggests that this could be an effective alternative to inpatient admission for women with moderate to severe symptoms where i.v. rehydration is required.

There is no population-based safety data regarding pregnancy outcomes following outpatient/day case management.

The estimated cost of an outpatient attendance, which would involve i.v. rehydration, i.v. antiemetics, appropriate blood and urine tests and an ultrasound scan, would be approximately £290.

Inpatient management costs

The estimated cost of a 2-night inpatient admission, which would involve i.v. rehydration, i.v./oral antiemetics, appropriate blood and urine test, an ultrasound scan, thromboprophylaxis and thiamine supplementation, was approximately £800. If a combination of, or alternative antiemetics were required this cost could rise to £850.

Corticosteroids

Evidence is limited but corticosteroids appear to reduce symptom severity, and appear to be more effective at reducing episodes of vomiting than metoclopramide and Phenergan.

Concerns remain about the safety of corticosteroids. In one meta-analysis, the pooled risk ratio for cohort and case-control studies combined revealed no increased risk of major malformations associated with first trimester exposure (cumulative OR 1.45, 95% CI 0.81 to 2.60). However, a subanalysis of case-control studies revealed an increase in the risk of the fetus developing an oral cleft palate (OR 3.35, 95% CI 1.97 to 5.69) and the results were homogeneous between studies.¹⁴⁷ However, other studies do not show this association with cleft palate formation.^{148,149}

Treatment with corticosteroids would generally be initiated following failure of other antiemetic therapies, when the woman is suffering from severe symptoms and during an inpatient admission. The estimated cost of a 5-day inpatient admission in which corticosteroid therapy was given was estimated at £2000–2100.

Enteral nutrition

Enteral feeding appears to be a potentially effective but extreme method of supporting women suffering from very severe symptoms, but no comparative studies were identified.

Costs were not available for enteral feeding but in a recent economic evaluation of total parental feeding the cost per patient of the intervention itself and excluding other hospital based costs was approximately £340 per patient (Kilonzo M, personal communication).

Summary

Overall, there are some data to guide the choice of therapy, but there is little information to guide the choice between therapies. The evidence reviewed suggests that treatments tend to improve symptoms quickly – over a small number of days. Therefore, if a woman gets insufficient relief from a first treatment then this suggests an alternative treatment could be tried. For some treatments there are no data currently available to support their use. Thus, while there may be few safety concerns with some of these treatments, lack of robust effectiveness data can nevertheless guide practitioners in the advice that is given to women.

One obvious concern is around safety. In part this is prompted by the legacy of thalidomide. No reliable evidence directly applicable to the target group of pregnant women was found although evidence from large population sources exists. These data have generally suggested that there is no evidence of any safety concerns with the medications, but this is not the same as ruling out any important differences in safety outcomes. In many cases the CIs are sufficiently wide to include clinically important differences both in favour and against the treatments. Although women need reassurance that treatments are safe, this should not offer false reassurance, therefore advice on safety should be tempered by these findings.

Chapter 21 Discussion

Clinical effectiveness and harms

This study aimed to systematically identify and assess the evidence on the clinical effectiveness and cost-effectiveness of treatments for severe NVP and HG. This review was complicated because there is no agreed point where severe NVP becomes HG, and because studies in this area were both poorly indexed and generally poorly described in terms of severity of symptoms. Scoping searches indicated that the total size of the evidence base in this area was, for a systematic review, relatively small. For this reason the search strategy adopted was unusually comprehensive and included terms related to NVP and HG, but made no restrictions around study design, intervention or language. As such it considers all studies related to NVP and HG.

The study aimed to conduct both a fixed- or random-effect model meta-analysis as well as a Bayesian mixed-treatment comparison. Full details of the proposed methods are reported in the review protocol (PROSPERO CRD42013006642). However, these planned analyses were not performed due to heterogeneity in interventions, trial populations, reporting and definitions of outcome measures and methods. As a consequence the data on effectiveness, fetal outcomes and adverse events were tabulated and reported in narrative fashion. The implication of this is that summary quantitative statistics of the relative performance of different treatments against each other are not available. Rather the focus has been on the consistency of the direction of effect and on the quality of the evidence available. Where possible we have attempted to draw out the implications for both women and practitioners.

In total, 11,830 papers were identified from the combination of standard electronic databases, specialist Chinese databases and various sources of grey literature. From these, 75 papers were identified for data extraction, based on a total of 73 separate studies. The key reasons for exclusion were duplicate papers already included; participant inclusion criteria for the identified study judged not relevant to our review; not including any of the pre-specified outcomes; or as ineligible study design (no comparator group) for effectiveness data or not a population-based case series for rarer maternal and fetal outcomes. The 73 included studies were made up of 64 RCTs and nine case series or non-randomised studies.

With respect to overall quality of the evidence, there was variation both in terms of the quality of the studies and the quality of the reporting. For almost half of all RCTs identified there was insufficient detail provided to permit clear judgement of risk of bias in a range of key areas. Overall, 33 RCTs were classed as having a low within-study risk of bias, 11 RCTs were classed as having a high within-study risk of bias, and the remainder ($n = 20$) were classed as unclear in this respect. The high proportion of studies at unclear risk of bias was due to poor reporting and a lack of detail, particularly in the methods section. All the case series or non-randomised studies were judged as weak methodologically.

The included studies were grouped into the three broad groups of interventions outlined in *Chapter 1*:

1. First-line 'over-the-counter' interventions, which can be recommended by HCPs, in information leaflets and on NHS approved websites, for women to try before seeking medical care. Alternative therapies were included in this group and there were 17 comparisons drawn from 43 studies.
2. Second-line interventions, usually delivered in primary health-care settings and prescribed by a GP, but may also involve referral of women with more severe symptoms for inpatient, outpatient or day case care in hospital. There were 16 comparisons from 20 studies.
3. Third-line interventions reserved for women in hospital with persistent or recurrent symptoms, despite second-line therapies, which are prescribed in hospital settings. There were four comparisons, drawn from 10 studies.

First-line 'over-the-counter' interventions

The most common comparisons drawn were acupressure versus placebo ($n = 8$); steroid versus usual treatment ($n = 7$); ginger versus placebo ($n = 7$); and ginger versus B6 ($n = 6$). Other comparators were considered in only one or two studies. A common finding was that symptoms in all arms (including placebo) improved from baseline.

The evidence on the use of ginger came from 16 RCTs^{13,42,60,63,67,70,74,77,82,89,92,96,102,103,110,113} which were predominantly at low or unclear risk of bias, with four exceptions.^{77,89,92,113} Seven studies compared ginger preparations with placebo^{60,74,82,89,96,110,113} and these generally reported a statistically significant improvement over a range of nausea and vomiting symptoms. However, these data are potentially unreliable because of the number of studies that were judged to have a high or unclear risk of bias. When the comparison was restricted just to those studies at low risk of bias the results were not conclusive. For the comparison of ginger with vitamin B6^{67,70,77,92,102,103} there are some higher-quality studies, but little evidence of a difference in effectiveness. There was a similar finding for the comparison of ginger with the other active treatments (doxylamine–pyridoxine⁶³ or antihistamine⁴² or metoclopramide⁸⁹). For the comparison of ginger versus acupressure,¹³ ginger again looked promising but the evidence was very limited. Overall, ginger might be better than placebo in reducing the severity of symptoms, but these data are limited to less severe symptoms.

A similar picture emerged for vitamin B6. Comparisons of B6 preparations with placebo^{41,100} generally reported evidence of reduced symptoms of nausea, especially for women with more severe symptoms and vomiting. Higher doses of vitamin B6 resulted in a greater improvement in symptoms. However, for other head-to-head comparisons of treatments there were few data identified that suggested a difference in performance.^{59,107,112} Furthermore, for the comparatively well researched acupuncture or acupressure (data were available from 18 trials^{43,61,62,66,73,78–80,83,87,91,94,98,101,104,109,111,115}), there is a suggestion that acupressure may reduce symptoms of nausea and retching in women with mild–moderate symptoms; however, the data are limited and inconclusive. A similar situation was observed for nerve stimulation^{73,98,109} and aromatherapy.^{76,97} Comparisons of traditional Chinese acupuncture and herbal medicine with Western medicine were at high risk of bias and impossible to emulate within the NHS.^{87,115}

No reliable direct evidence was identified in the review on the impact on maternal weight; fetal outcomes [fetal or neonatal death, congenital abnormalities; low birthweight (< 2.5 kg), preterm birth (before 37 weeks' gestation) or small for gestational age (birthweight < 10th centile)]; or adverse events (e.g. pregnancy complications). The identified studies were all too small to provide any reliable data. Indirect evidence came from a variety of large population studies and these data suggested that there was no evidence of any increased risk of adverse fetal or maternal outcomes.

Overall, it is disappointing that the evidence base to guide women in the choice of therapy and practitioners in the advice is so severely lacking. What evidence did exist mainly related to those with milder symptoms and not severe NVP/HG.

Second-line interventions prescribed by general practitioners in primary care settings

For antihistamines, their use resulted in an improvement compared with placebo or no treatment over a range of symptoms.^{68,71,90} Diclectin (a vitamin B6 and antihistamine combination), which is not currently available within the UK, also appears to be more effective than placebo.^{65,84,117} Further, pre-emptive treatment with Diclectin before symptoms of NVP begin in women at high risk of severe NVP recurrence appears to result in a reduced risk of moderate/severe NVP compared with women who take Diclectin once symptoms begin.¹¹⁷

Dopamine antagonists were used in one trial which was judged to be at low risk of bias¹⁰⁶ and one poor-quality non-randomised study.¹²² These studies provided limited evidence suggesting that promethazine is as effective as metoclopramide in reducing the symptoms of NVP. Five trials^{57,72,75,81,105}

and one case series study¹²¹ compared serotonin antagonists (ondansetron) against a range of alternatives. Three trials tested ondansetron against metoclopramide: symptoms were classified as mild to moderate in two trials^{75,81} and severe in one trial.⁵⁷ The identified studies comparing ondansetron with metoclopramide had mixed results, with both drugs improving symptoms. However, the evidence from one study⁸¹ with low risk of bias found ondansetron more effective at reducing vomiting compared with metoclopramide after 4 days. There was some suggestion that ondansetron appears more effective at reducing nausea than vitamin B6 plus doxylamine, but with equivocal evidence for vomiting. However, evidence from two trials^{72,105} comparing ondansetron with antihistamines among women whose symptoms were classified as being moderate to severe found there was no evidence of a significant difference between treatments. The data from one small study⁸⁵ also suggests that the use of transdermal clonidine patches may be effective for the treatment of severe NVP/HG.

With respect to safety, the only data available was indirect sources from large population-based observational studies relating to pregnancy rather than NVP or HG. These sources found no evidence of increased risk of congenital malformations, miscarriages or stillbirths (see *Chapter 20* for details). For ondansetron, concerns have been raised regarding the risk of cardiac arrhythmias (QT prolongation), relating to the administration of large doses of i.v. ondansetron. However, a recent systematic review¹⁴⁶ concluded that prior screening of an ECG and electrolytes should be limited to high risk patients receiving ondansetron intravenously.

Second- and third-line interventions delivered in secondary care settings

For interventions provided in secondary care, one issue is whether they should be provided as inpatient therapy, or can be provided in a day case or outpatient setting. For women with moderate to severe symptoms day case care is feasible and acceptable, and the data suggest that day case management is as effective at improving severity scores as inpatient management for some women.

In terms of specific therapies, there were two studies^{69,108} identified that compared i.v. fluids. One¹⁰⁸ compared different compositions of i.v. solution (dextrose + saline vs. saline only, which was at low risk of bias), and one⁶⁹ compared i.v. fluids containing vitamins with diazepam. The findings suggest that i.v. fluid improves reported symptoms and that dextrose saline may be more effective at improving nausea over time for those with moderate nausea. The lower concentration of sodium in dextrose saline may exacerbate any pre-existing hyponatraemia. High doses/concentrations of dextrose solutions may increase the risk of Wernicke's encephalopathy. However, concentrations in dextrose saline are unlikely to provoke this response. Diazepam appears to be more effective than i.v. fluids at reducing nausea on day 2, but there was no evidence post treatment for those with moderate/severe nausea or HG.

The use of corticosteroids was better researched with seven studies^{58,64,93,99,114,116,125} identified (three at low risk of bias,^{64,93,99} three where the risk of bias was unclear/high^{58,114,116} and one weak case series study¹²⁵). The evidence suggested a trend towards improved symptoms with steroids compared with placebo, but the results were not statistically significant. There was a small amount of evidence suggesting steroids were more effective at reducing vomiting episodes than Phenergan suppositories or metoclopramide. Nevertheless, the overall evidence base on effectiveness is limited and there are concerns over safety. As reported in *Chapter 20*, a subanalysis of case-control studies identified an increase in the risk of the fetus developing an oral cleft palate (OR 3.35, 95% CI 1.97 to 5.69) and the results were homogeneous between studies,¹⁴⁷ although other studies did not show this association.^{148,149} Few data were found on the use of assisted feeding in this patient group, but the limited data showed that enteral feeding is an effective, but extreme, method of supporting women suffering from very severe symptoms as a last resort.

In summary, the evidence on effectiveness of all treatments was severely limited by the poor quality of the evidence available. Although there appears to be some evidence that some treatments (ginger preparations, vitamin antihistamines, metoclopramide, B6) were better than placebo for mild disease, there is little on the effectiveness of treatments for more severe disease. Evidence on differences in effectiveness was available for few other comparisons.

Cost-effectiveness

As part of the systematic review, a review of the economic evaluations was planned but no studies that met the inclusion criteria were identified. Likewise, an economic evaluation based on a modelling exercise was planned. Given the paucity of the effectiveness data this proved not to be possible. As a consequence, a simpler analysis was conducted that estimated the cost of the intervention and then used these data to consider the difference in effectiveness that would be implied if a more costly intervention was used instead of a less costly intervention. This form of analysis is based on the theoretical conditions required for an efficient allocation of resources. The implied relative effectiveness estimated along with information on cost were then set alongside the limited evidence on effectiveness.

The results of the economic analysis indicate that the cost of treating patients with NVP/HG increases rapidly with severity. The ranges of costs are as follows: patient-initiated first-line interventions, cost between £0.12 and £90 per week; patient-initiated first-line interventions following a GP visit, cost between £45 and £47 per week; and clinician-prescribed second-line interventions (e.g. antihistamines), cost between £45 and £106 per week, including the cost of the GP visit itself.

Some women are referred to hospital by primary care/community HCPs after failure of self-help and/or primary care interventions. Women may also self-refer themselves to hospital maternity services (via day/maternity assessment units), often without any prior intervention. This opens up a range of other potential interventions and potentially greatly increases costs. For example, for a clinician-prescribed second-line intervention when a woman attends hospital as a day case the cost would be between almost £300 per day. Whereas if the woman were admitted to hospital as an inpatient, the cost for a 2-day admission would be approximately £800. For those women with the most severe or intractable symptoms, combinations of medications may be administered and should the woman be admitted for 5 days, the cost was estimated to be between £2000 and £2100. However, the actual medication costs as a component of total cost is relatively small, with the majority of cost incurred for the inpatient stay itself.

These data on costs are not of much practical value without information on effectiveness (although they may help in assessing budget impacts). As noted, data on effectiveness are very limited. Nevertheless, what data there is can be interpreted, along with the cost data. For patient-initiated first-line interventions, the cost comparison ratios ranged from 1.01 : 1 for the comparison between vitamin B12 and vitamin B6 (i.e. vitamin B12 would only need to be 1% more effective than vitamin B6 to be considered cost-effective) to 41 : 1 for the comparison between hypnotherapy and ginger. In cost terms there is little difference between possible medications and some evidence that ginger and vitamin B6 may provide some relief. Therefore, in economic terms the results suggest that we are indifferent between these treatments. There is also evidence that acupuncture may be effective, but the cost of acupuncture consultations and treatments range from £35 to £70. Comparing this with the cost of a medication like ginger or vitamin B6, therefore, choosing acupuncture over either ginger or vitamin B6, would imply that acupuncture is at least 13 times better (taking the costs most in favour of acupuncture, £35 for acupuncture and £2.60 for the weekly cost of ginger or vitamin B6). A judgement is required whether or not this is plausible. If it is not judged plausible then this suggests that acupuncture should not be adopted as an initial treatment, but it does not rule it out as a second treatment if the first does not provide sufficient symptom relief.

The relatively small medication costs in comparison to the cost of accessing primary or secondary care had a direct impact on the results of the implied value method of economic evaluation. For the majority of cost comparisons in the remaining packages of care, the difference in cost was so small that only a marginal increase in clinical effectiveness would be required in order for the more costly interventions to be considered a worthwhile use of resources. Nevertheless, the same sort of judgement described above still applies.

A final cost comparison considered was between inpatient management and day case management. The results of this analysis indicated that inpatient management would need to be 1.5 times more effective than day case management. The question for decision-makers is whether or not this difference is plausible

given that the evidence from the systematic review provided no evidence of a difference in effectiveness between inpatient and day case care.

Overall, however, even though costs for the different treatments can be estimated, it must be emphasised that the evidence on cost-effectiveness is constrained by the very limited data available on effectiveness.

Strengths and limitations

The main strength of this review was the comprehensive and systematic approach taken to review the literature. The searches sought to be exhaustive, and have included the major electronic databases, grey literature sources and non-English-language databases. The search strategy itself was broadly defined and included terms related to NVP and HG, but made no restrictions on language, study design or type of intervention. Taken together, the searches should have identified all relevant studies. It is possible, however, that there exist some relevant data that remains hidden because of non-publication, but it is questionable about what this might add given the generally very small size of identified studies and their often very limited methodology. Thus, it seems implausible that a large high-quality study was missed.

The process of assembling data on interventions was also rigorous. As noted above, no studies were excluded on the basis of language, and where necessary translations were sought. Risk of bias assessments were conducted for included RCTs and for non-randomised studies using high-quality tools. Data were extracted for pre-specified outcomes onto a standard form to prevent biases caused by selective data extraction. The choice of outcomes for the systematic review was itself based on the advice of our expert clinical panel and from advice from women themselves and patient representative groups.

The study included a pre-planned economic component and although the proposed economic modelling was not possible, information were rigorously assembled and presented in such a way as to help facilitate judgements about alternative therapies. As such, although less sophisticated than the proposed model, it is entirely consistent with the role of economic evaluation as an aid to decision-making.

The review itself identified 73 studies but very few were available for most comparisons and not every study contributed data to each outcome, including our pre-specified primary outcome of severity of symptoms (such as PUQE, RINVR, McGill Nausea Questionnaire, NVPI, VASs). This was compounded by the differences in reporting of studies that precluded the opportunity for meta-analysis. As a consequence, we were restricted to a narrative review that was not able to produce summary measures that quantified the effect on outcomes. Rather we were restricted to reporting on the likely direction of effect. This problem will remain and there is a need for future studies to use standardised measures of severity and include patient-centric outcomes such as QoL. The development of a standardised set of outcomes that would be measured and reported for all studies is required.

One set of outcome measures that are likely to be very important to women and practitioners are around safety. In this study we sought to assemble data on fetal outcomes and adverse events to both the mother and child. No reliable data meeting our inclusion criteria were identified. To partially overcome this limitation we looked at large population-based observational studies. This evidence is indirect in that it relates to pregnancy but not specifically to NVP. Nevertheless, it does provide some reassurance that many of the treatments are not clearly associated with an increase in adverse events. However, by the same token, these data do not rule out differences that would be considered important by the women themselves or by practitioners.

With respect to the economic element of the research, this element shares many of the strengths and limitations already described. The main manifestation of this limitation is that the planned economic modelling was not possible. An alternative approach was used and although this may provide some useful data for decision-making its value is limited.

Chapter 22 Conclusions

Introduction

In developing these conclusions, we worked with a range of individuals and organisations to ensure their relevancy to patients and practitioners. Women who had suffered from NVP and/or HG and representatives of key patient advocacy groups in the field were members of our Project Steering Group, and were consulted in shaping both the review implications and our recommendations for future research. Our Steering Group and Project Management Group also included a number of clinicians working with women sufferers of NVP and/or HG in a range of health-care settings, representing midwifery, general practice, obstetrics, teratology, paediatrics and clinical pharmacology.

Implications for women and for practitioners

- Nausea and vomiting in pregnancy occurs frequently. Even without treatment, the symptoms for some women will resolve. For women who do experience more persistent problems there are some simple approaches that they could adopt themselves that might improve symptoms and/or QoL. These could include increasing oral fluid intake, eating small frequent meals, eating bland foods/protein-predominant meals, avoiding spicy, odorous and fatty foods and stopping iron-containing multivitamins. However, for others there are a range of treatments available to women where there is some evidence that they might help.
- Where symptoms are mild, there are a number of first-line over-the-counter or self-purchased therapies where there is some evidence that they work. These therapies may also be considered as initial treatments if a woman sees a doctor, nurse, midwife or other HCP. These treatments are ginger supplements and vitamin B6. These are generally low cost and there is no evidence that they are a risk to the health of the mother or baby. There is no proof that a more expensive version of either ginger or vitamin B6 is any better than a lower cost option. Higher doses of vitamin B6 were found to be more effective than lower doses in reducing symptoms (0.64 mg twice daily vs. 1.28 mg per day). There is also some evidence that acupressure may be effective, but the evidence is very limited. The available evidence suggests that any benefit derived from these over-the-counter therapies will be evident within 3–4 days. Thus, women need to know that, in such cases, it is worth trying something else or consulting with a GP as there are other treatments available via prescription. There is no evidence that these treatments are unsafe to use. Further details on dosage and effects are in *Chapters 4–7* and *Tables 8–11*.
- Where symptoms are mild to moderate and/or if the above over-the-counter therapies have not proved helpful to women, a number of second-line interventions are available via prescription. Of the drugs that a GP might prescribe there is evidence that an antihistamine can help reduce symptoms (either alone or combined with vitamin B6) compared with no treatment. Limited data suggests that metoclopramide and promethazine may also help when symptoms are moderate. Droperidol may also be effective, especially at higher doses. Available evidence indicates these treatments are likely to be safe, but more research is needed to clarify this (further details on dosage and effects are in *Chapter 9* and *Table 13*).
- A GP may also prescribe ondansetron when other treatments have failed. This treatment may be effective for some women in reducing symptoms. The drug appears to be safe in pregnancy but experience is limited and more research is needed. Where women are given large doses of ondansetron and have a risk of some cardiac conditions, they may need an ECG and to have their blood chemistry checked (further details on dosage and effects are in *Chapter 8* and *Table 12*).

- Where symptoms are more severe or persistent, based on assessment using a validated measurement scale, care may be provided in hospital, and for these women a further range of treatments are available. Evidence suggests that where available these treatments can be provided as an outpatient or day case patient rather than requiring an admission, and usually involve rehydration with i.v. fluids. Dextrose saline appears to be more effective at reducing nausea when compared with normal saline, but must be administered with care as excessive dextrose can exacerbate the potential problem of Wernicke's encephalopathy (see *Chapter 12* and *Table 16* for more details).
- Treatment with corticosteroids does work but would generally only be used following failure of other treatments, when the woman is suffering from severe symptoms and during an inpatient admission. This is because doctors and other HCPs are cautious about the safety of these treatments and the side effect they may have for both the mother and the fetus. Therefore, doctors would like to use other options first if possible. Ideally, more evidence is needed comparing corticosteroids to other antisickness medications (further details on dosage and effects are in *Chapter 15* and *Table 19*).
- Where symptoms are extremely severe, and when the prolonged effects of severe nausea and vomiting have made women extremely ill, assisted feeding either by tube directly into the stomach or, in very rare cases, by i.v. catheter may be effective. As there are risks associated with this type of treatment, it is restricted to women suffering from very severe symptoms (see *Chapter 16* and *Table 20* for more details).

Recommendations for research

Trajectory of research

Any recommendations for further research need to take into account the trajectory of *new* research. Research on treatments and management of NVP/HG is ongoing: since the date of the last systematic update of published research for this study (September 2014) new evidence has been published. Although not assembled systematically, scoping searches conducted in early December 2014 identified two potentially relevant studies. The first was a review of the use of gabapentin in pregnancy¹⁵⁰ and the second was a RCT comparing day care with inpatient management.¹⁵¹ Neither study was formally assessed, but the authors' conclusions were that further clinical trials were needed on the use of gabapentin for HG, and that day case care was acceptable and reduced length of stay. These conclusions are consistent with the findings reported in this review.

There are also a number of studies recorded as ongoing on searches of ClinicalTrials.gov (www.clinicaltrials.gov) and the International Clinical Trials Registry Platform (<http://apps.who.int/trialsearch/default.aspx>) (searches of both were conducted in December 2014). Ongoing studies are investigating:

- (a) gabapentin versus ondansetron for HG in a double-blind RCT of 80 women with HG (NCT02163434, due to finish in May 2018)
- (b) Diclectin as a pre-emptive treatment for NVP versus treatment with Diclectin when symptoms occur in a RCT (NCT00293644, due to end in April 2015)
- (c) rapid hydration with hospital admission for HG (ISRCTN24659467)
- (d) enteral feeding with oral rehydration for the treatment of HG.

Research recommendations

Given the trajectory of research and the evidence gaps identified by this study, in order of priority, the following recommendations for research are made:

- A well-designed multicentre RCT to determine which second-line, hospital-prescribed therapy should be adopted as mainstream provision for the treatment of NVP in the UK NHS. This could compare day case or inpatient delivery of (a) a combination of antiemetic therapy (antihistamines or dopamine receptor antagonists or serotonin receptor antagonists) with i.v. rehydration (rapid i.v. hydration or standard inpatient care) as required; against (b) alternative antiemetic therapy (antihistamines or dopamine receptor antagonists or serotonin receptor antagonists) with i.v. rehydration (rapid i.v. hydration or standard inpatient care).

- A well-designed multicentre RCT to test the use of subsequent treatments, such as steroids, as a third-line therapy. This could examine, for example, the effectiveness of corticosteroids versus serotonin receptor antagonists (ondansetron). In this trial, the indication for steroid use could also be investigated. For example, failure of successful treatment following three admissions for i.v. rehydration and medication. The dose, duration of treatment and effectiveness could also be examined.
- A well-designed multicentre RCT to determine which second-line, GP-prescribed therapy should be adopted as mainstream provision for the treatment of NVP in UK primary care. This could compare the effectiveness and cost-effectiveness of vitamin B6–antihistamine combination against, for example, a dopamine receptor antagonist.
- Six key factors should be incorporated within any future research trial of second-line, third-line or GP-initiated NVP/HG therapies:
 - i. Stakeholder co-design: all stakeholders and, in particular, the women themselves, should be involved in the design of future studies to ensure that future research produces information of relevance to them as well as to health services.
 - ii. Embedded process evaluation in order to examine the views of women participants on the intervention/therapy being trialled, study how the intervention is implemented in practice, investigate contextual factors that affect the delivery of an intervention, and study the way effects vary in subgroups.
 - iii. Mental health and well-being outcomes: we know from other sources that severe NVP has an impact on mental health both ante- and post-natally. Therefore consideration needs to be given as to how to capture the impact on women regarding these factors. A variety of approaches are possible, but it may for example include qualitative work and/or the use of existing validated tools such as (a) QoL indices such as Short Form questionnaire-36 items (total and subscales); (b) measures of mental health and well-being such as Edinburgh Postnatal Depression Scale; and (c) satisfaction with care as measured by Client Satisfaction Questionnaire-8.
 - iv. Validated symptom severity scale: given the problems identified in this review with the lack of consistent outcome measures in the existing evidence base, entry to future trials should be based on PUQE criteria as an objective means of establishing severity, with subsequent outcome measurements also based on the same score.
 - v. Standard reporting criteria for adverse events, maternal and fetal outcomes: all studies of drugs administered during the first trimester of pregnancy should have obligatory reporting of rates of adverse events and maternal/fetal outcomes to the UKTIS central database to support the availability of reliable prospective controlled data. In addition, all trials of NVP/HG interventions in the UK NHS should include sufficiently long-term participant follow-up periods to enable the capture of relevant maternal and fetal outcomes given the safety profile of the therapies compared. These outcome data should be analysed to determine whether or not there are any associations between particular therapies and/or characteristics of women, in order to guide further research into stratified care.
 - vi. Economic evaluation of NVP/HG therapies: as there may be trade-offs between cost and effects, future studies should also include an economic evaluation.
- A large simple RCT of self-medication. The choice of interventions should be informed by the preferences of women about which treatments are of most relevance to them. Points 1–5 listed above also apply, but the inclusion of an economic component would also need to be justified given the anticipated low cost of the intervention and that a more effective treatment would save subsequent management costs.

Acknowledgements

We thank Professor Gideon Koren (Director, the Motherisk Program; Professor of Pediatrics, Pharmacology, Pharmacy, Medicine and Medical Genetics at The University of Toronto; Senior Scientist at The Research Institute Hospital for Sick Children; and The Ivey Chair in Molecular Toxicology, at the University of Western Ontario Canada) for advice on clinical aspects of the research. We thank our patient and public representatives, Dr Nicolette Rousseau and Juliet Hall, for providing ongoing advice and support as part of the Review Steering Group. We thank Dr Margaret O'Hara and the other women volunteers from PSS for sharing their views and experiences on the review topic, and for the contribution of survey data to the chapter on patient issues. We thank the UKTIS for the provision of enquiry data relating to medications for HG/NVP. We thank Astrid McIntyre for secretarial support and assistance with obtaining full-text papers. The views expressed are those of the authors and not necessarily those of the funding bodies. Any errors are the responsibility of the authors.

Contributions of authors

Amy O'Donnell (Research Associate) and **Catherine McParlin** (Senior Research Midwife) took the role as co-first authors. They screened the search results, assessed full-text studies for inclusion, undertook data extraction and quality assessment, drafted the introduction and background chapter, methods, overview of included studies, individual intervention results chapters and co-ordinated the review.

Stephen C Robson (Professor of Fetal Medicine), **Catherine Nelson-Piercy** (Professor of Obstetric Medicine), **Justine Norman** (GP) and **Laura Yates** (Consultant in Clinical Genetics and Head of Teratology) provided expert advice on clinical aspects of the review.

Fiona Beyer (Information Scientist) developed and ran the search strategies, reconciled search results, obtained papers and managed the reference database.

Eoin Moloney drafted the chapter on the economic evaluation, supervised by **Luke Vale** (Professor of Health Economics).

Andrew Bryant undertook quality assessment and data extraction, contributed to the drafting of the overview chapter, individual results chapters, with particular responsibility for the sections on risk of bias including the production of the SoF tables, supervised by **Colin Muirhead** (Lecturer in Medical Statistics).

Jennifer Bradley (Research Assistant) and **Emma Simpson** (Research Assistant) undertook data extraction and quality assessment of included papers, supervised by **Amy O'Donnell**.

Dorothy Newbury-Birch (Lecturer in Public Health) provided expert advice on the systematic review process.

Brian Swallow (Expert Advisor) provided expert advice in relation to the patient perspective.

Stephen C Robson and **Luke Vale** were coprinciple investigators and take overall responsibility for the study.

All authors assisted in preparing the manuscript, reading and commenting on drafts, and reading the final draft.

Publication

McParlin C, O'Donnell A, Robson SC, Beyer F, Moloney E, Bryant A, *et al.* Treatments for hyperemesis gravidarum and nausea and vomiting in pregnancy: a systematic review. *JAMA* 2016;**316**:1392–401.

Data sharing statement

All available data can be obtained from the corresponding author.

References

1. Niebyl JR. Clinical practice. Nausea and vomiting in pregnancy. *N Engl J Med* 2010;**363**:1544–50. <http://dx.doi.org/10.1056/NEJMcp1003896>
2. Jarvis S, Nelson-Piercy C. Management of nausea and vomiting in pregnancy. *BMJ* 2011;**342**:d3606. <http://dx.doi.org/10.1136/bmj.d3606>
3. Attard CL, Kohli MA, Coleman S, Bradley C, Hux M, Atanackovic G, et al. The burden of illness of severe nausea and vomiting of pregnancy in the United States. *Am J Obstet Gynecol* 2002;**186**:S220–7. <http://dx.doi.org/10.1067/mob.2002.122605>
4. Jueckstock JK, Kaestner R, Mylonas I. Managing hyperemesis gravidarum: a multimodal challenge. *BMC Med* 2010;**8**:46. <http://dx.doi.org/10.1186/1741-7015-8-46>
5. Trogstad LIS, Stoltenberg C, Magnus P, Skjaerven R, Irgens LM. Recurrence risk in hyperemesis gravidarum. *Br J Obstet Gynaecol* 2005;**112**:1641–5. <http://dx.doi.org/10.1111/j.1471-0528.2005.00765.x>
6. Vikanes AV, Skjaerven R, Grijibovski AM, Gunnes N, Vangen S, Magnus P. Recurrence of hyperemesis gravidarum across generations: population based cohort study. *BMJ* 2010;**340**:c2050. <http://dx.doi.org/10.1097/ogx.0b013e3182021d24>
7. Derbent AU, Yanik FF, Simavli S, Atasoy L, Urun E, Kuscu UE, et al. First trimester maternal serum PAPP-A and free beta-HCG levels in hyperemesis gravidarum. *Prenat Diagn* 2011;**31**:450–3. <http://dx.doi.org/10.1002/pd.2715>
8. Goodwin TM, Poursharif B, Korst LM, MacGibbon KW, Romero R, Fejzo MS. Secular trends in the treatment of hyperemesis gravidarum. *Am J Perinatol* 2008;**25**:141–7. <http://dx.doi.org/10.1055/s-2008-1040344>
9. Ismail SK, Kenny L. Review on hyperemesis gravidarum. *Baillieres Best Pract Res Clin Gastroenterol* 2007;**21**:755–69. <http://dx.doi.org/10.1016/j.bpg.2007.05.008>
10. Yamazaki K, Sato K, Shizume K, Kanaji Y, Ito Y, Obara T, et al. Potent thyrotropic activity of human chorionic gonadotrophin variants in terms of 125I incorporation and de novo synthesized thyroid hormone release in human thyroid follicles. *J Clin Endocrinol Metab* 1995;**80**:473–9.
11. Verberg MF, Gillott DJ, Al-Fardan N, Grudzinskas JG. Hyperemesis gravidarum, a literature review. *Hum Reprod Update* 2005;**11**:527–39. <http://dx.doi.org/10.1093/humupd/dmi021>
12. Sandven I, Abdelnoor M, Nesheim B-I, Melby KK. Helicobacter pylori infection and hyperemesis gravidarum: a systematic review and meta-analysis of case-control studies. *Acta Obstet Gynecol Scand* 2009;**88**:1190–200. <http://dx.doi.org/10.3109/00016340903284927>
13. Saberi F, Sadat Z, Abedzadeh-Kalahroudi M, Taebi M. Acupressure and ginger to relieve nausea and vomiting in pregnancy: a randomized study. *Iran Red Crescent Med J* 2013;**15**:854–61. <http://dx.doi.org/10.5812/ircmj.12984>
14. Mazzotta P, Maltepe C, Navioz Y, Magee LA, Koren G. Attitudes, management and consequences of nausea and vomiting of pregnancy in the United States and Canada. *Int J Gynaecol Obstet* 2000;**70**:359–65. [http://dx.doi.org/10.1016/S0020-7292\(00\)00255-1](http://dx.doi.org/10.1016/S0020-7292(00)00255-1)
15. Smith C, Crowther C, Beilby J, Dandeaux J. The impact of nausea and vomiting on women: a burden of early pregnancy. *Aust N Z J Obstet Gynaecol* 2000;**40**:397–401. <http://dx.doi.org/10.1111/j.1479-828X.2000.tb01167.x>

16. Poursharif B, Korst LM, MacGibbon KW, Fejzo MS, Romero R, Goodwin TM. Elective pregnancy termination in a large cohort of women with hyperemesis gravidarum. *Contraception* 2007;**76**:451–6. <http://dx.doi.org/10.1016/j.contraception.2007.08.009>
17. Tan PC, Zaidi SN, Azmi N, Omar SZ, Khong SY. Depression, anxiety, stress and hyperemesis gravidarum: temporal and case controlled correlates. *PLOS ONE* 2014;**9**:e92036. <http://dx.doi.org/10.1371/journal.pone.0092036>
18. McCarthy FP, Khashan AS, North RA, Moss-Morris R, Baker PN, Dekker G, *et al.* A prospective cohort study investigating associations between hyperemesis gravidarum and cognitive behavioural and emotional well-being in pregnancy. *PLOS ONE* 2011;**6**:e27678. <http://dx.doi.org/10.1371/journal.pone.0027678>
19. Ayyavoo A, Derraik JGB, Hofman PL, Cutfield WS. Hyperemesis gravidarum and long-term health of the offspring. *Am J Obstet Gynecol* 2014;**210**:521–5. <http://dx.doi.org/10.1016/j.ajog.2013.11.035>
20. NHS Information Centre. *Hospital Episode Statistics for England. In-Patient Statistics – Diagnosis*. Leeds: NHS Information Centre; 2012.
21. Locock L, Alexander J, Rozmovitz L. Women’s responses to nausea and vomiting in pregnancy. *Midwifery* 2008;**24**:143–52. <http://dx.doi.org/10.1016/j.midw.2006.12.001>
22. Veenendaal MV, van Abeelen AF, Painter RC, van der Post JA, Roseboom TJ. Consequences of hyperemesis gravidarum for offspring: a systematic review and meta-analysis. *BJOG* 2011;**118**:1302–13. <http://dx.doi.org/10.1111/j.1471-0528.2011.03023.x>
23. Bolin M, Akerud H, Cnattingius S, Stephansson O, Wikstrom AK. Hyperemesis gravidarum and risks of placental dysfunction disorders: a population-based cohort study. *Br J Obstet Gynaecol* 2013;**120**:541–7. <http://dx.doi.org/10.1111/1471-0528.12132>
24. Koren G, Boskovic R, Hard M, Maltepe C, Navioz Y, Einarson A. Motherisk-PUQE (pregnancy-unique quantification of emesis and nausea) scoring system for nausea and vomiting of pregnancy. *Am J Obstet Gynecol* 2002;**186**:S228–31. <http://dx.doi.org/10.1067/mob.2002.123054>
25. Koren G, Piwko C, Ahn E, Boskovic R, Maltepe C, Einarson A. Validation studies of the Pregnancy Unique-Quantification of Emesis (PUQE) scores. *J Obstet Gynaecol* 2005;**25**:241–4. <http://dx.doi.org/10.1080/01443610500060651>
26. Lacasse A, Rey E, Ferreira E, Morin C, Berard A. Validity of a modified Pregnancy-Unique Quantification of Emesis and Nausea (PUQE) scoring index to assess severity of nausea and vomiting of pregnancy. *Am J Obstet Gynecol* 2008;**198**:71.e1–71.e7. <http://dx.doi.org/10.1016/j.ajog.2007.05.051>
27. Rhodes V, Watson P, Johnson M. Development of reliable and valid measures of nausea and vomiting. *Cancer Nurs* 1984;**7**:33–41. <http://dx.doi.org/10.1097/00002820-198402000-00003>
28. Zhou Q, O’Brien B, Soeken K. Rhodes Index of nausea and vomiting-Form 2 in pregnant women. A confirmatory factor analysis. *Nurs Res* 2001;**50**:251–7. <http://dx.doi.org/10.1097/00006199-200107000-00009>
29. O’Brien B, Relyea MJ, Taerum T. Efficacy of P6 acupressure in the treatment of nausea and vomiting during pregnancy. *Am J Obstet Gynecol* 1996;**174**:708–15. [http://dx.doi.org/10.1016/S0002-9378\(96\)70454-4](http://dx.doi.org/10.1016/S0002-9378(96)70454-4)
30. Melzack R, Rosberger Z, Hollingsworth M. New approaches to measuring nausea. *CMAJ* 1985;**133**:755–8.

31. Lacroix R, Eason E, Melzack R. Nausea and vomiting during pregnancy: a prospective study of its frequency, intensity, and patterns of change. *Am J Obstet Gynecol* 2000;**182**:931–7. [http://dx.doi.org/10.1016/S0002-9378\(00\)70349-8](http://dx.doi.org/10.1016/S0002-9378(00)70349-8)
32. Swallow BL, Lindow SW, Masson EA, Hay DM. Women with nausea and vomiting in pregnancy demonstrate worse health and are adversely affected by odours. *J Obstet Gynaecol* 2005;**25**:544–9. <http://dx.doi.org/10.1080/01443610500230783>
33. Swallow BL, Lindow SW, Masson EA, Hay DM. Development of an instrument to measure nausea and vomiting in pregnancy. *J Obstet Gynaecol* 2002;**22**:481–5. <http://dx.doi.org/10.1080/0144361021000003582>
34. Chandra K, Magee L, Einarson A, Koren G. Nausea and vomiting in pregnancy: results of a survey that identified interventions used by women to alleviate their symptoms. *J Psychosom Obstet Gynaecol* 2003;**24**:71–5. <http://dx.doi.org/10.3109/01674820309042804>
35. Thomson M, Corbin R, Leung L. Effects of ginger for nausea and vomiting in early pregnancy: a meta-analysis. *J Am Board Fam Med* 2014;**27**:115–22. <http://dx.doi.org/10.3122/jabfm.2014.01.130167>
36. Ashkenazi-Hoffnung L, Merlob P, Stahl B, Klinger G. Evaluation of the efficacy and safety of bi-daily therapy with pyridoxine and doxylamine for nausea and vomiting of pregnancy. *MAJ* 2013;**15**:23–6.
37. American Congress of Obstetricians and Gynecologists. American Congress of Obstetricians and Gynecologists (ACOG) practice bulletin #52: nausea and vomiting of pregnancy. *Obstet Gynecol* 2004;**103**:803–16.
38. Maina A, Todros T. A novel approach to hyperemesis gravidarum: evaluation by a visual analogue scale score and treatment with transdermal clonidine. *Obstetric Medicine* 2011;**4**:156–9. <http://dx.doi.org/10.1258/om.2011.110050>
39. Guttuso TJr, Robinson LK, Amankwah KS. Gabapentin use in hyperemesis gravidarum: a pilot study. *Early Hum Dev* 2010;**86**:65–6. <http://dx.doi.org/10.1016/j.earlhumdev.2009.11.003>
40. O'Donnell A, Vale L, Moloney E, Muirhead C, Bryant A, Beyer F, et al. *A Systematic Review and Economic Modelling of the Relative Clinical- and Cost-Effectiveness of Interventions for Hyperemesis gravidarum*. PROSPERO 2013:CRD42013006642 URL: www.crd.york.ac.uk/PROSPERO/display_record.asp?ID=CRD42013006642 (accessed on 4 September 2014).
41. Vutyavanich T, Wongtra-ngan S, Ruangsri R. Pyridoxine for nausea and vomiting of pregnancy: a randomized, double-blind, placebo-controlled trial. *Am J Obstet Gynecol* 1995;**173**:881–4. [http://dx.doi.org/10.1016/0002-9378\(95\)90359-3](http://dx.doi.org/10.1016/0002-9378(95)90359-3)
42. Pongrojpraw D, Somprasit C, Chanthasenanont A. A randomized comparison of ginger and dimenhydrinate in the treatment of nausea and vomiting in pregnancy. *J Med Assoc Thai* 2007;**90**:1703–9.
43. Can Gurkan O, Arslan H. Effect of acupuncture on nausea and vomiting during pregnancy. *Complement Ther Clin Pract* 2008;**14**:46–52. <http://dx.doi.org/10.1016/j.ctcp.2007.07.002>
44. Pham B, Klassen TP, Lawson ML, Moher D. Language of publication restrictions in systematic reviews gave different results depending on whether the intervention was conventional or complementary. *J Clin Epidemiol* 2005;**58**:769–76. <http://dx.doi.org/10.1016/j.jclinepi.2004.08.021>
45. GRADE Working Group. Grading quality of evidence and strength of recommendations. *BMJ* 2004;**328**:1490–4. <http://dx.doi.org/10.1136/bmj.328.7454.1490>

46. Higgins J, Green S, editors. *Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0*. The Cochrane Collaboration; 2011. URL: www.cochrane-handbook.org (accessed 16 March 2015).
47. National Collaborating Centre for Methods and Tools. *Quality Assessment Tool for Quantitative Studies Method*. Hamilton, ON: McMaster University; 2008. URL: www.nccmt.ca/registry/view/eng/15.html (accessed April 2011).
48. Armijo-Olivo S, Stiles CR, Hagen NA, Biondo PD, Cummings GG. Assessment of study quality for systematic reviews: a comparison of the Cochrane Collaboration Risk of Bias Tool and the Effective Public Health Practice Project Quality Assessment Tool: methodological research. *J Eval Clin Pract* 2010;**18**:12–18. <http://dx.doi.org/10.1111/j.1365-2753.2010.01516.x>
49. Thomas BH, Ciliska D, Dobbins M, Micucci S. A process for systematically reviewing the literature: providing the research evidence for public health nursing interventions. *Worldviews Evid Based Nurs* 2004;**1**:176–84. <http://dx.doi.org/10.1111/j.1524-475X.2004.04006.x>
50. Guyatt GH, Oxman AD, Kunz R, Brozek J, Alonso-Coello P, Rind D, et al. GRADE guidelines 6. Rating the quality of evidence – imprecision. *J Clin Epidemiol* 2011;**64**:1283–93. <http://dx.doi.org/10.1016/j.jclinepi.2011.01.012>
51. Guyatt GH, Oxman AD, Kunz R, Woodcock J, Brozek J, Helfand M, et al. GRADE guidelines 8. Rating the quality of evidence – indirectness. *J Clin Epidemiol* 2011;**64**:1303–10. <http://dx.doi.org/10.1016/j.jclinepi.2011.04.014>
52. Guyatt GH, Oxman AD, Kunz R, Woodcock J, Brozek J, Helfand M, et al. GRADE guidelines 7. Rating the quality of evidence – inconsistency. *J Clin Epidemiol* 2011;**64**:1294–302. <http://dx.doi.org/10.1016/j.jclinepi.2011.03.017>
53. Guyatt GH, Oxman AD, Montori V, Vist G, Kunz R, Brozek J, et al. GRADE guidelines 5. Rating the quality of evidence – publication bias. *J Clin Epidemiol* 2011;**64**:1277–82. <http://dx.doi.org/10.1016/j.jclinepi.2011.01.011>
54. Guyatt GH, Oxman AD, Vist G, Kunz R, Brozek J, Alonso-Coello P, et al. GRADE guidelines 4. Rating the quality of evidence – study limitations (risk of bias). *J Clin Epidemiol* 2011;**64**:407–15. <http://dx.doi.org/10.1016/j.jclinepi.2010.07.017>
55. Jones B, Roger J, Lane PW, Lawton A, Fletcher C, Cappelleri JC, et al. Statistical approaches for conducting network meta-analysis in drug development. *Pharmaceut Stat* 2011;**10**:523–31. <http://dx.doi.org/10.1002/pst.533>
56. Salanti G, Higgins JP, Ades AE, Ioannidis JP. Evaluation of networks of randomized trials. *Stat Methods Med Res* 2008;**17**:279–301. <http://dx.doi.org/10.1177/0962280207080643>
57. Abas MN, Tan PC, Azmi N, Omar SZ. Ondansetron compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2014;**123**:1272–9. <http://dx.doi.org/10.1097/AOG.0000000000000242>
58. Adamczak J, Kasdaglis J, Rinehart B, Antebi Y, Wolf E, Terrone D. A prospective randomized trial of solumedrol dose pack vs. Phenergan for the treatment of symptomatic nausea and vomiting in pregnancy. *Am J Obstet Gynecol* 2007;**197**:S88. <http://dx.doi.org/10.1016/j.ajog.2007.10.292>
59. Babaei AH, Foghaha MH. A randomized comparison of vitamin B6 and dimenhydrinate in the treatment of nausea and vomiting in early pregnancy. *Iran J Nurs Midwifery Res* 2014;**19**:199–202.
60. Basirat Z, Moghadamnia AA, Kashifard M, Sarifi-Razavi A. The effect of ginger biscuit on nausea and vomiting in early pregnancy. *Acta Med Iran* 2009;**47**:51–6.

61. Bayreuther J, Lewith GT, Pickering R. A double-blind cross-over study to evaluate the effectiveness of acupressure at pericardium 6 (P6) in the treatment of early morning sickness (EMS). *Complement Ther Med* 1994;**2**:70–6. [http://dx.doi.org/10.1016/0965-2299\(94\)90002-7](http://dx.doi.org/10.1016/0965-2299(94)90002-7)
62. Belluomini J, Litt RC, Lee KA, Katz M. Acupressure for nausea and vomiting of pregnancy: a randomized, blinded study. *Obstet Gynecol* 1994;**84**:245–8.
63. Biswas SC, Dey R, Kamliya GS, Bal R, Hazra A, Tripathi SK. A single-masked, randomized, controlled trial of ginger extract in the treatment of nausea and vomiting of pregnancy. *JIMSA* 2011;**24**:167–9.
64. Bondok RS, El Sharnouby NM, Eid HE, Abd Elmaksoud AM. Pulsed steroid therapy is an effective treatment for intractable hyperemesis gravidarum. *Crit Care Med* 2006;**34**:2781–3. <http://dx.doi.org/10.1097/01.CCM.0000242156.15757.70>
65. Capp S, Oliveira L, Carstairs S, You W. Ondansetron versus doxylamine/pyridoxine for treatment of nausea and vomiting in pregnancy: a prospective randomized double-blind trial. *Am J Obstet Gynecol* 2014;**210**:S39. <http://dx.doi.org/10.1016/j.ajog.2013.10.092>
66. Carlsson CP, Axemo P, Bodin A, Carstensen H, Ehrenroth B, Madegard-Lind I, *et al*. Manual acupuncture reduces hyperemesis gravidarum: a placebo-controlled, randomized, single-blind, crossover study. *J Pain Symptom Manage* 2000;**20**:273–9. [http://dx.doi.org/10.1016/S0885-3924\(00\)00185-8](http://dx.doi.org/10.1016/S0885-3924(00)00185-8)
67. Chittumma P, Kaewkiattikun K, Wiriyasiriwach B. Comparison of the effectiveness of ginger and vitamin B6 for treatment of nausea and vomiting in early pregnancy: a randomized double-blind controlled trial. *J Med Assoc Thai* 2007;**90**:15–20.
68. Diggory PL, Tomkinson JS. Nausea and vomiting in pregnancy. A trial of meclizine dihydrochloride with and without pyridoxine. *Lancet* 1962;**2**:370–2. [http://dx.doi.org/10.1016/S0140-6736\(62\)90228-3](http://dx.doi.org/10.1016/S0140-6736(62)90228-3)
69. Ditto A, Morgante G, la Marca A, De Leo V. Evaluation of treatment of hyperemesis gravidarum using parenteral fluid with or without diazepam. A randomized study. *Gynecol Obstet Invest* 1999;**48**:232–6. <http://dx.doi.org/10.1159/000010189>
70. Ensiyeh J, Sakineh MA. Comparing ginger and vitamin B6 for the treatment of nausea and vomiting in pregnancy: a randomised controlled trial. *Midwifery* 2009;**25**:649–53. <http://dx.doi.org/10.1016/j.midw.2007.10.013>
71. Erez S, Schifrin BS, Dirim O. Double-blind evaluation of hydroxyzine as an antiemetic in pregnancy. *J Reprod Med* 1971;**7**:35–7.
72. Eftekhari N, Mehralhasani Y. A comparison of ondansetron and promethasin in treating hyperemesis gravidarum. *JKUMS* 2013;**20**:354–65.
73. Evans AT, Samuels SN, Marshall C, Bertolucci LE. Suppression of pregnancy-induced nausea and vomiting with sensory afferent stimulation. *J Reprod Med* 1993;**38**:603–6.
74. Fischer-Rasmussen W, Kjaer SK, Dahl C, Asping U. Ginger treatment of hyperemesis gravidarum. *Eur J Obstet Gynecol Reprod Biol* 1991;**38**:19–24. [http://dx.doi.org/10.1016/0028-2243\(91\)90202-V](http://dx.doi.org/10.1016/0028-2243(91)90202-V)
75. Ghahiri AA, Abdi F, Mastoo R, Ghasemi M. The effect of ondansetron and metoclopramide in nausea and vomiting of pregnancy. *JIMS* 2011;**29**:259–65.
76. Ghani RMA, Ibrahim ATA. The effect of aromatherapy inhalation on nausea and vomiting in early pregnancy: a pilot randomized controlled trial. *J Nat Sci Res* 2013;**3**:10–22.

77. Haji Seid Javadi E, Salehi F, Mashrabi O. Comparing the effectiveness of vitamin B6 and ginger in treatment of pregnancy-induced nausea and vomiting. *Obstet Gynecol Int* 2013;927834. <http://dx.doi.org/10.1155/2013/927834>
78. Heazell A, Thorneycroft J, Walton V, Etherington I. Acupressure for the in-patient treatment of nausea and vomiting in early pregnancy: a randomized control trial. *Am J Obstet Gynecol* 2006;**194**:815–20. <http://dx.doi.org/10.1016/j.ajog.2005.08.042>
79. Hsu E, Pei V, Shofer FS, Abbuhl SB. A prospective randomized controlled trial of acupressure vs. sham for pregnancy-related nausea and vomiting in the emergency department. *Acad Emerg Med* 2003;**10**:437. <http://dx.doi.org/10.1197/aemj.10.5.437>
80. Jamigorn M, Phupong V. Acupressure and vitamin B6 to relieve nausea and vomiting in pregnancy: a randomized study. *Arch Gynecol Obstet* 2007;**276**:245–9. <http://dx.doi.org/10.1007/s00404-007-0336-2>
81. Kashifard M, Basirat Z, Kashifard M, Golsorkhtabar-Amiri M, Moghaddamnia A. Ondansetron or metoclopramide? Which is more effective in severe nausea and vomiting of pregnancy? A randomized trial double-blind study. *Clin Exp Obstet Gynecol* 2013;**40**:127–30.
82. Keating A, Chez RA. Ginger syrup as an antiemetic in early pregnancy. *Altern Ther Health Med* 2002;**8**:89–91.
83. Knight B, Mudge C, Openshaw S, White A, Hart A. Effect of acupuncture on nausea of pregnancy: a randomized, controlled trial. *Obstet Gynecol* 2001;**97**:184–8. [http://dx.doi.org/10.1016/S0029-7844\(00\)01152-2](http://dx.doi.org/10.1016/S0029-7844(00)01152-2)
84. Koren G, Clark S, Hankins GD, Caritis SN, Miodovnik M, Umans JG, et al. Effectiveness of delayed-release doxylamine and pyridoxine for nausea and vomiting of pregnancy: a randomized placebo controlled trial. *Am J Obstet Gynecol* 2010;**203**:571.e1–7. <http://dx.doi.org/10.1016/j.ajog.2010.07.030>
85. Maina A, Arrotta M, Cicogna L, Donvito V, Mischinelli M, Todros T, et al. Transdermal clonidine in the treatment of severe hyperemesis. A pilot randomised control trial: CLONEMESI. *BJOG* 2014;**121**:1556–62. <http://dx.doi.org/10.1111/1471-0528.12757>
86. Maltepe C, Koren G. The management of nausea and vomiting of pregnancy and hyperemesis gravidarum – a 2013 update. *J Popul Ther Clin Pharmacol* 2013;**20**:e184–92.
87. Mao ZN, Liang CE. Observation on therapeutic effect of acupuncture on hyperemesis gravidarum. *Zhongguo Zhen Jiu* 2009;**29**:973–6.
88. McParlin C, Carrick-Sen D, Steen IN, Taylor P, Robson SC. Hyperemesis in pregnancy study: a randomised controlled trial of midwife-led 'outpatient' care. *Arch Dis Child Fetal Neonatal Ed* 2008;**93**:Fa9.
89. Mohammadbeigi R, Shahgeibi S, Soufizadeh N, Rezaie M, Farhadifar F. Comparing the effects of ginger and metoclopramide on the treatment of pregnancy nausea. *Pak J Biol Sci* 2011;**14**:817–20. <http://dx.doi.org/10.3923/pjbs.2011.817.820>
90. Monias M. Evaluation of cyclizine with pyridoxine in vomiting of pregnancy. *Mil Med* 1957;**121**:403–4.
91. Naeimi Rad M, Lamyian M, Heshmat R, Jaafarabadi MA, Yazdani S. A randomized clinical trial of the efficacy of KID21 point (youmen) acupressure on nausea and vomiting of pregnancy. *Iran Red Crescent Med J* 2012;**14**:697–701. <http://dx.doi.org/10.5812/ircmj.2153>
92. Narenji F, Delavar M, Rafiei M. Comparison the effects of the ginger fresh root and vitamin B6 on the nausea and vomiting in pregnancy. *IJOGI* 2012;**15**:39–43.

93. Nelson-Piercy C, Fayers P, Swiet M. Randomised, double-blind, placebo-controlled trial of corticosteroids for the treatment of hyperemesis gravidarum. *BJOG* 2001;**108**:9–15. [http://dx.doi.org/10.1016/s0306-5456\(00\)00017-6](http://dx.doi.org/10.1016/s0306-5456(00)00017-6)
94. Neri I, Allais G, Schiapparelli P, Blasi I, Benedetto C, Facchinetti F. Acupuncture versus pharmacological approach to reduce Hyperemesis gravidarum discomfort. *Minerva Ginecol* 2005;**57**:471–5.
95. Oliveira LG, Capp S, You WB, Carstairs SD. Ondansetron versus doxylamine/pyridoxine for treatment of nausea and vomiting in first trimester pregnancy: a prospective randomized double-blind controlled study. *Acad Emerg Med* 2013;**1**:S101. <http://dx.doi.org/10.1097/AOG.0000000000000479>
96. Ozgoli G, Goli M, Simbar M. Effects of ginger capsules on pregnancy, nausea, and vomiting. *J Altern Complement Med* 2009;**15**:243–6. <http://dx.doi.org/10.1089/acm.2008.0406>
97. Pasha H, Behmanesh F, Mohsenzadeh F, Hajahmadi M, Moghadamnia AA. Study of the effect of mint oil on nausea and vomiting during pregnancy. *Iran Red Crescent Med J* 2012;**14**:727–30. <http://dx.doi.org/10.5812/ircmj.3477>
98. Rosen T, de Veciana M, Miller HS, Stewart L, Rebarber A, Slotnick RN. A randomized controlled trial of nerve stimulation for relief of nausea and vomiting in pregnancy. *Obstet Gynecol* 2003;**102**:129–35. [http://dx.doi.org/10.1016/S0029-7844\(03\)00375-2](http://dx.doi.org/10.1016/S0029-7844(03)00375-2)
99. Safari HR, Fassett MJ, Souter IC, Alsulyman OM, Goodwin TM. The efficacy of methylprednisolone in the treatment of hyperemesis gravidarum: a randomized, double-blind, controlled study. *Am J Obstet Gynecol* 1998;**179**:921–4. [http://dx.doi.org/10.1016/S0002-9378\(98\)70189-9](http://dx.doi.org/10.1016/S0002-9378(98)70189-9)
100. Sahakian V, Rouse D, Sipes S, Rose N, Niebyl J. Vitamin B6 is effective therapy for nausea and vomiting of pregnancy: a randomized, double-blind placebo-controlled study. *Obstet Gynecol* 1991;**78**:33–6.
101. Smith C, Crowther C, Beilby J. Acupuncture to treat nausea and vomiting in early pregnancy: a randomized controlled trial. *Birth* 2002;**29**:1–9. <http://dx.doi.org/10.1046/j.1523-536X.2002.00149.x>
102. Smith C, Crowther C, Willson K, Hotham N, McMillian V. A randomized controlled trial of ginger to treat nausea and vomiting in pregnancy. *Obstet Gynecol* 2004;**103**:639–45. <http://dx.doi.org/10.1097/01.AOG.0000118307.19798.ec>
103. Sripramote M, Lekhyananda N. A randomized comparison of ginger and vitamin B6 in the treatment of nausea and vomiting in pregnancy. *J Med Assoc Thai* 2003;**86**:846–53.
104. Steele NM, French J, Gatherer-Boyles J, Newman S, Leclair S. Effect of acupressure by sea-bands on nausea and vomiting of pregnancy. *J Obstet Gynecol Neonatal Nurs* 2001;**30**:61–70. <http://dx.doi.org/10.1111/j.1552-6909.2001.tb01522.x>
105. Sullivan CA, Johnson CA, Roach H, Martin RW, Stewart DK, Morrison JC. A pilot study of intravenous ondansetron for hyperemesis gravidarum. *Am J Obstet Gynecol* 1996;**174**:1565–8. [http://dx.doi.org/10.1016/S0002-9378\(96\)70607-5](http://dx.doi.org/10.1016/S0002-9378(96)70607-5)
106. Tan PC, Khine PP, Vallikkannu N, Omar SZ. Promethazine compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2010;**115**:975–81. <http://dx.doi.org/10.1097/AOG.0b013e3181d99290>
107. Tan PC, Yow CM, Omar SZ. A placebo-controlled trial of oral pyridoxine in hyperemesis gravidarum. *Gynecol Obstet Invest* 2009;**67**:151–7. <http://dx.doi.org/10.1159/000181182>

108. Tan PC, Norazilah MJ, Omar SZ. Dextrose saline compared with normal saline rehydration of hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2013;**121**:291–8. <http://dx.doi.org/10.1097/AOG.0b013e31827c5e99>
109. Veciana M, Stewart L, Miller H, Slotnick R, Rebarber A, Rosen T. Multicenter randomized controlled trial of nerve stimulation therapy for the relief of nausea and vomiting in pregnancy. *Am J Obstet Gynecol* 2001;**185**:S182. [http://dx.doi.org/10.1016/S0002-9378\(01\)80402-6](http://dx.doi.org/10.1016/S0002-9378(01)80402-6)
110. Vutyavanich T, Kraissarin T, Ruangsri R. Ginger for nausea and vomiting in pregnancy: randomized, double-masked, placebo-controlled trial. *Obstet Gynecol* 2001;**97**:577–82. [http://dx.doi.org/10.1016/S0029-7844\(00\)01228-X](http://dx.doi.org/10.1016/S0029-7844(00)01228-X)
111. Werntoft E, Dykes AK. Effect of acupressure on nausea and vomiting during pregnancy. A randomized, placebo-controlled, pilot study. *J Reprod Med* 2001;**46**:835–9.
112. Wibowo N, Purwosunu Y, Sekizawa A, Farina A, Tambunan V, Bardosono S. Vitamin B6 supplementation in pregnant women with nausea and vomiting. *Int J Gynaecol Obstet* 2012;**116**:206–10. <http://dx.doi.org/10.1016/j.ijgo.2011.09.030>
113. Willetts KE, Ekangaki A, Eden JA. Effect of a ginger extract on pregnancy-induced nausea: a randomised controlled trial. *Aust N Z J Obstet Gynaecol* 2003;**43**:139–44. <http://dx.doi.org/10.1046/j.0004-8666.2003.00039.x>
114. Yost NP, McIntire DD, Wians FH Jr, Ramin SM, Balko JA, Leveno KJ. A randomized, placebo-controlled trial of corticosteroids for hyperemesis due to pregnancy. *Obstet Gynecol* 2003;**102**:1250–4. <http://dx.doi.org/10.1016/j.obstetgynecol.2003.08.013>
115. Zhang HH. Observation on therapeutic effect of acupuncture and moxibustion on hyperemesis gravidarum. *Zhongguo Zhen Jiu* 2005;**25**:469–70.
116. Ziaei S, Hosseiny FS, Faghihzadeh S. The efficacy low dose of prednisolone in the treatment of hyperemesis gravidarum. *Acta Obstet Gynecol Scand* 2004;**83**:272–5. <http://dx.doi.org/10.1111/j.0001-6349.2004.0141.x>
117. Maltepe C, Koren G. Preemptive treatment of nausea and vomiting of pregnancy: results of a randomized controlled trial. *Obstet Gynecol Int* 2013;809787. <http://dx.doi.org/10.1155/2013/809787>
118. Smith C, Crowther C, Beilby J. Pregnancy outcome following women's participation in a randomised controlled trial of acupuncture to treat nausea and vomiting in early pregnancy. *Complement Ther Med* 2002;**10**:78–83. <http://dx.doi.org/10.1054/ctim.2002.0523>
119. Biswas SC, Bal R, Kamilya GS, Mukherjee J, Hazra A, Narender S, et al. A single-masked, randomized, controlled trial of ginger extract in the treatment of nausea and vomiting of pregnancy. 49th All India Congress of Obstetrics and Gynaecology: Cochin, Kerala State, India; 6–9 January 2006. URL: <http://onlinelibrary.wiley.com/doi/10.1002/1471-0528.2004.00236.x> (accessed 28 November 2014).
120. Alalade AO, Khan R, Dawlatly B. Day-case management of hyperemesis gravidarum: feasibility and clinical efficacy. *J Obstet Gynaecol* 2007;**27**:363–4. <http://dx.doi.org/10.1080/01443610701327396>
121. Einarson A, Maltepe C, Navioz Y, Kennedy D, Tan MP, Koren G. The safety of ondansetron for nausea and vomiting of pregnancy: a prospective comparative study. *BJOG* 2004;**111**:940–3. <http://dx.doi.org/10.1111/j.1471-0528.2004.00236.x>
122. Ferreira E, Bussieres JF, Turcotte V, Duperron L, Ouellet G. Case-control study comparing droperidol plus diphenhydramine with conventional treatment in hyperemesis gravidarum. *JPT* 2003;**19**:349–54. <http://dx.doi.org/10.1177/875512250301900602>

123. Hsu JJ, Clark-Glena R, Nelson DK, Kim CH. Nasogastric enteral feeding in the management of hyperemesis gravidarum. *Obstet Gynecol* 1996;**88**:343–6. [http://dx.doi.org/10.1016/0029-7844\(96\)00174-3](http://dx.doi.org/10.1016/0029-7844(96)00174-3)
124. Markose MT, Ramanathan K, Vijayakumar J. Reduction of nausea, vomiting, and dry retches with P6 acupressure during pregnancy. *Int J Gynaecol Obstet* 2004;**85**:168–9. <http://dx.doi.org/10.1016/j.ijgo.2003.09.008>
125. Moran P, Taylor R. Management of hyperemesis gravidarum: the importance of weight loss as a criterion for steroid therapy. *QJM* 2002;**95**:153–8. <http://dx.doi.org/10.1093/qjmed/95.3.153>
126. Saha S, Loranger D, Pricolo V, Degli-Esposti S. Feeding jejunostomy for the treatment of severe hyperemesis gravidarum: a case series. *J Parenter Enteral Nutr* 2009;**33**:529–34. <http://dx.doi.org/10.1177/0148607109333000>
127. Koren G, Maltepe C. Preemptive Diclectin therapy for the management of nausea and vomiting of pregnancy and hyperemesis gravidarum. *Am J Obstet Gynecol* 2013;**208**:S20. <http://dx.doi.org/10.1016/j.ajog.2012.10.205>
128. Safari HR, Fassett MJ, Souter IC, Goodwin TM. Randomized double-blind trial of methylprednisolone versus promethazine in the treatment of hyperemesis gravidarum. *Am J Obstet Gynecol* 1998;**178**:S60.
129. Yang LC, Jawan B, Chen CN, Ho RT, Chang KA, Lee JH. Comparison of P6 acupoint injection with 50% glucose in water and intravenous droperidol for prevention of vomiting after gynecological laparoscopy. *Acta Anaesthesiol Scand* 1993;**37**:192–4. <http://dx.doi.org/10.1111/j.1399-6576.1993.tb03699.x>
130. Evers S, Goossens M, de Vet H, van Tulder M, Ament A. Criteria list for assessment of methodological quality of economic evaluations: consensus on health economic criteria. *Int J Technol Assess Health Care* 2005;**21**:240–5.
131. Philips Z, Bojke L, Sculpher M, Claxton K, Golder S. Good practice guidelines for decision-analytic modelling in health technology assessment: a review and consolidation of quality assessment. *Pharmacoeconomics* 2006;**24**:355–71. <http://dx.doi.org/10.2165/00019053-200624040-00006>
132. Joint Formulary Committee. *British National Formulary*. 65th ed. London: BMJ Group and Pharmaceutical Press; 2014.
133. Department of Health. *Payment by Results in the NHS: Tariff for 2012 to 2013*. URL: www.gov.uk/government/publications/confirmation-of-payment-by-results-Pbr-arrangements-for-2012-13 (accessed 28 November 2014).
134. Curtis L. *Unit Costs of Health and Social Care 2013*. University of Kent: Personal Social Services Research Unit; 2013.
135. Grimshaw J, Thomas R, MacLennan G, Fraser C, Ramsay C, Vale L, et al. Effectiveness and efficiency of guideline dissemination and implementation strategies. *Health Technol Assess* 2004;**8**(6). <http://dx.doi.org/10.3310/hta8060>
136. Portnoi G, Chng L, Karimi-Tabesh L, Koren G, Tan MP, Einarson A. Prospective comparative study of the safety and effectiveness of ginger for the treatment of nausea and vomiting in pregnancy. *Am J Obstet Gynecol* 2003;**190**:1140–374. [http://dx.doi.org/10.1067/s0002-9378\(03\)00649-5](http://dx.doi.org/10.1067/s0002-9378(03)00649-5)
137. Heitmann K, Nordeng H, Holst L. Safety of ginger use in pregnancy: results from a large population-based cohort study. *Eur J Clin Pharmacol* 2013;**69**:269–77. <http://dx.doi.org/10.1007/s00228-012-1331-5>
138. Einarson T, Leeder J, Koren G. A method for metaanalysis of epidemiological studies. *Drug Intell Clin Pharm* 1988;**22**:813–24.

139. McKeigue PM, Lamm SH, Linn S, Kutcher JS. Bendectin and birth defects: I. A meta-analysis of the epidemiologic studies. *Teratology* 1994;**50**:27–37. <http://dx.doi.org/10.1002/tera.1420500105>
140. Seto A, Einarson T, Koren G. Pregnancy outcome following first trimester exposure to antihistamines: meta-analysis. *Am J Perinatol* 1997;**14**:119–24. <http://dx.doi.org/10.1055/s-2007-994110>
141. Mazzotta P, Magee LA. A risk-benefit assessment of pharmacological and nonpharmacological treatments for nausea and vomiting of pregnancy. *Drugs* 2000;**59**:781–800. <http://dx.doi.org/10.2165/00003495-200059040-00005>
142. Matok I, Gorodischer R, Koren G, Sheiner E, Wiznitzer A, Levy A. The safety of metoclopramide use in the first trimester of pregnancy. *N Engl J Med* 2009;**360**:2528–35. <http://dx.doi.org/10.1056/NEJMoa0807154>
143. Pasternak B, Svanstrom H, Molgaard-Nielsen D, Melbye M, Hviid A. Metoclopramide in pregnancy and risk of major congenital malformations and fetal death. *JAMA* 2013;**310**:1601–11. <http://dx.doi.org/10.1001/jama.2013.278343>
144. Anderka M, Mitchell AA, Louik C, Werler MM, Hernandez-Diaz S, Rasmussen SA, *et al.* Medications used to treat nausea and vomiting of pregnancy and the risk of selected birth defects. *Birth Defects Res A Clin Mol Teratol* 2012;**94**:22–30. <http://dx.doi.org/10.1002/bdra.22865>
145. Pasternak B, Svanstrom H, Hviid A. Ondansetron in pregnancy and risk of adverse fetal outcomes. *N Engl J Med* 2013;**368**:814–23. <http://dx.doi.org/10.1056/NEJMoa1211035>
146. Freedman SB, Uleryk E, Rumantir M, Finkelstein Y. Ondansetron and the risk of cardiac arrhythmias: a systematic review and postmarketing analysis. *Ann Emerg Med* 2014;**64**:19–25, e1–6. <http://dx.doi.org/10.1016/j.annemergmed.2013.10.026>
147. Park-Wyllie L, Mazzotta P, Pastuszak A, Moretti M, Beique L, Hunnisett L, *et al.* Birth defects after maternal exposure to corticosteroids: prospective cohort study and meta-analysis of epidemiological studies. *Teratology* 2000;**62**:385–92. [http://dx.doi.org/10.1002/1096-9926\(200012\)62:6<385::AID-TERA5>3.0.CO;2-Z](http://dx.doi.org/10.1002/1096-9926(200012)62:6<385::AID-TERA5>3.0.CO;2-Z)
148. Czeizel AE, Rockenbauer M. Population-based case-control study of teratogenic potential of corticosteroids. *Teratology* 1997;**56**:335–40. [http://dx.doi.org/10.1002/\(SICI\)1096-9926\(199711\)56:5<335::AID-TERA7>3.0.CO;2-W](http://dx.doi.org/10.1002/(SICI)1096-9926(199711)56:5<335::AID-TERA7>3.0.CO;2-W)
149. Tata LJ, Lewis SA, McKeever TM, Smith CJP, Doyle P, Smeeth L, *et al.* Effect of maternal asthma, exacerbations and asthma medication use on congenital malformations in offspring: a UK population-based study. *Thorax* 2008;**63**:981–987. <http://dx.doi.org/10.1136/thx.2008.098244>
150. Guttuso T Jr, Shaman M, Thornburg LL. Potential maternal symptomatic benefit of gabapentin and review of its safety in pregnancy. *Eur J Obstet Gynecol Reprod Biol* 2014;**181**:280–3. <http://dx.doi.org/10.1016/j.ejogrb.2014.08.013>
151. McCarthy FP, Murphy A, Khashan AS, McElroy B, Spillane N, Marchocki Z, *et al.* Day care compared with inpatient management of nausea and vomiting of pregnancy: a randomized controlled trial. *Obstet Gynecol* 2014;**124**:743–8. <http://dx.doi.org/10.1097/AOG.0000000000000449>
152. UK Teratology Information Service (UKTIS). *Use of Ginger in Pregnancy*. London: UKTIS; 2013.
153. UK Teratology Information Service (UKTIS). *Use of Vitamin B12 in Pregnancy*. London: UKTIS; 2013.
154. UK Teratology Information Service (UKTIS). *Use of Vitamin B6 in Pregnancy*. London: UKTIS; 2011.
155. UK Teratology Information Service (UKTIS). *Use of Promethazine in Pregnancy*. London: UKTIS; 2010.
156. UK Teratology Information Service (UKTIS). *Use of Ondansetron in Pregnancy*. London: UKTIS; 2014.

Appendix 1 Examples of hyperemesis gravidarum/nausea and vomiting in pregnancy assessment tools

This assessment has been reproduced with permission from Professor Gideon Koren, Director, The Motherisk Program; Professor of Pediatrics, Pharmacology, Pharmacy, Medicine and Medical Genetics at The University of Toronto; Senior Scientist at The Research Institute Hospital for Sick Children; and The Ivey Chair in Molecular Toxicology at the University of Western Ontario Canada, 2014, personal communication.

Motherisk PUQE-24 scoring system

Please fill out the Motherisk PUQE Scoring System for the last 24 hours (*please tick box and write total score*)

1. In the last 24 hours, for how long have you felt nauseated or sick at your stomach,	Not at all (1)	1 hour or less (2)	2-3 hours (3)	4-6 hours (4)	More than 6 hours (5)	Total hrs	Mild: ≤ 6 Moderate: 7-12 Severe: ≥13
2. In the last 24 hours, have you vomited or thrown up,	7 or more times (5)	5-6 (4)	3-4 (3)	1-2 (2)	I did not throw up (1)	Total #	
3. In the last 24 hours, how many times have you had retching or dry heaves without bringing anything up,	No time (1)	1-2 (2)	3-4 (3)	5-6 (4)	7 or more (5)	Total #	Total score: _____

How many hours have you slept out of 24 hours? _____ Why?

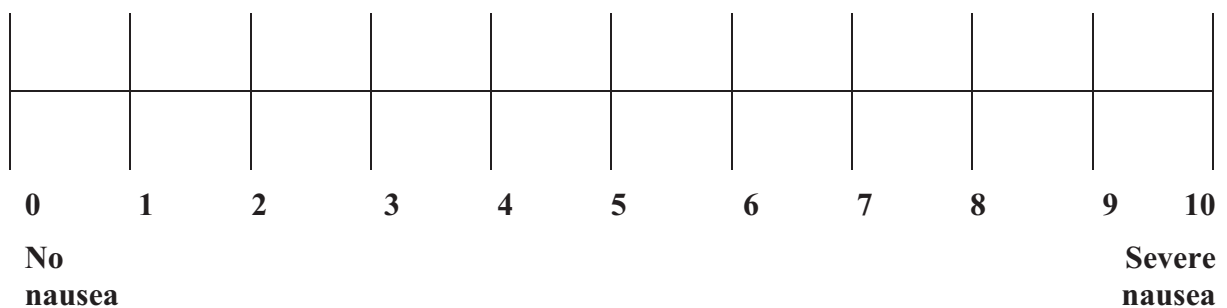
On a scale of 0-10, how would you rate your *Well Being*?

0 (Worst possible) _____ 10 (The best you felt before pregnancy)

Can you tell me what causes you to feel that way?

Likert scale example (6-point)

	Frequency					
	0 (not at all)	1 (occasionally)	2 (3-6 days during the week)	3 (daily)	4 (more than once a day)	5 (all the time)
(1) How often have you felt like being sick (nauseous) in the past week?						
(2) How often have you retched (but without actually being sick) in the past week?						
(3) How often have you been physically sick during the past week?						

Visual Analogue Scale

Appendix 2 Data abstraction form: clinical effectiveness

Reviewer ID:

Data extraction date:

GENERAL INFORMATION	
Report title:	
First author / contact details	
Publication year	
Publication status:	Full-text paper <input type="checkbox"/> Conference abstract <input type="checkbox"/> Personal communication <input type="checkbox"/> Other unpublished reports <input type="checkbox"/>
Journal yy:vol(issue):pp	
Language (if non-English):	
Study IDs of any linked reports:	
Study funding sources (including role of funders)	
Possible conflicts of interest (for study authors)	
STUDY ELIGIBILITY	
Type of study:	RCT <input type="checkbox"/> Non-randomised comparative study <input type="checkbox"/> Case series <input type="checkbox"/>
Type of intervention:	<p>Dietary / lifestyle:</p> <p>Vitamin B6 <input type="checkbox"/> Vitamin B12 <input type="checkbox"/> Ginger <input type="checkbox"/> Acupuncture <input type="checkbox"/> Acupressure <input type="checkbox"/> Hypnosis <input type="checkbox"/></p> <p>Antiemetic drugs:</p> <p>Antihistamines <input type="checkbox"/> Dopamine antagonists <input type="checkbox"/> 5-HT receptor antagonists <input type="checkbox"/> Corticosteroids <input type="checkbox"/> Doxylamine-Pyridoxine <input type="checkbox"/> Other (details below) <input type="checkbox"/> Intravenous fluids: <input type="checkbox"/></p> <p>Enteral and total parenteral nutrition</p> <p>Enteral feeding <input type="checkbox"/> Total parenteral nutrition <input type="checkbox"/></p> <p>Other Intervention <input type="checkbox"/> _____</p>
Comparator:	No treatment <input type="checkbox"/> Treatment as usual (details below) <input type="checkbox"/> Alternative intervention (details below) <input type="checkbox"/> _____
Comparator not applicable:	

Women with severe symptoms	<input type="checkbox"/>
Participants:	
Gestational age ≤ 20 weeks	<input type="checkbox"/>
Symptom severity:	
PUQE score ≥ 13	<input type="checkbox"/>
Rhodes score ≥ 33	<input type="checkbox"/>
Author defined scale (provide details below)	<input type="checkbox"/>
Percentage experiencing symptoms $>80\%$	<input type="checkbox"/>
Primary outcomes:	
Severity of symptoms:	
PUQE	<input type="checkbox"/>
Rhodes Index	<input type="checkbox"/>
McGill Nausea Questionnaire	<input type="checkbox"/>
Visual Analogue scales	<input type="checkbox"/>
Nausea and Vomiting of Pregnancy Instrument	<input type="checkbox"/>
Secondary outcomes:	
Maternal-physical:	
Admission/readmission rate	<input type="checkbox"/>
Length of hospital stay	<input type="checkbox"/>
Antiemetic / other medication use	<input type="checkbox"/>
Amount/duration IV fluid administration	<input type="checkbox"/>
Enteral/total Parenteral nutrition	<input type="checkbox"/>
Adverse events	<input type="checkbox"/>
Economic costs (hospital/medical care)	<input type="checkbox"/>
Adverse pregnancy outcomes	<input type="checkbox"/>
Weight loss	<input type="checkbox"/>
Therapeutic termination of pregnancy	<input type="checkbox"/>
Other author defined NVP scale	<input type="checkbox"/>
Maternal –psychosocial:	
Quality of life (eg. SF-12/SF-36 score)	<input type="checkbox"/>
General Health Questionnaire	<input type="checkbox"/>
Pregnancy specific QoL instrument	<input type="checkbox"/>
NVP specific questionnaire	<input type="checkbox"/>
Satisfaction with care	<input type="checkbox"/>
Direct costs to woman/family	<input type="checkbox"/>
Time lost from work	<input type="checkbox"/>
Edinburgh post natal depression score	<input type="checkbox"/>
Fetal/Neonatal:	
Congenital abnormality	<input type="checkbox"/>
Low birth weight ($<2.5\text{kg}$)	<input type="checkbox"/>
Small for gestational age ($<10^{\text{th}}$ centile)	<input type="checkbox"/>
Pre-term birth (<37 weeks gestation)	<input type="checkbox"/>
5 minute APGAR	<input type="checkbox"/>
Stillbirth/IUD	<input type="checkbox"/>
Neonatal death	<input type="checkbox"/>
Spontaneous miscarriage	<input type="checkbox"/>
Admission to special care baby unit	<input type="checkbox"/>
Long term infant outcomes	<input type="checkbox"/>
INCLUDE	<input type="checkbox"/>
EXCLUDE	<input type="checkbox"/>
Reasons for exclusion:	

DO NOT PROCEED IF STUDY EXCLUDED FROM REVIEW

ADDITIONAL STUDY INFORMATION			
Population and setting			
	Intervention	Comparator	Location in text
Population description			
Setting (including country / location / social context etc)			
Inclusion exclusion criteria			
Method/s of recruitment			
Informed consent obtained	Yes <input type="checkbox"/> No <input type="checkbox"/> Unclear <input type="checkbox"/>	Yes <input type="checkbox"/> No <input type="checkbox"/> Unclear <input type="checkbox"/>	
Notes:			
Methods			
	Descriptions as stated in report / paper		Location in text
Aim of study			
Design (no of arms)			
Unit of allocation (by individuals, cluster / groups)			
Start date			
End date			
Total study duration			
Ethical approval needed / obtained for study	Yes <input type="checkbox"/>	No <input type="checkbox"/> Unclear <input type="checkbox"/>	
Notes:			
Participant characteristics			
	Intervention	Comparator	Location in text
Number of patients enrolled:			
- Randomised (RCTs only), n (%)			
- Included (RCTs only), n (%)			
- Completed, n (%)			

- Available for follow-up, n (%)			
- Withdrew/lost to follow-up, with reasons, n (%)			
- Number analysed, n (%)			
Age (mean/median, SD/range)			
Ethnicity, n (%)			
Smoking status (give n (%) of smokers)			
BMI at baseline, (mean/median, SD/range)			
Weight at baseline (mean/median, SD range)			
Singleton pregnancy only (if no, give n (%))			
Gestational age at onset (week: mean/median, SD/range)			
Gestational age at primary admission (week: mean/median, SD/range)			
Gestational age at study entry / randomisation (week: mean/median, SD/range)			
Primiparas only (if no, give %)			
Obstetric history (previous NVP)			
Pre-existing medical conditions (please specify)			
NVP / HG Severity:			
- PUQE (Mean/median, SD/range)			
- Rhodes Index (Mean/median, SD/range)			
- McGill Nausea Questionnaire (Mean/median, SD/range)			

- Nausea and Vomiting of Pregnancy Instrument (Mean/median, SD/range)			
- Other scale (details)			
Other baseline characteristics (please specify):			
INTERVENTION GROUPS (<i>copy and paste table for each intervention group and comparator</i>)			
Intervention Group 1			
	Description in text	Location in text	
Group name			
No. randomised to group			
Theoretical basis (include key references)			
Description of intervention (include sufficient detail for replication eg content, dose, components)			
Duration of treatment period			
Timing of treatment (eg frequency, duration of each episode)			
Delivery (eg mechanism, medium, intensity, fidelity)			
Providers (eg number, profession, training, gender / ethnicity / age if relevant)			
Co-interventions			
Economic variables (eg intervention cost, changes in other others as result of intervention)			
Notes:			

OUTCOMES (<i>copy and paste table for each outcome</i>)		
Outcome 1		
	Description as stated in report / paper	Location in text
Outcome name		
Time points measured		
Time points reported		
Outcome definition (with diagnostic criteria if relevant)		
Personal measuring/ reporting		
Unit of measurement (if relevant)		
Scales: upper and lower limits (indicate whether high or low score is good)		
Is outcome/tool validated?	Yes <input type="checkbox"/> No <input type="checkbox"/> Unclear <input type="checkbox"/>	
Imputations of missing data (eg assumptions made for ITT analysis)		
Assumed risk estimate (eg baseline or population risk note in background)		
Power		
Notes:		
RESULTS		
Dichotomous outcomes		
	Description as stated in report/paper	Location in text
Comparison		
Outcome		
Subgroup		
Timepoint (specify whether from start or end of intervention)		
Results	Intervention	Comparison

	No. events	No. participants	No. events	No. participants			
No. missing participants and reasons							
No. participants moved from other group and reasons							
Any other results reported							
Unit of analysis (by individuals, cluster/groups or body parts)							
Statistical methods used							
Notes:							
Continuous outcome							
	Description as stated in report/paper				Location in text		
Comparison							
Outcome							
Subgroup							
Timepoint (specify whether from start or end of intervention)							
Post-intervention or change from baseline?							
Results	Intervention			Comparison			
	Mean	SD (or other variance)	No. participants	Mean	SD (or other variance)	No. participants	

No. missing participants and reasons				
No. participants moved from other group and reasons				
Any other results reported				
Unit of analysis (individuals, cluster/groups or body parts)				
Statistical methods used				
Notes:				
Other outcome				
	Description as stated in report/paper			Location in text
Comparison				
Outcome				
Subgroup				
Timepoint (specify whether from start or end of intervention)				
Results	Intervention result	SD (or other variance)	Control result	SD (or other variance)
	Overall results		SE (or other variance)	
No. participants	Intervention		Control	
No. missing participants and reasons				
No. participants moved from other group and reasons				

Any other results reported		
Unit of analysis (by individuals, cluster/groups or body parts)		
Statistical methods used		
Notes:		
Conclusion as reported by the authors of the study		
Additional information and comments		

Appendix 3 Risk of bias for randomised controlled trials

Domain	Description	Review authors' judgement
Sequence generation	Describe the method used to generate the allocation sequence in sufficient detail to allow an assessment of whether or not it should produce comparable groups	Was the allocation sequence adequately generated?
Allocation concealment	Describe the method used to conceal the allocation sequence in sufficient detail to determine whether or not intervention allocations could have been foreseen in advance of, or during, enrolment	Was allocation adequately concealed?
Blinding of participants, personnel and outcome assessors	Describe all measures used, if any, to blind study participants and personnel from knowledge of which intervention a participant received. Provide any information relating to whether the intended blinding was effective	Was knowledge of the allocated intervention adequately prevented during the study?
<i>Assessments should be made for each main outcome (or class of outcomes)</i>		
Incomplete outcome data	Describe the completeness of outcome data for each main outcome, including attrition and exclusions from the analysis. State whether attrition and exclusions were reported, the numbers in each intervention group (compared with total randomised participants), reasons for attrition/exclusions where reported, and any reinclusions in analyses performed by the review authors	Were incomplete outcome data adequately addressed?
<i>Assessments should be made for each main outcome (or class of outcomes)</i>		
Selective outcome reporting	State how the possibility of selective outcome reporting was examined by the review authors, and what was found	Are reports of the study free of suggestion of selective outcome reporting?
Other sources of bias	State any important concerns about bias not addressed in the other domains in the tool If particular questions/entries were pre-specified in the review's protocol, responses should be provided for each question/entry	Was the study apparently free of other problems that could put it at a high risk of bias?

Possible approach for summary assessments outcome (across domains) within and across studies

Risk of bias	Interpretation	Within a study	Across studies
Low risk of bias	Plausible bias unlikely to seriously alter the results	Low risk of bias for all key domains	Most information is from studies at low risk of bias
Unclear risk of bias	Plausible bias that raises some doubt about the results	Unclear risk of bias for one or more key domains	Most information is from studies at low or unclear risk of bias
High risk of bias	Plausible bias that seriously weakens confidence in the results	High risk of bias for one or more key domains	The proportion of information from studies at high risk of bias is sufficient to affect the interpretation of the results

Criteria for judging risk of bias in the 'risk of bias' assessment tool

Sequence generation

Was the allocation sequence adequately generated? [Short form: *Adequate sequence generation?*]

Criteria for a judgement of 'YES' (i.e. low risk of bias)

The investigators describe a random component in the sequence generation process such as:

- referring to a random number table; using a computer random number generator; coin tossing; shuffling cards or envelopes; throwing dice; drawing of lots; minimisation^a

Criteria for the judgement of 'NO' (i.e. high risk of bias)

The investigators describe a non-random component in the sequence generation process. Usually, the description would involve some systematic, non-random approach, for example:

- sequence generated by odd or even date of birth
- sequence generated by some rule based on date (or day) of admission
- sequence generated by some rule based on hospital or clinic record number

Other non-random approaches happen much less frequently than the systematic approaches mentioned above and tend to be obvious. They usually involve judgement or some method of non-random categorisation of participants, for example:

- allocation by judgement of the clinician
- allocation by preference of the participant
- allocation based on the results of a laboratory test or a series of tests
- allocation by availability of the intervention

Criteria for the judgement of 'UNCLEAR' (uncertain risk of bias)

Insufficient information about the sequence generation process to permit judgement of 'YES' or 'NO'

Allocation concealment

Was allocation adequately concealed? [Short form: *Allocation concealment?*]

Criteria for a judgement of 'YES' (i.e. low risk of bias)

Participants and investigators enrolling participants could not foresee assignment because one of the following, or an equivalent method, was used to conceal allocation:

- central allocation (including telephone, web-based, and pharmacy-controlled, randomisation)
- sequentially numbered drug containers of identical appearance
- sequentially numbered, opaque, sealed envelopes

Criteria for the judgement of 'NO' (i.e. high risk of bias)

Participants or investigators enrolling participants could possibly foresee assignments and thus introduce selection bias, such as allocation based on:

- using an open random allocation schedule (e.g. a list of random numbers)
- assignment envelopes were used without appropriate safeguards (e.g. if envelopes were unsealed or non-opaque or not sequentially numbered)
- alternation or rotation
- date of birth
- case record number
- any other explicitly unconcealed procedure

Criteria for the judgement of 'UNCLEAR' (uncertain risk of bias)

Insufficient information to permit judgement of 'YES' or 'NO'. This is usually the case if the method of concealment is not described or not described in sufficient detail to allow a definite judgement; for example, if the use of assignment envelopes is described, but it remains unclear whether envelopes were sequentially numbered, opaque and sealed

Blinding of participants, personnel and outcome assessors

Was knowledge of the allocated interventions adequately prevented during the study? [Short form: *Blinding?*]

Criteria for a judgement of 'YES' (i.e. low risk of bias)	Any one of the following: <ul style="list-style-type: none"> • no blinding, but the review authors judge that the outcome and the outcome measurement are not likely to be influenced by lack of blinding • blinding of participants and key study personnel ensured, and unlikely that the blinding could have been broken • either participants or some key study personnel were not blinded, but outcome assessment was blinded and the non-blinding of others unlikely to introduce bias
Criteria for the judgement of 'NO' (i.e. high risk of bias)	Any one of the following: <ul style="list-style-type: none"> • no blinding or incomplete blinding, and the outcome or outcome measurement is likely to be influenced by lack of blinding • blinding of key study participants and personnel attempted, but likely that the blinding could have been broken • either participants or some key study personnel were not blinded, and the non-blinding of others likely to introduce bias
Criteria for the judgement of 'UNCLEAR' (uncertain risk of bias)	Any one of the following: <ul style="list-style-type: none"> • insufficient information to permit judgement of 'YES' or 'NO' • the study did not address this outcome

Incomplete outcome data

Were incomplete outcome data adequately addressed? [Short form: *Incomplete outcome data addressed?*]

Criteria for a judgement of 'YES' (i.e. low risk of bias)	Any one of the following: <ul style="list-style-type: none"> • no missing outcome data • reasons for missing outcome data unlikely to be related to true outcome (for survival data, censoring unlikely to be introducing bias) • missing outcome data balanced in numbers across intervention groups, with similar reasons for missing data across groups • for dichotomous outcome data, the proportion of missing outcomes compared with observed event risk not enough to have a clinically relevant impact on the intervention effect estimate • for continuous outcome data, plausible effect size (difference in means or standardised difference in means) among missing outcomes not enough to have a clinically relevant impact on observed effect size • missing data have been imputed using appropriate methods
Criteria for the judgement of 'NO' (i.e. high risk of bias)	Any one of the following: <ul style="list-style-type: none"> • reason for missing outcome data likely to be related to true outcome, with either imbalance in numbers or reasons for missing data across intervention groups • for dichotomous outcome data, the proportion of missing outcomes compared with observed event risk enough to induce clinically relevant bias in intervention effect estimate • For continuous outcome data, plausible effect size (difference in means or standardised difference in means) among missing outcomes enough to induce clinically relevant bias in observed effect size • 'as-treated' analysis done with substantial departure of the intervention received from that assigned at randomisation • potentially inappropriate application of simple imputation
Criteria for the judgement of 'UNCLEAR' (uncertain risk of bias)	Any one of the following: <ul style="list-style-type: none"> • insufficient reporting of attrition/exclusions to permit judgement of 'YES' or 'NO' (e.g. number randomised not stated, no reasons for missing data provided) • The study did not address this outcome

Selective outcome reporting

Are reports of the study free of suggestion of selective outcome reporting? [Short form: *Free of selective reporting?*]

Criteria for a judgement of 'YES' (i.e. low risk of bias)

Any of the following:

- the study protocol is available and all of the study's pre-specified (primary and secondary) outcomes that are of interest in the review have been reported in the pre-specified way
- the study protocol is not available but it is clear that the published reports include all expected outcomes, including those that were pre-specified (convincing text of this nature may be uncommon)

Criteria for the judgement of 'NO' (i.e. high risk of bias)

Any one of the following:

- not all of the study's pre-specified primary outcomes have been reported
- one or more primary outcomes is reported using measurements, analysis methods or subsets of the data (e.g. subscales) that were not pre-specified
- one or more reported primary outcomes were not pre-specified (unless clear justification for their reporting is provided such as an unexpected adverse effect)
- one or more outcomes of interest in the review are reported incompletely so that they cannot be entered in a meta-analysis
- the study report fails to include results for a key outcome that would be expected to have been reported for such a study

Criteria for the judgement of 'UNCLEAR' (uncertain risk of bias)

Insufficient information to permit judgement of 'YES' or 'NO'. It is likely that the majority of studies will fall into this category

Other potential threats to validity

Was the study apparently free of other problems that could put it at a risk of bias? [Short form: *Free of other bias?*]

Criteria for a judgement of 'YES' (i.e. low risk of bias)

The study appears to be free of other sources of bias

Criteria for the judgement of 'NO' (i.e. high risk of bias)

There is at least one important risk of bias. For example, the study:

- had a potential source of bias related to the specific study design used; or
- stopped early due to some data-dependent process (including a formal-stopping rule); or
- had extreme baseline imbalance; or
- has been claimed to have been fraudulent; or
- had some other problem

Criteria for the judgement of 'UNCLEAR' (uncertain risk of bias)

There may be a risk of bias, but there is either:

- insufficient information to assess whether or not an important risk of bias exists; or
- insufficient rationale or evidence that an identified problem will introduce bias

a Minimisation may be implemented without a random element, and this is considered to be equivalent to being random.

Appendix 4 Quality of case series studies

EPHPP Quality Assessment Tool for Quantitative Studies

COMPONENT RATINGS

A) SELECTION BIAS

(Q1) Are the individuals selected to participate in the study likely to be representative of the target population?

- 1 Very likely
- 2 Somewhat likely
- 3 Not likely
- 4 Can't tell

(Q2) What percentage of selected individuals agreed to participate?

- 1 80 - 100% agreement
- 2 60 – 79% agreement
- 3 less than 60% agreement
- 4 Not applicable
- 5 Can't tell

RATE THIS SECTION	STRONG	MODERATE	WEAK
See dictionary	1	2	3

B) STUDY DESIGN

Indicate the study design

- 1 Randomised controlled trial
- 2 Controlled clinical trial

- 3 Cohort analytic (two group pre + post)
- 4 Case-control
- 5 Cohort (one group pre + post (before and after))
- 6 Interrupted time series
- 7 Other specify _____
- 8 Can't tell

Was the study described as randomized? If NO, go to Component C.

No Yes

If Yes, was the method of randomization described? (See dictionary)

No Yes

If Yes, was the method appropriate? (See dictionary)

No Yes

RATE THIS SECTION	STRONG	MODERATE	WEAK
See dictionary	1	2	3

C) CONFOUNDERS

(Q1) Were there important differences between groups prior to the intervention?

Yes

No

Can't tell

The following are examples of confounders:

Race

Sex

Marital status/family

Age

SES (income or class)

Education

Health status

Pre-intervention score on outcome measure

(Q2) If yes, indicate the percentage of relevant confounders that were controlled (either in the design (e.g. stratification, matching) or analysis)?

80 – 100% (most)

60 – 79% (some)

Less than 60% (few or none)

Can't Tell

RATE THIS SECTION	STRONG	MODERATE	WEAK
See dictionary	1	2	3

D) BLINDING

(Q1) Was (were) the outcome assessor(s) aware of the intervention or exposure status of participants?

Yes

No

Can't tell

(Q2) Were the study participants aware of the research question?

Yes

No

Can't tell

RATE THIS SECTION	STRONG	MODERATE	WEAK
See dictionary	1	2	3

E) DATA COLLECTION METHODS

(Q1) Were data collection tools shown to be valid?

Yes

No

Can't tell

(Q2) Were data collection tools shown to be reliable?

Yes

No

Can't tell

RATE THIS SECTION	STRONG	MODERATE	WEAK
See dictionary	1	2	3

F) WITHDRAWALS AND DROP-OUTS

(Q1) Were withdrawals and drop-outs reported in terms of numbers and/or reasons per group?

Yes

No

Can't tell

Not Applicable (i.e. one time surveys or interviews)

(Q2) Indicate the percentage of participants completing the study. (If the percentage differs by groups, record the lowest).

80 -100%

60 - 79%

less than 60%

Can't tell

Not Applicable (i.e. Retrospective case-control)

RATE THIS SECTION	STRONG	MODERATE	WEAK	
See dictionary	1	2	3	Not Applicable

G) INTERVENTION INTEGRITY

(Q1) What percentage of participants received the allocated intervention or exposure of interest?

80 -100%

60 - 79%

less than 60%

Can't tell

(Q2) Was the consistency of the intervention measured?

Yes

No

Can't tell

(Q3) Is it likely that subjects received an unintended intervention (contamination or co-intervention) that may influence the results?

Yes

No

Can't tell

H) ANALYSES

(Q1) Indicate the unit of allocation (circle one)

community organisation/institution practice/office individual

(Q2) Indicate the unit of analysis (circle one)

community organisation/institution practice/office individual

(Q3) Are the statistical methods appropriate for the study design?

Yes

No

Can't tell

(Q4) Is the analysis performed by intervention allocation status (i.e. intention to treat) rather than the actual intervention received? (per protocol)

Yes

No

Can't tell

GLOBAL RATING

COMPONENT RATINGS

Please transcribe the information from the boxes on pages 1-4 onto this page. See dictionary on how to rate this section.

A	SELECTION BIAS	STRONG	MODERATE	WEAK	
		1	2	3	
B	STUDY DESIGN	STRONG	MODERATE	WEAK	
		1	2	3	
C	CONFOUNDERS	STRONG	MODERATE	WEAK	
		1	2	3	
D	BLINDING	STRONG	MODERATE	WEAK	
		1	2	3	
E	DATA COLLECTION METHOD	STRONG	MODERATE	WEAK	
		1	2	3	
F	WITHDRAWALS AND DROPOUTS	STRONG	MODERATE	WEAK	
					Not applicable

GLOBAL RATING FOR THIS PAPER (circle one):

- 1 STRONG (no WEAK ratings)
- 2 MODERATE (one WEAK rating)
- 3 WEAK (two or more WEAK ratings)

With both reviewers discussing the ratings:

Is there a discrepancy between the two reviewers with respect to the component (A-F) ratings?

No Yes

If yes, indicate the reason for the discrepancy

- 1 Oversight
- 2 Differences in interpretation of criteria
- 3 Differences in interpretation of study

Final decision of both reviewers (circle one):

- 1 STRONG
- 2 MODERATE
- 3 WEAK

Appendix 5 Included papers

Abas 2014

Abas MN, Tan PC, Azmi N, Omar SZ. Ondansetron compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2014;**123**:1272–9.

Adamczak 2007

Adamczak J, Kasdaglis J, Rinehart B, Antebi Y, Wolf E, Terrone D. A prospective randomized trial of solumedrol dose pack vs. Phenergan for the treatment of symptomatic nausea and vomiting in pregnancy. *Am J Obstet Gynecol* 2007;**197**:S88.

Alalade 2007

Alalade AO, Khan R, Dawlatly B. Day-case management of hyperemesis gravidarum: feasibility and clinical efficacy. *J Obstet Gynaecol* 2007;**27**:363–4.

Ashkenazi-Hoffnung 2013

Ashkenazi-Hoffnung L, Merlob P, Stahl B, Klinger G. Evaluation of the efficacy and safety of bi-daily combination therapy with pyridoxine and doxylamine for nausea and vomiting of pregnancy. *Isr Med Assoc J* 2013;**15**:23–6.

Babaei 2014

Babaei AH, Foghaha MH. A randomized comparison of vitamin B6 and dimenhydrinate in the treatment of nausea and vomiting in early pregnancy. *Iran J Nurs Midwifery Res* 2014;**19**:199–202.

Basirat 2009

Basirat Z, Moghadamnia AA, Kashifard M, Sarifi-Razavi A. The effect of ginger biscuit on nausea and vomiting in early pregnancy. *Acta Medica Iranica* 2009;**47**:51–6.

Bayreuther 1994

Bayreuther J, Lewith GT, Pickering R. A double-blind cross-over study to evaluate the effectiveness of acupressure at pericardium 6 (P6) in the treatment of early morning sickness (EMS). *Complement Ther Med* 1994;**2**:70–6.

Belluomini 1994

Belluomini J, Litt RC, Lee KA, Katz M. Acupressure for nausea and vomiting of pregnancy: a randomized, blinded study. *Obstet Gynecol* 1994;**84**:245–8.

Biswas 2011

Biswas SC, Dey R, Kamliya GS, Bal R, Hazra A, Tripathi SK. A single-masked, randomized, controlled trial of ginger extract in the treatment of nausea and vomiting of pregnancy. *JIMSA* 2011;**24**:167–9.

Bondok 2006

Bondok RS, El Sharnouby NM, Eid HE, Abd Elmaksoud AM. Pulsed steroid therapy is an effective treatment for intractable hyperemesis gravidarum. *Crit Care Med* 2006;**34**:2781–3.

Capp 2014

Capp S, Oliveira L, Carstairs S, You W. Ondansetron versus doxylamine/pyridoxine for treatment of nausea and vomiting in pregnancy: a prospective randomized double-blind trial. *Am J Obstet Gynecol* 2014;**210**:S39.

Can Gurkan 2008

Can Gurkan O, Arslan H. Effect of acupressure on nausea and vomiting during pregnancy. *Complement Ther Clin Pract* 2008;**14**:46–52.

Carlsson 2000

Carlsson CP, Axemo P, Bodin A, Carstensen H, Ehrenroth B, Madegard-Lind I, *et al.* Manual acupuncture reduces hyperemesis gravidarum: a placebo-controlled, randomized, single-blind, crossover study. *J Pain Symptom Manage* 2000;**20**:273–9.

Chittumma 2007

Chittumma P, Kaewkiattikun K, Wiriyasiriwach B. Comparison of the effectiveness of ginger and vitamin B6 for treatment of nausea and vomiting in early pregnancy: a randomized double-blind controlled trial. *J Med Assoc Thai* 2007;**90**:15–20.

Diggory 1962

Diggory PL, Tomkinson JS. Nausea and vomiting in pregnancy. A trial of meclozine dihydrochloride with and without pyridoxine. *Lancet* 1962;**2**:370–2.

Ditto 1999

Ditto A, Morgante G, la Marca A, De Leo V. Evaluation of treatment of hyperemesis gravidarum using parenteral fluid with or without diazepam. A randomized study. *Gynecol Obstet Invest* 1999;**48**:232–6.

Eftekhari 2013

Eftekhari N, Mehralhasani Y. A comparison of ondansetron and promethasin in treating hyperemesis gravidarum. *JKUMS* 2013;**20**:354–65.

Einarson 2004

Einarson A, Maltepe C, Navioz Y, Kennedy D, Tan MP, Koren G. The safety of ondansetron for nausea and vomiting of pregnancy: a prospective comparative study. *BJOG* 2004;**111**:940–3.

Ensiyeh 2009

Ensiyeh J, Sakineh MA. Comparing ginger and vitamin B6 for the treatment of nausea and vomiting in pregnancy: a randomised controlled trial. *Midwifery* 2009;**25**:649–53.

Erez 1971

Erez S, Schifrin BS, Dirim O. Double-blind evaluation of hydroxyzine as an antiemetic in pregnancy. *J Reprod Med* 1971;**7**:35–7.

Evans 1993

Evans AT, Samuels SN, Marshall C, Bertolucci LE. Suppression of pregnancy-induced nausea and vomiting with sensory afferent stimulation. *J Reprod Med* 1993;**38**:603–6.

Ferreira 2003

Ferreira E, Bussieres JF, Turcotte V, Duperron L, Ouellet G. Case-control study comparing droperidol plus diphenhydramine with conventional treatment in hyperemesis gravidarum. *JPT* 2003;**19**:349–54.

Fischer-Rasmussen 1991

Fischer-Rasmussen W, Kjaer SK, Dahl C, Asping U. Ginger treatment of hyperemesis gravidarum. *Eur J Obstet Gynecol Reprod Biol* 1991;**38**:19–24.

Ghahiri 2011

Ghahiri AA, Abdi F, Mastoo R, Ghasemi M. The effect of ondansetron and metoclopramide in nausea and vomiting of pregnancy. *JIMS* 2011;**29**:259–65.

Ghani 2013

Ghani RMA, Ibrahim ATA. The effect of aromatherapy inhalation on nausea and vomiting in early pregnancy: a pilot randomized controlled trial. *J Nat Sci Res* 2013;**3**:10–22.

Guttuso 2010

Guttuso T Jr, Robinson LK, Amankwah KS. Gabapentin use in hyperemesis gravidarum: a pilot study. *Early Hum Dev* 2010;**86**:65–6.

Haji Seid Javadi 2013

Haji Seid Javadi E, Salehi F, Mashrabi O. Comparing the effectiveness of vitamin B6 and ginger in treatment of pregnancy-induced nausea and vomiting. *Obstet Gynecol Int* 2013;927834.

Heazell 2006

Heazell A, Thorneycroft J, Walton V, Etherington I. Acupressure for the inpatient treatment of nausea and vomiting in early pregnancy: a randomized control trial. *Am J Obstet Gynecol* 2006;**194**:815–20.

Hsu 1996

Hsu JJ, Clark-Glena R, Nelson DK, Kim CH. Nasogastric enteral feeding in the management of hyperemesis gravidarum. *Obstet Gynecol* 1996;**88**:343–6.

Hsu 2003

Hsu E, Pei V, Shofer FS, Abbuhl SB. A prospective randomized controlled trial of acupressure vs sham for pregnancy-related nausea and vomiting in the emergency department. *Acad Emerg Med* 2003;**10**:437.

Jamigorn 2007

Jamigorn M, Phupong V. Acupressure and vitamin B6 to relieve nausea and vomiting in pregnancy: a randomized study. *Arch Gynecol Obstet* 2007;**276**:245–9.

Kashifard 2013

Kashifard M, Basirat Z, Kashifard M, Golsorkhtabar-Amiri M, Moghaddamnia A. Ondansetrone or metoclopramide? Which is more effective in severe nausea and vomiting of pregnancy? A randomized trial double-blind study. *Clin Exp Obstet Gynecol* 2013;**40**:127–30.

Keating 2002

Keating A, Chez RA. Ginger syrup as an antiemetic in early pregnancy. *Altern Ther Health Med* 2002;**8**:89–91.

Knight 2001

Knight B, Mudge C, Openshaw S, White A, Hart A. Effect of acupuncture on nausea of pregnancy: a randomized, controlled trial. *Obstet Gynecol* 2001;**97**:184–8.

Koren 2010

Koren G, Clark S, Hankins GD, Caritis SN, Miodovnik M, Umans JG, *et al.* Effectiveness of delayed-release doxylamine and pyridoxine for nausea and vomiting of pregnancy: a randomized placebo controlled trial. *Am J Obstet Gynecol* 2010;**203**:571.e1–7.

Koren 2013

Koren G, Maltepe C. Preemptive Diclectin therapy for the management of nausea and vomiting of pregnancy and hyperemesis gravidarum. *Am J Obstet Gynecol* 2013;**208**:S20.

Maina 2014

Maina A, Arrotta M, Cicogna L, Donvito V, Mischinelli M, Todros T, *et al.* Transdermal clonidine in the treatment of severe hyperemesis. A pilot randomised control trial: CLONEMESI. *BJOG* 2014;**121**:1556–62.

Maltepe 2013

Maltepe C, Koren G. Preemptive treatment of nausea and vomiting of pregnancy: results of a randomized controlled trial. *Obstet Gynecol Int* 2013;809787.

Mao 2009

Mao ZN, Liang CE. [Observation on therapeutic effect of acupuncture on hyperemesis gravidarum.] *Zhongguo Zhenjiu* 2009;**29**:973–6.

Markose 2004

Markose MT, Ramanathan K, Vijayakumar J. Reduction of nausea, vomiting, and dry retches with P6 acupressure during pregnancy. *Int J Gynaecol Obstet* 2004;**85**:168–9.

McParlin 2008

Primary reference

McParlin C, Carrick-Sen D, Steen IN, Taylor P, Robson SC. Hyperemesis in pregnancy study: a randomised controlled trial of midwife-led 'outpatient' care. *Arch Dis Child Fetal Neonatal Ed* 2008;**93**:Fa9.

Secondary reference

McParlin C, Carrick-Sen D, Steen IN, Taylor P, Robson SC. Hyperemesis in pregnancy study: a randomised controlled trial of midwife-led 'outpatient' care. Unpublished.

Mohammadbeigi 2011

Mohammadbeigi R, Shahgeibi S, Soufizadeh N, Rezaie M, Farhadifar F. Comparing the effects of ginger and metoclopramide on the treatment of pregnancy nausea. *Pak J Biol Sci* 2011;**14**:817–20.

Monias 1957

Monias M. Evaluation of cyclizine with pyridoxine in vomiting of pregnancy. *Mil Med* 1957;**121**:403–4.

Moran 2002

Moran P, Taylor R. Management of hyperemesis gravidarum: the importance of weight loss as a criterion for steroid therapy. *QJM* 2002;**95**:153–8.

Naeimi Rad 2012

Naeimi Rad M, Lamyian M, Heshmat R, Jaafarabadi MA, Yazdani S. A randomized clinical trial of the efficacy of KID21 point (youmen) acupressure on nausea and vomiting of pregnancy. *Iran* 2012;**14**:697–701.

Narenji 2012

Narenji F, Delavar M, Rafiei M. Comparison the effects of the ginger fresh root and vitamin B6 on the nausea and vomiting in pregnancy. *IJOGI* 2012;**15**:39–43.

Nelson-Piercy 2001

Nelson-Piercy C, Fayers P, Swiet M. Randomised, double-blind, placebo-controlled trial of corticosteroids for the treatment of hyperemesis gravidarum. *BJOG* 2001;**108**:9–15.

Neri 2005

Neri I, Allais G, Schiapparelli P, Blasi I, Benedetto C, Facchinetti F. Acupuncture versus pharmacological approach to reduce hyperemesis gravidarum discomfort. *Minerva Ginecol* 2005;**57**:471–5.

Oliveira 2013

Oliveira LG, Capp S, You WB, Carstairs SD. Ondansetron versus doxylamine/pyridoxine for treatment of nausea and vomiting in first trimester pregnancy: a prospective randomized double-blind controlled study. *Acad Emerg Med* 2013;**1**:S101.

Ozgoli 2009

Ozgoli G, Goli M, Simbar M. Effects of ginger capsules on pregnancy, nausea, and vomiting. *J Altern Complement Med* 2009;**15**:243–6.

Pasha 2012

Pasha H, Behmanesh F, Mohsenzadeh F, Hajahmadi M, Moghadamnia AA. Study of the effect of mint oil on nausea and vomiting during pregnancy. *Iran Red Crescent Med J* 2012;**14**:727–30.

Pongroj paw 2007

Pongroj paw D, Somprasit C, Chanthasenanont A. A randomized comparison of ginger and dimenhydrinate in the treatment of nausea and vomiting in pregnancy. *J Med Assoc Thai* 2007;**90**:1703–9.

Rosen 2003

Rosen T, de Veciana M, Miller HS, Stewart L, Rebarber A, Slotnick RN. A randomized controlled trial of nerve stimulation for relief of nausea and vomiting in pregnancy. *Obstet Gynecol* 2003;**102**:129–35.

Saberi 2013

Saberi F, Sadat Z, Abedzadeh-Kalahroudi M, Taebi M. Acupressure and ginger to relieve nausea and vomiting in pregnancy: a randomized study. *Iran Red Crescent Med J* 2013;**15**:854–61.

Safari 1998

Safari HR, Fassett MJ, Souter IC, Alsulyman OM, Goodwin TM. The efficacy of methylprednisolone in the treatment of hyperemesis gravidarum: a randomized, double-blind, controlled study. *Am J Obstet Gynecol* 1998;**179**:921–4.

Saha 2009

Saha S, Loranger D, Pricolo V, Degli-Esposti S. Feeding jejunostomy for the treatment of severe hyperemesis gravidarum: a case series. *J Parenter Enteral Nutr* 2009;**33**:529–34.

Sahakian 1991

Sahakian V, Rouse D, Sipes S, Rose N, Niebyl J. Vitamin B6 is effective therapy for nausea and vomiting of pregnancy: a randomized, double-blind placebo-controlled study. *Obstet Gynecol* 1991;**78**:33–6.

Smith 2002

Smith C, Crowther C, Beilby J. Acupuncture to treat nausea and vomiting in early pregnancy: a randomized controlled trial. *Birth* 2002;**29**:1–9.

Smith 2004

Smith C, Crowther C, Willson K, Hotham N, McMillian V. A randomized controlled trial of ginger to treat nausea and vomiting in pregnancy. *Obstet Gynecol* 2004;**103**:639–45.

Sripramote 2003

Sripramote M, Lekhyananda N. A randomized comparison of ginger and vitamin B6 in the treatment of nausea and vomiting in pregnancy. *J Med Assoc Thai* 2003;**86**:846–53.

Steele 2001

Steele NM, French J, Gatherer-Boyles J, Newman S, Leclaire S. Effect of acupuncture by sea-bands on nausea and vomiting of pregnancy. *J Obstet Gynecol Neonatal Nurs* 2001;**30**:61–70.

Sullivan 1996

Sullivan CA, Johnson CA, Roach H, Martin RW, Stewart DK, Morrison JC. A pilot study of intravenous ondansetron for hyperemesis gravidarum. *Am J Obstet Gynecol* 1996;**174**:1565–8.

Tan 2009

Tan PC, Yow CM, Omar SZ. A placebo-controlled trial of oral pyridoxine in hyperemesis gravidarum. *Gynecol Obstet Invest* 2009;**67**:151–7.

Tan 2010

Tan PC, Khine PP, Vallikkannu N, Omar SZ. Promethazine compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2010;**115**:975–81.

Tan 2013

Tan PC, Norazilah MJ, Omar SZ. Dextrose saline compared with normal saline rehydration of hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2013;**121**:291–8.

Veciana 2001

Veciana M, Stewart L, Miller H, Slotnick R, Rebarber A, Rosen T. Multicenter randomized controlled trial of nerve stimulation therapy for the relief of nausea and vomiting in pregnancy. *Am J Obstet Gynecol* 2001;**185**:S182.

Vutyavanich 1995

Vutyavanich T, Wongtra-ngan S, Ruangsri R. Pyridoxine for nausea and vomiting of pregnancy: a randomized, double-blind, placebo-controlled trial. *Am J Obstet Gynecol* 1995;**173**:881–4.

Vutyavanich 2001

Vutyavanich T, Kraisarin T, Ruangsri R. Ginger for nausea and vomiting in pregnancy: randomized, double-masked, placebo-controlled trial. *Obstet Gynecol* 2001;**97**:577–82.

Wentoft 2001

Werntoft E, Dykes AK. Effect of acupuncture on nausea and vomiting during pregnancy. A randomized, placebo-controlled, pilot study. *J Reprod Med* 2001;**46**:835–9.

Wibowo 2012

Wibowo N, Purwosunu Y, Sekizawa A, Farina A, Tambunan V, Bardosono S. Vitamin B6 supplementation in pregnant women with nausea and vomiting. *Int J Gynaecol Obstet* 2012;**116**:206–10.

Willetts 2003

Willetts KE, Ekangaki A, Eden JA. Effect of a ginger extract on pregnancy-induced nausea: a randomised controlled trial. *Aust N Z J Obstet Gynaecol* 2003;**43**:139–44.

Yost 2003

Yost NP, McIntire DD, Wians FH Jr, Ramin SM, Balko JA, Leveno KJ. A randomized, placebo-controlled trial of corticosteroids for hyperemesis due to pregnancy. *Obstet Gynecol* 2003;**102**:1250–4.

Zhang 2005

Zhang HH. [Observation on therapeutic effect of acupuncture and moxibustion on hyperemesis gravidarum.] *Zhongguo Zhenjiu* 2005;**25**:469–70.

Ziaei 2004

Ziaei S, Hosseiny FS, Faghihzadeh S. The efficacy low dose of prednisolone in the treatment of hyperemesis gravidarum. *Acta Obstet Gynecol Scand* 2004;**83**:272–5.

Appendix 6 Excluded papers and reasons for exclusion

Required study design not met ($n = 70$)

Abas MN, Tan PC, Azmi N, Omar SZ. Ondansetron compared with metoclopramide for hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2013;**123**:1272–9.

Agren A, Berg M. Tactile massage and severe nausea and vomiting during pregnancy – women’s experiences. *Scand J Caring Sci* 2006;**20**:169–76.

Ajufo I. Four-year retrospective review of day admission service provision for hyperemesis gravidarum (HG) at St Thomas’ Hospital London [UK]. *BJOG* 2013;**120**:534–5.

Bailit JL. Hyperemesis gravidarum: epidemiologic findings from a large cohort. *Am J Obstet Gynecol* 2005;**193**:811–14.

Balestri F. [Treatment of hyperemesis gravidarum with combined inositol and pyridoxine.] *Gazz Med Ital* 1955;**114**:225–7.

Bond S. Large study finds metoclopramide can be safely used for nausea and vomiting in early pregnancy. *J Midwifery Womens Health* 2009;**54**:510–11.

Boone SA, Shields KM. Treating pregnancy-related nausea and vomiting with ginger. *Ann Pharmacother* 2005;**39**:1710–13.

Boskovic R, Einarson A, Maltepe C, Wolpin J, Koren G. Diclectin therapy for nausea and vomiting of pregnancy: effects of optimal dosing. *J Obstet Gynaecol Can* 2003;**25**:830–3.

Bozzo P, Koren G, Nava-Ocampo AA, Einarson A. The incidence of nausea and vomiting of pregnancy (NVP): a comparison between depressed women treated with antidepressants and non-depressed women. *Clin Invest Med* 2006;**29**:347–50.

Brill JR. Acupressure for nausea and vomiting of pregnancy: a randomized, blinded study (4). *Obstet Gynecol* 1995;**85**:159–60.

Brodbaek HB, Damkier P. [The treatment of hyperemesis gravidarum with chlorobutanol-caffeine rectal suppositories in Denmark: practice and evidence.] *Ugeskr Laeger* 2007;**169**:2122–3.

Bufano G, Cervellin G, Calderini MC, Coscelli C. [Enteral nutrition in hyperemesis gravidarum.] *Rivista Italiana di Nutrizione Parenterale ed Enterale* 1992;**10**:260–5.

Buttino L Jr, Coleman SK, Bergauer NK, Gambon C, Stanziano GJ. Home subcutaneous metoclopramide therapy for hyperemesis gravidarum. *J Perinatol* 2000;**20**:359–62.

Charlin V, Borghesi L, Hasbun J, Von Mulenbrock R, Moreno MI. Parenteral nutrition in hyperemesis gravidarum. *Nutrition* 1993;**9**:29–32; discussion 68.

Crawford-Faucher A. Which drug is more effective for treating hyperemesis gravidarum? *Am Fam Physician* 2011;**83**:842.

Fejzo MS, Magtira A, Schoenberg FP, MacGibbon K, Mullin P, Romero R, *et al.* Antihistamines and other prognostic factors for adverse outcome in hyperemesis gravidarum. *Eur J Obstet Gynecol Reprod Biol* 2013;**170**:71–6.

Fuchs K, Paldi E, Abramovici H, Peretz BA. Treatment of hyperemesis gravidarum by hypnosis. *Int J Clin Exp Hyp* 1980;**28**:313–23.

Gordon GH. Dextrose saline compared with normal saline rehydration of hyperemesis gravidarum: a randomized controlled trial. *Obstet Gynecol* 2013;**121**:1360.

Heitmann K, Holst L, Maltepe C, Lupattelli A, Nordeng H. Treatment of nausea during pregnancy – results from a multinational, cross-sectional, internet-based study. *Res Soc Adm Pharm* 2014;**10**:e57–8.

Gill SK, Maltepe C, Mastali K, Koren G. The effect of acid-reducing pharmacotherapy on the severity of nausea and vomiting of pregnancy. *Obstet Gynecol Int* 2009;585269.

Golaszewski T, Frigo P, Mark HE, Rattay F, Schaller A. [Treatment of hyperemesis gravidarum by electrostimulation of the vestibular apparatus.] *Z Geburtshilfe Neonatol* 1995;**199**:107–10.

Goodwin TM, Poursharif B, Korst LM, MacGibbon KW, Romero R, Fejzo MS. Secular trends in the treatment of hyperemesis gravidarum. *Am J Perinatol* 2008;**25**:141–7.

Gulley RM, Pleog NV, Gulley JM. Treatment of hyperemesis gravidarum with nasogastric feeding. *Nutr Clin Pract* 1993;**8**:33–5.

Hordern CE, Medina Lucena H, Stanley KP, Sule MM. HARP – Hyperemesis Ambulatory Rehydration Project: development, implementation and improvement to minimise hospital stay. *BJOG* 2013;**120**:432–3.

Hyde E. Acupressure therapy for morning sickness. A controlled clinical trial. *J Nurse Midwifery* 1989;**34**:171–8.

Jednak MA, Shadigian EM, Kim MS, Woods ML, Hooper FG, Owyang C, *et al.* Protein meals reduce nausea and gastric slow wave dysrhythmic activity in first trimester pregnancy. *Am J Physiol* 1999;**277**:G855–61.

Jewell D, Young G, Hall PF. Review: antiemetic drugs reduce nausea in early pregnancy. *Evid-Based Med* 2002;**7**:155.

Jungmayr P. [Pregnancy hyperemesis: a cohort study confirms safety of metoclopramide.] *Deutsche Apotheker Zeitung* 2009;**149**:42–3.

Kawakami SI, Furuiki Y, Kubota T, Souda Y, Souda T, Kyono K, *et al.* Effect of Chinese Herbal Medicine Suppositories for Hyperemesis Gravidarum by Using an Index for Nausea and Vomiting of Pregnancy. In Koren G, Bishai R, editors. *Nausea and Vomiting of Pregnancy: State of the Art 2000*. Toronto: Hospital Sick Children, Motherisk Program; 2000. pp. 122–7.

Khan TN, Karpate S, Shehmar M. Hyperemesis day centre audit. *BJOG* 2013;**120**:527–8.

King TL, Murphy PA. Evidence-based approaches to managing nausea and vomiting in early pregnancy. *J Midwifery Womens Health* 2009;**54**:430–44.

Koren G. Appraisal of drug therapy for nausea and vomiting of pregnancy: I. The placebo effect – methodological and practical considerations. *Can J Clin Pharmacol* 2000;**7**:135–7.

- Lacasse A, Rey E, Ferreira E, Morin C, Bérard A. Validity of a modified Pregnancy-Unique Quantification of Emesis and Nausea (PUQE) scoring index to assess severity of nausea and vomiting of pregnancy. *Am J Obstet Gynecol* 2008;**198**:71.e1–7.
- Lacasse A, Berard A. Validation of the nausea and vomiting of pregnancy specific health related quality of life questionnaire. *Health Qual Life Outcomes* 2008;**6**:32.
- Long MA, Simone SS, Tucher JJ. Outpatient treatment of hyperemesis gravidarum with stimulus control and imagery procedures. *J Behav Ther Exp Psychiatry* 1986;**17**:105–9.
- Low KG. Nausea and vomiting in pregnancy: a review of the research. *J Gender Culture Health* 1996;**1**:151–72.
- Magee LA, Chandra K, Mazzotta P, Stewart D, Koren G, Guyatt GH. Development of a health-related quality of life instrument for nausea and vomiting of pregnancy. *Am J Obstet Gynecol* 2002;**186**:S232–8.
- Maina A, Todros T. A novel approach to hyperemesis gravidarum: evaluation by a visual analogue scale score and treatment with transdermal clonidine. *Obstet Med* 2011;**4**:156–9.
- Matok I, Umans J, Feghali MN, Clark S, Caritis S, Miodovnik M, et al. Characteristics of women with nausea and vomiting of pregnancy who chose to continue compassionate use of placebo after a randomised trial. *J Obstet Gynaecol* 2013;**33**:557–60.
- Maxwell CI, Hilden K, Delegge MH, Kinikini MM, Olafsson S, Fang JC. Feasibility of percutaneous direct jejunal feeding tubes in pregnant patients. *Gastrointest Endosc* 2010;**71**:AB268–9.
- McCauley L, Coleman S, Jacques D, Palmer B, Stanziano G. Safety and efficacy of ondansetron therapy for nausea and vomiting of pregnancy. *Obstet Gynecol* 2002;**99**:24S.
- Milkovich L, van den Berg BJ. An evaluation of the teratogenicity of certain anti-nauseant drugs. *Am J Obstet Gynecol* 1976;**125**:244–8.
- Munch S. A qualitative analysis of physician humanism: women's experiences with hyperemesis gravidarum. *J Perinatol* 2000;**20**:540–7.
- Munch S, Schmitz MF. The Hyperemesis Beliefs Scale (HBS): a new instrument for assessing beliefs about severe nausea and vomiting in pregnancy. *J Psychosom Obstet Gynaecol* 2007;**28**:219–29.
- Nelson-Piercy C, De Swiet M. Corticosteroids for the treatment of hyperemesis gravidarum. *Br J Obstet Gynaecol* 1994;**101**:1013.
- Norheim AJ, Pedersen EJ, Fonnebo V, Berge L. Acupressure treatment of morning sickness in pregnancy. A randomised, double-blind, placebo-controlled study. *Scand J Prim Health Care* 2001;**19**:43–7.
- Nulman I, Rovet J, Barrera M, Knittel-Keren D, Feldman BM, Koren G. Long-term neurodevelopment of children exposed to maternal nausea and vomiting of pregnancy and dicyclanil. *J Pediatr* 2009;**155**:45–50.e1–2.
- Onghena G, De ROM. [Clinical study of prochlorperazine in hyperemesis gravidarum.] *Brux Med* 1960;**40**:1551–4.
- Palatty PL, Haniadka R, Valder B, Arora R, Baliga MS. Ginger in the prevention of nausea and vomiting: a review. *Clin Rev Food Sci Nutr* 2013;**53**:659–69.

Pasquinucci C. [Association of vitamins B 1 and B 6 in high doses in therapy of hyperemesis gravidarum.] *Ann Ostet Ginecol Med Perinat* 1965;**87**:65–70.

Pasternak B, Svanstrom H, Molgaard-Nielsen D, Melbye M, Hviid A. Metoclopramide in pregnancy and risk of major congenital malformations and fetal death. *JAMA* 2013;**310**:1601–11.

Pasternak B, Svanstrom H, Hviid A. Ondansetron in pregnancy and risk of adverse fetal outcomes. *N Engl J Med* 2013;**368**:814–23.

Portnoi G, Chng LA, Karimi-Tabesh L, Koren G, Tan MP, Einarson A. Prospective comparative study of the safety and effectiveness of ginger for the treatment of nausea and vomiting in pregnancy. *Am J Obstet Gynecol* 2003;**189**:1374–7.

Puangrichareern A, Mahasukhon S. Effectiveness of auricular acupressure in the treatment of nausea and vomiting in early pregnancy. *J Med Assoc Thai* 2008;**91**:1633–8.

Seidner HM. Nausea and vomiting of pregnancy; preliminary trial of bonamine in sixteen patients. *Ill Med J* 1956;**109**:20–1.

Shrim A, Boskovic R, Maltepe C, Navios Y, Garcia-Bournissen F, Koren G. Pregnancy outcome following use of large doses of vitamin B6 in the first trimester. *J Obstet Gynaecol* 2006;**26**:749–51.

Slotnick RN. Safe, successful nausea suppression in early pregnancy with P-6 acustimulation. *J Reprod Med* 2001;**46**:811–14.

Smith C, Crowther C. The placebo response and effect of time in a trial of acupuncture to treat nausea and vomiting in early pregnancy. *Complement Ther Med* 2002;**10**:210–16.

Smith C, Crowther C, Beilby J. Pregnancy outcome following women's participation in a randomised controlled trial of acupuncture to treat nausea and vomiting in early pregnancy. *Complement Ther Med* 2002;**10**:78–83.

Sorensen HT, Nielsen GL, Christensen K, Tage-Jensen U, Ekbom A, Baron J. Birth outcome following maternal use of metoclopramide. The Euromap study group. *Br J Clin Pharmacol* 2000;**49**:264–8.

Stainton MC, Neff EJ. The efficacy of SeaBands for the control of nausea and vomiting in pregnancy. *Health Care Women Int* 1994;**15**:563–75.

Subramaniam R, Soh EB, Dhillon HK, Abidin HZ. Total parenteral nutrition (TPN) and steroid usage in the management of hyperemesis gravidarum. *Aust N Z J Obstet Gynaecol* 1998;**38**:339–41.

Taylor R. Successful management of hyperemesis gravidarum using steroid therapy. *QJM* 1996;**89**:103–7.

Tompsett H, Ajala T, Dixon G, Kelly T. The implementation of an ambulatory care bundle for the treatment of hyperemesis gravidarum. *BJOG* 2013;**120**:477.

Trovik J, Haram K, Berstad A, Flaatten H. [Nasoenteral tube feeding in hyperemesis gravidarum. An alternative to parenteral nutrition.] *Tidsskr Nor Laegeforen* 1996;**116**:2442–4.

Vaisman N, Kaidar R, Levin I, Lessing JB. Nasojejunal feeding in hyperemesis gravidarum – a preliminary study. *Clin Nutr* 2004;**23**:53–7.

Wang Y, Lin Y, Wu X. The effect of total parenteral nutrition on hyperemesis gravidarum and pregnancy ending. *Clin Med China* 2001;**8**:63–4.

Weinstein BB, Wohl Z, Mitchell GJ, Sustental GF. Oral administration of pyridoxime hydrochloride in the treatment of nausea and vomiting of pregnancy. *Amur J Obstet Gynecol* 1944;**47**:389–94.

Westfall RE, Janssen PA, Lucas P, Capler R. Survey of medicinal cannabis use among childbearing women: patterns of its use in pregnancy and retroactive self-assessment of its efficacy against 'morning sickness'. *Complement Ther Clin Pract* 2006;**12**:27–33.

Whittaker R. *Randomised, Double-Blind, Placebo-Controlled Trial of Corticosteroids for the Treatment of Hyperemesis Gravidarum*. 2003. URL: <http://onlinelibrary.wiley.com/doi/10.1002/14651914.cochrane.clcentral/articles/519/CN-00473519/frame.html>

Required participant inclusion criteria not met (n = 34)

Atanackovic G, Navioz Y, Moretti ME, Koren G. The safety of higher than standard dose of doxylamine-pyridoxine (Diclectin) for nausea and vomiting of pregnancy. *J Clin Pharmacol* 2001;**41**:842–5.

Bethea RC. Evaluation of a combination of meclizine and pyridoxine in the control of nausea and vomiting in pregnancy. *Int Rec Med* 1960;**173**:283–7.

Bisley BL, Kay CR. Anti-emetic effects of a phenothiazine compared with an antihistamine. *Practitioner* 1964;**193**:358–60.

Boneva RS, Moore CA, Botto L, Wong LY, Erickson JD. Nausea of Pregnancy, Antinausea Preparations and Congenital Heart Defects: A Population-Based Case–Control Study. In Koren G, Bishai R, editors. *Nausea and Vomiting of Pregnancy: State of the Art 2000*. Toronto: Hospital Sick Children, Motherisk Program; 2000. pp. 60–72.

Christer C. Manual acupuncture at pc6 reduces hyperemesis gravidarum—a placebo-controlled randomised crossover study. *Akupunktur* 1998;**26**:261–2.

Coleman L, O'Sullivan M, Dilloway L, Sinha A, Epee M. An innovative ambulatory care service for women suffering with hyperemesis gravidarum. *BJOG* 2014;**121**:19.

Cottin J, Beghin D, Jonville-Bera AP, Boyer M, Zenut M, Damase-Michel C, *et al*. First trimester exposure to domperidone: a comparative prospective study. *Fundam Clin Pharm* 2013;**27**:44.

Czeizel AE, Dudas I, Fritz G, Tecsoi A, Hanck A, Kunovits G. The effect of periconceptional multivitamin-mineral supplementation on vertigo, nausea and vomiting in the first trimester of pregnancy. *Arch Gynecol Obstet* 1992;**251**:181–5.

Dorsey CW. The use of pyridoxine and suprarenal cortex combined in the treatment of nausea and vomiting of pregnancy. *Am J Obstet Gynecol* 1949;**58**:1073–8.

Duggar CR, Carlan SJ. The efficacy of methylprednisolone in the treatment of hyperemesis gravidarum: a randomized double-blind controlled study. *Obstet Gynecol* 2001;**97**:45.

Dundee JW, Sourial FB, Ghaly RG, Bell PF. P6 acupressure reduces morning sickness. *J R Soc Med* 1988;**81**:456–7.

- Durham MP. Clinical trial of buclizine hydrochloride for vomiting of pregnancy. *Br Med J* 1956;**2**:1276–7.
- Eriksson G, Jullig D, Tysk U. [Postadoxin-a new remedy for nausea and emesis gravidarum. A clinical test with the double-blind method.] *Sven Lakartidn* 1961;**58**:1722–5.
- Fletcher SJ, Waterman H, Nelson L, Carter LA, Chuang LH, Roberts C, *et al.* The effectiveness and cost-effectiveness of a holistic assessment and individualised package of care of women with hyperemesis gravidarum: randomised controlled trial. *BJOG* 2013;**120**:552–3.
- Ge Q. The therapeutic effect of intravenous nutritional solution combined with psychological interventions on 100 cases of hyperemesis gravidarum. *China J Health Psychol* 2012;**9**:1322–4.
- Gill SK, Maltepe C, Koren G. The effectiveness of discontinuing iron-containing prenatal multivitamins on reducing the severity of nausea and vomiting of pregnancy. *J Obstet Gynaecol* 2009;**29**:13–16.
- Hart BF, McConnell WT, Pickett AN. Vitamin and endocrine therapy in nausea and vomiting of pregnancy. *Am J Obstet Gynecol* 1944;**48**:251–3.
- He XL, Zhong G, He Y. [Clinical observation on treatment of hyperemesis gravidarum by integrative Chinese and Western medicine and its influence on serum motilin.] *Zhongguo Zhong Xi Yi Jie He Za Zhi* 2009;**29**:872–4.
- Henker FO III. Psychotherapy as adjunct in treatment of vomiting during pregnancy. *South Med J* 1976;**69**:1585–7.
- Klauser CK, Fox NS, Istwan N, Rhea D, Rebarber A, Desch C, *et al.* Treatment of severe nausea and vomiting of pregnancy with subcutaneous medications. *Am J Perinatol* 2011;**28**:715–21.
- Koren G, Maltepe C. Pre-emptive therapy for severe nausea and vomiting of pregnancy and hyperemesis gravidarum. *J Obstet Gynaecol* 2004;**24**:530–3.
- Lacasse A, Lagoutte A, Ferreira E, Berard A. Metoclopramide and diphenhydramine in the treatment of hyperemesis gravidarum: effectiveness and predictors of rehospitalisation. *Eur J Obstet Gynecol Reprod Biol* 2009;**143**:43–9.
- Lacasse A, Lagoutte A, Ferreira E, Berard A. Metoclopramide and diphenhydramine in the treatment of hyperemesis gravidarum: effectiveness, safety, and predictors of rehospitalisation. *Pharmacoepidem Dr S* 2007;**16**:S146–S7.
- Liu S-J. Clinical study on TCM and acupuncture combined treating hyperemesis gravidarum. *LNUTCM* 2007;**6**:145–6.
- Modares M, Besharat S, Kian FR, Besharat S, Mahmoudi M, Sourmaghi HS. Effect of ginger and chamomile capsules on nausea and vomiting in pregnancy. *J Gorgan Uni Med Sci* 2012;**14**:e46–e50.
- Mulla NP. Evaluation of trimethobenzamide with pyridoxine in nausea and vomiting of pregnancy. *Clin Med* 1963;**70**:2052–4.
- Newlinds JS. Nausea and vomiting in pregnancy: a trial of thiethylperazine. *Med J Aust* 1964;**1**:234–6.
- O'Brien B, Relyea MJ, Taerum T. Efficacy of P6 acupressure in the treatment of nausea and vomiting during pregnancy. *Am J Obstet Gynecol* 1996;**174**:708–15.

Price JJ, Barry MC. A double blind study of fluphenazine with pyridoxine. *Pa Med J* 1964;**67**:37–40.

Shin HS, Song YA, Seo S. Effect of Nei-Guan point (P6) acupuncture on ketonuria levels, nausea and vomiting in women with hyperemesis gravidarum. *J Adv Nurs* 2007;**59**:510–19.

Shin HS, Song YA. [The effect of P6 acupuncture for symptom control in pregnant women having hyperemesis gravidarum.] *Taehan Kanho Hakhoe Chi* 2005;**35**:593–601.

Sullivan CL. Treatment of nausea and vomiting of pregnancy with chlorpromazine; a report of 100 cases. *Postgrad Med* 1957;**22**:429–32.

Yavari Kia P, Safajou F, Shahnazi M, Nazemiyeh H. The effect of lemon inhalation aromatherapy on nausea and vomiting of pregnancy: a double-blinded, randomized, controlled clinical trial. *Iran* 2014;**16**:e14360.

Zhu Z-N, Fa Y-H, Wang C-R. Injection plus acupuncture clinical observation hyperemesis gravidarum 44 cases. *J Pract Tradit Chinese Intern Med* 2011;**6**:129–30.

Required outcomes not reported (n = 34)

Albertazzi E, Bellodi P, Guerzoni G. Phosphate carbohydrate solutions in the treatment of vomiting in pregnancy. *Curr Ther Res Clin E* 1989;**46**:1059–67.

Anderka M, Mitchell AA, Louik C, Werler MM, Hernandez-Diaz S, Rasmussen SA, *et al.* Medications used to treat nausea and vomiting of pregnancy and the risk of selected birth defects. *Birth Defects Res Part A Clin Mol Teratol* 2012;**94**:22–30.

Andersen JT, Jimenez-Solem E, Andersen NL, Poulsen HE. Ondansetron use in early pregnancy and the risk of congenital malformations—a register based nationwide cohort study. *Pharmacoepidem Dr S* 2013;**22**:13–14.

Aselton P, Jick H, Chentow SJ, Perera DR, Hunter JR, Rothman KJ. Pyloric stenosis and maternal Bendectin exposure. *Am J Epidemiol* 1984;**120**:251–6.

Asker C, Norstedt Wikner B, Kallen B. Use of antiemetic drugs during pregnancy in Sweden. *Eur J Clin Pharmacol* 2005;**61**:899–906.

Babae PAG. Determining the effect of pyridoxine on nausea and vomiting in pregnancy. *J Matern Fetal Neonatal Med* 2010;**23**:129.

Balsamo SB, Sampaio LF, Nobre Nde S, Rozas A, Barros HC, Neme B. [Clinical trial of thiethylperazine (Torecan).] *Hospital (Rio J)* 1966;**70**:451–7.

Berkovitch M, Mazzota P, Greenberg R, Elbirt D, Addis A, Schuler-Faccini L, *et al.* Metoclopramide for nausea and vomiting of pregnancy: a prospective multicenter international study. *Am J Perinatol* 2002;**19**:311–16.

Bsat FA, Hoffman DE, Seubert DE. Comparison of three outpatient regimens in the management of nausea and vomiting in pregnancy. *J Perinatol* 2003;**23**:531–5.

Costantine MM, Matok I, Chioffi G, Clark S, Miodovnik M, Umans JG, *et al.* Determinants of adherence to delayed-release doxylamine and pyridoxine in patients with nausea and vomiting of pregnancy. *Ther Drug Monit* 2012;**34**:569–73.

- de Aloysio D, Penacchioni P. Morning sickness control in early pregnancy by Neiguan point acupressure. *Obstet Gynecol* 1992;**80**:852–4.
- Farazmand T, Khadem N. Comparison of the effect of methylprednisolone and promethazine in the treatment of hyperemesis gravidarum (2001–2002). *Int J Gynecol Obstet* 2009;**107**:S523.
- Folk JJ, Leslie-Brown HF, Nosovitch JT, Silverman RK, Aubry RH. Hyperemesis gravidarum: outcomes and complications with and without total parenteral nutrition. *J Reprod Med* 2004;**49**:497–502.
- Fuchs K, Paldi E, Abramovici H, Peretz BA. Treatment of hyperemesis gravidarum by hypnosis. *Int J Clin Exp Hyp* 1980;**28**:313–23.
- Gant H, Reinken L, Dapunt O, Scholz K. Vitamin B-6 depletion in hyperemesis gravidarum. *Wiener Klinische Wochenschrift* 1975;**87**:510–13.
- Gawande S, Vaidya M, Tadke R, Kirpekar V, Bhawe S. Progressive muscle relaxation in hyperemesis gravidarum. *SAFOG* 2011;**3**:28–32.
- Habek D, Barbir A, Habek JC, Janculiak D, Bobic-Vukovic M. Success of acupuncture and acupressure of the Pc 6 acupoint in the treatment of hyperemesis gravidarum. *Forsch Komplementarmed Klass Naturheilkd* 2004;**11**:20–3.
- Hasbun J, Charlin V, Von Mulhenbrock R, Munoz H, Yuri C. [Total parenteral nutrition in severe hyperemesis gravidarum.] *Rev Chil Obstet Ginecol* 1994;**59**:378–82.
- Heazell AEP, Langford N, Judge JK, Heazell MA, Downey GP. The use of levomepromazine in hyperemesis gravidarum resistant to drug therapy – a case series. *Reprod Toxicol* 2005;**20**:569–72.
- Holmgren C, Aagaard-Tillery KM, Silver RM, Porter TF, Varner M. Hyperemesis in pregnancy: an evaluation of treatment strategies with maternal and neonatal outcomes. *Am J Obstet Gynecol* 2008;**198**:56.e1–4.
- Kutcher JS, Engle A, Firth J, Lamm SH. Bendectin and birth defects. II: Ecological analyses. *Birth Defects Res Part A Clin Mol Teratol* 2003;**67**:88–97.
- Lacasse A, Rey E, Ferreira E, Morin C, Berard A. Nausea and vomiting of pregnancy: what about quality of life? *BJOG* 2008;**115**:1484–93.
- Levine MG, Esser D. Total parenteral nutrition for the treatment of severe hyperemesis gravidarum: maternal nutritional effects and fetal outcome. *Obstet Gynecol* 1988;**72**:102–7.
- Lombardi DG, Istwan NB, Rhea DJ, O'Brien JM, Barton JR. Measuring outpatient outcomes of emesis and nausea management in pregnant women. *Manag Care* 2004;**13**:48–52.
- Matok I, Gorodischer R, Koren G, Sheiner E, Wiznitzer A, Levy A. The safety of metoclopramide use in the first trimester of pregnancy. *N Engl J Med* 2009;**360**:2528–35.
- Moessner GF. Clinical evaluation of a meprobamate–promazine combination for control of nausea and vomiting in pregnancy; a preliminary report. *West J Surg Obstet Gynecol* 1959;**67**:180–2.
- Paridokht A, Gholamreza B. Determine effect VB6 on nausea and vomiting of pregnancy. *J Matern Fetal Neonatal Med* 2010;**23**:127–8.

Reyhani M, Eskandari M. Evaluation of the achillea millefolium effect in nausea and vomiting of early pregnancy. *Iran J Pharmaceut Res* 2013;**12**:232.

Rotman J. [Clinical trial of plitican in vomiting of pregnancy.] *Semaine des Hopitaux* 1986;**62**:138–40.

Tabatabaie A, Sekhvat L, Mojibian M. A randomized, placebo-controlled trial of corticosteroids for hyperemesis gravidarum. *J Matern Fetal Neonatal Med* 2008;**21**:225.

Tasci Y, Demir B, Dilbaz S, Haberal A. Use of diazepam for hyperemesis gravidarum. *J Matern Fetal Neonatal Med* 2009;**22**:353–6.

Wheatley D. Treatment of pregnancy sickness. *Br J Obstet Gynaecol* 1977;**84**:444–7.

Xu C-Z. The clinical study of 654-II in treating hyperemesis gravidarum. *Hebei Medicine* 2005;**10**:923–4.

Ylikorkala O, Kauppila A, Ollanketo ML. Intramuscular ACTH or placebo in the treatment of hyperemesis gravidarum. *Acta Obstet Gynecol Scand* 1979;**58**:453–5.

Appendix 7 UK Teratology Information Service enquiries and follow-ups relating to hyperemesis gravidarum/nausea and vomiting in pregnancy medication

UK Teratology Information Service follow-up data

Ginger

The UKTIS has followed up a single prospective case of ginger exposure during pregnancy. Ginger exposure occurred at an unknown stage of pregnancy and the outcome was a live-born infant with no reported congenital malformations or neonatal problems.¹⁵²

Vitamin B12

The UKTIS has followed up 21 cases of vitamin B12 (hydroxocobalamin and cyanocobalamin) exposure during pregnancy. There were 20 prospective therapeutic exposures and one retrospective case.¹⁵³

Prospective therapeutic exposure data

The frequency of congenital malformations in live-born infants (0/21; 0%, 95% CI 0% to 19.2%) was not significantly higher than the background rate. However, due to the small numbers of cases and wide CIs due to the consequent uncertainty, the conclusions that can be drawn from these data are limited.

Retrospective therapeutic exposure data

The UKTIS occasionally receives retrospective information on the outcome of pregnancies. Although retrospective reports are biased in that adverse outcomes are more likely to be reported, these data may be useful in identifying patterns of malformations that may be suggestive of a teratogenic syndrome and are therefore analysed periodically.

The UKTIS have retrospective follow-up data for one pregnancy following therapeutic exposure to vitamin B12 at 5 weeks. The infant had microcephaly and dysmorphic facial features.

Vitamin B6 (pyridoxine)

The UKTIS has followed up 23 cases of pyridoxine exposure during pregnancy including 18 prospective therapeutic exposures, three overdoses and two retrospective cases.¹⁵⁴

Prospective therapeutic exposure data

The frequency of congenital malformations in live-born infants (3/15; 20%, 95% CI 5.3% to 48.6%) was significantly higher than the background rate; however, the numbers involved are very small and no pattern of malformation was observed. One of the malformations reported is an inherited disorder (Apert syndrome) and not expected to be linked to the maternal exposure.

Prospective overdose data

The UKTIS have collected prospective follow up data on three pregnancies in which pyridoxine was taken in overdose (amounts ingested ranged from 1250 mg to 3500 mg). The outcomes were two healthy babies and one elective termination.

Retrospective therapeutic exposure data

The UKTIS have retrospective follow-up data for two pregnancies following therapeutic exposure to pyridoxine. In the first case, exposure to an unreported amount of pyridoxine occurred between 11 and

15 weeks' gestation. The outcome was an infant with multiple congenital anomalies including a bent ulna with a malformed hand, polydactyly on the other hand, and one leg with talipes and abnormal digits. In the second case maternal use of 50-mg pyridoxine per day throughout pregnancy resulted in an infant delivered with polydactyly. In addition to the pyridoxine, 20-mg paroxetine was taken each day throughout pregnancy, and 600-mg rifampicin and 300-mg isoniazid was administered per day from 0 to 12 weeks' gestation.

Promethazine

The UKTIS has followed up 67 cases of promethazine exposure during pregnancy. There were 45 prospective therapeutic exposures, 15 overdoses and seven retrospective cases.¹⁵⁵

Prospective therapeutic exposure data

The frequency of congenital malformations in live-born infants (2/41; 4.9%, 95% CI 0.85% to 17.8%) was not significantly higher than the background rate.

Retrospective exposure data

The UKTIS have retrospective follow-up data for seven pregnancies following exposure to promethazine, six therapeutic and one overdose. Malformations were reported in all cases but showed no consistent pattern suggestive of teratogenesis.

Ondansetron

The UKTIS has followed up 48 cases of ondansetron exposure during pregnancy. There were 42 prospective therapeutic exposures and six retrospective cases.¹⁵⁶

Prospective therapeutic exposure data

The frequency of live-born infants with one or more major congenital malformation (1/34; 2.9%, 95% CI 0.15% to 17.1%) was not significantly higher than the background rate (2–3%).

Retrospective therapeutic exposure data

The UKTIS occasionally receives retrospective information on the outcome of pregnancies. These data are analysed periodically to look for patterns of malformations (*Table 42*). No specific pattern has been observed for ondansetron. The UKTIS have retrospective follow-up data for six pregnancies following therapeutic exposure to ondansetron.

TABLE 42 UK Teratology Information Service enquires relating to HG/NVP medications

Drugs	Total number of enquiries		Annual enquiries by 5-year period									
	Total number of enquiries for exposure	Enquiries for which indication was HG	HG enquiries for which pregnancy outcome known ^a	Pre 1990	1990–4	1995–9	2000–4	2005–9	2010–14 ^b			
Calcium carbonate	188	1	0	–	–	–	–	1	–	–	–	
Cetirizine	356	6	3	–	–	1	–	3	–	–	2	
Cyclizine (Valoid)	433	221	58	–	–	7	36	79	–	–	99	
Dimenhydrinate (Dramamine®, Prestige Brands Holdings, Inc.)	4	1	1	–	–	–	–	–	–	–	1	
Diphenhydramine	68	2	1	–	1	1	–	–	–	–	–	
Doxylamine	12	3	1	–	–	–	–	1	–	–	2	
Doxylamine succinate-pyridoxine hydrochloride	2	1	0	–	–	–	–	1	–	–	–	
Hydroxyzine	62	2	0	–	–	–	1	1	–	–	–	
Hydrocortisone	265	19	4	–	–	–	6	12	–	–	1	
Methylprednisolone	58	2	0	–	–	–	1	–	–	–	1	
Metoclopramide	278	108	33	–	2	7	17	43	–	–	39	
Ondansetron	235	140	33	–	–	8	17	53	–	–	62	
Peppermint	76	1	0	–	–	–	–	1	–	–	–	
Prednisolone	559	21	8	–	–	1	5	10	–	–	5	
Prochlorperazine (Stemetil, Sanofi)	426	207	43	–	9	11	35	92	–	–	60	
Promethazine (Avomine, Manx Healthcare Ltd, Phenergan)	405	93	28	1	7	4	16	31	–	–	34	
Pyridoxine (vitamin B6)	100	12	6	1	–	–	–	8	–	–	3	
Ranitidine	377	35	17	–	2	7	8	10	–	–	8	
Thiamine	97	18	7	–	–	2	3	12	–	–	1	
Vitamin B12 (cyanocobalamin)	144	2	2	–	–	–	–	–	–	–	2	
Total enquiries	–	–	219 ^a	2	21	49	145	358	–	–	320 ^a	

^a Data for 2010–14 are incomplete.

^b Incomplete 5-year period as 2014 ongoing, some of these pregnancies will not yet have ended so follow-up rate likely to be slightly higher.

Appendix 8 Secondary outcome data

Study	Pregnancy outcomes	Other outcomes
Abas 2014 ⁵⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference in rates of overall treatment curtailment; treatment curtailment attributable to adverse events; or need for continuation with open-label i.v. antiemetic ($p > 0.05$) No statistically significant difference in length of hospital stay between trial arms [intervention: 1.9 (IQR 1.5–2.4) days vs. comparator: 2.0 (IQR 1.7–2.7) days; $p = 0.1$] Significant differences were found in self-reported drowsiness ($p = 0.011$) and dry mouth ($p = 0.003$) in favour of ondansetron, but other minor side effects (difficulty sleeping, dizziness, diarrhoea, headache, palpitations, and skin rash) were reported in similar proportions across the trial arms ($p > 0.5$)
Adamczak 2007 ⁵⁸	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Alalade 2007 ¹²⁰	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> All patients managed as day cases were discharged within 24 hours None of them were readmitted A total of 51% of patients were treated and discharged between 12 and 24 hours
Ashkenazi-Hoffnung 2013 ³⁶	<ul style="list-style-type: none"> One miscarriage control group, two in treatment group. No difference in mean birthweight. One congenital abnormality in control group (hypospadias). Late preterm birth: B6 combination 5/28 (18%) vs. metoclopramide 0/28 (0%); $p = 0.03$ 	<ul style="list-style-type: none"> Not reported
Babaei 2014 ⁵⁹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Occurrence of drowsiness significant lower in the vitamin B6 group compared with the dimenhydrinate group [5 (4.5%) vs. 36 (53%); $p < 0.01$] No other adverse effect was observed in either group during the 1-week follow-up
Basirat 2009 ⁶⁰	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> 28/32 (87.5%) of ginger group vs. 21/30 (70%) of placebo group reported a subjective improvement in overall symptoms ($p = 0.043$)
Bayreuther 1994 ⁵¹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Belluomini 1994 ⁶²	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Biswas 2011 ⁶³	<ul style="list-style-type: none"> No reported stillbirths, congenital anomalies, neonatal or fetal complications 	<ul style="list-style-type: none"> No difference in reported overall well-being between groups ($p > 0.05$)
Bondok 2006 ⁶⁴	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No patients from the hydrocortisone group but six from the metoclopramide group were readmitted to the intensive care unit for recurrence of severe vomiting ($p < 0.001$)
Capp 2014 ⁶⁵	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference between the groups with respect to sedation or constipation ($p > 0.05$)
Can Gurkan 2008 ⁴³	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Carlsson 2000 ⁶⁶	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Chittumma 2007 ⁶⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Minor side effects such as sedation, heartburn, arrhythmia were reported, but the difference between groups was not significant (% experiencing side effects: ginger 25.4%, vitamin B6 23.8%; $p = 0.80$)

Study	Pregnancy outcomes	Other outcomes
Diggory 1962 ⁶⁸	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Duration of symptoms (weeks): group 1 = 6.4; group 2 = 7.1; group 3 = 3.8; group 4 = 3.5 No statistically significant differences between groups 1 and 2, or 3 and 4, but both groups 3 and 4 significantly improved compared with 1 and 2 ($p < 0.001$ in all cases)
Ditto 1999 ⁶⁹	<ul style="list-style-type: none"> No statistically significant differences reported in terms of gestation at delivery; preterm delivery; caesarean section rate; mean birthweight or neonatal abnormalities (p-value not reported) 	<ul style="list-style-type: none"> Readmission rate was less in diazepam group 1 (4%) than the comparison group 6 (27%). Difference was described as significant but p-value was not reported
Eftekhari 2013 ⁷²	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference in the women's views regarding the effectiveness of treatments using author-defined scales Symptoms of dizziness were significantly higher in the promethazine group ($p = 0.001$)
Einarson 2004 ¹²¹	<ul style="list-style-type: none"> No statistically significant differences reported in terms of miscarriage, terminations, live births, stillbirths, major malformations, mean birthweight or gestational age at birth ($p > 0.05$) 	<ul style="list-style-type: none"> Not reported
Ensiyeh 2009 ⁷⁰	<ul style="list-style-type: none"> Ginger group reported two miscarriages; vitamin B6 group reported one miscarriage No congenital abnormalities or neonatal problems were reported in either group 	<ul style="list-style-type: none"> On day 7, 29/35 women in the ginger group reported an improvement in nausea symptoms, compared with 23/34 in the vitamin B6 group, but this was not statistically significant ($p = 0.52$)
Erez 1971 ⁷¹	<ul style="list-style-type: none"> No statistically significant differences reported in terms of miscarriage, perinatal outcomes and fetal outcomes between groups ($p > 0.05$) 	<ul style="list-style-type: none"> Slight drowsiness was reported by 7% of the hydroxyzine-treated patients
Evans 1993 ⁷³	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Ferreira 2003 ¹²²	<ul style="list-style-type: none"> No statistically significant differences in spontaneous/elective medical abortions, preterm births, live births or major malformations reported ($p > 0.05$) 	<ul style="list-style-type: none"> Lower proportion of patients readmitted in group B compared with the two other groups, but this was not statistically significant ($p = 0.17$) No statistically significant difference reported in terms of average length of stay (days) (group A 3.53 ± 1.69, group B 2.85 ± 1.19, group C 3.25 ± 1.59; $p > 0.05$)
Fischer-Rasmussen 1991 ⁷⁴	<ul style="list-style-type: none"> One miscarriage and one termination were reported Mean birthweight 3585 g (range 2450–5150 g) Mean gestation at delivery: 39.9 weeks (range 36–41 weeks) No congenital abnormalities reported 	<ul style="list-style-type: none"> 19 women (70.4%) stated preference to ginger ($p = 0.003$)
Ghahiri 2011 ⁷⁵	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference in reported headaches, dizziness, sedation or anxiety ($p > 0.05$)
Ghani 2013 ⁷⁶	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Mood and fatigue score improved in both groups [mood score (energy) increased (positive outcome) from 1.94 (SD 1.54) to 4.62 (SD 0.69) in treatment group ($p < 0.001$) and fatigue severity score decreased (positive outcome) from 50.68 (SD 7.66) to 44.92 (SD 6.83) in treatment group ($p < 0.001$)]. Results were not reported for the control group

Study	Pregnancy outcomes	Other outcomes
Guttuso 2010 ³⁹	<ul style="list-style-type: none"> Two congenital defects were reported among seven of the subjects' infants (tethered spinal cord and hydronephrosis) The gestational ages when gabapentin was initiated were 8 and 9 weeks respectively 	<ul style="list-style-type: none"> Subjects gained an average 0.85 lb from baseline to day 21 The initial three subjects enrolled as inpatients were discharged after a mean of 2.3 days after initiating gabapentin therapy None of the subjects required admission/readmission for hyperemesis after starting gabapentin Four subjects experienced mild-moderate side effects of sleepiness or dizziness when first starting gabapentin
Haji Seid Javadi 2013 ⁷⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No side effects reported
Heazell 2006 ⁷⁸	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Median number of women staying > 4 days: (1) intervention 11; (2) comparator 18 ($p < 0.05$, not clear if this is exact p-value) Median length of stay in days: (1) intervention 3 (range 2-4); (2) comparator 3 (range 2-5) ($p > 0.05$) Number of antiemetic doses: (1) intervention 7.1 (range 3-10); (2) comparator 7.4 (range 4-9.8) ($p > 0.05$) Number of antiemetic doses per day: (1) intervention 2.5 (range 1.4-3); (2) comparator 2.3 (range 1.5-2.8) ($p > 0.05$) Total amount of i.v. fluid (l): (1) intervention: 4 (range 2-7); (2) comparator 5 (range 3-6) ($p > 0.05$)
Hsu 1996 ¹²³	<ul style="list-style-type: none"> Gestational age at delivery ranged from 37 to 41 weeks Birthweight ranged from 2766 to 3949 g 	<ul style="list-style-type: none"> Mean reported weight gain: 12.6 lb (range 3-22 lb)
Hsu 2003 ⁷⁹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Subsequent antiemetic administration: (1) intervention 72%; (2) comparator 75% ($p = 1$) Length of emergency department stay: (1) intervention 6.3 hours; (2) comparator 5.5 hours ($p = 0.3$)
Jamigorn 2007 ⁸⁰	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> There was no statistically significant change in maternal weight between the two groups ($p > 0.05$): (1) acupuncture group pre-pregnancy weight (kg) 52.4 (± 7.0); weight on the participation day (kg) 53.5 (± 7.6); weight at the end of the trial (kg) 55.1 (± 7.4); (2) comparator group: pre-pregnancy weight (kg) 49.9 (± 4.7); weight on the participation day (kg) 50.7 (± 4.5); weight at the end of the trial (kg) 50.7 (± 4.6)
Kashifard 2013 ⁸¹	<ul style="list-style-type: none"> All mothers and infants were healthy at the time of birth 	<ul style="list-style-type: none"> None of the patients showed any side effects of the offered medicines
Keating 2002 ⁸²	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Three women in the ginger group lost between 0.57 and 1 kg at 4 weeks compared with weight losses between 0.34 and 0.9 kg for four women in the placebo group in the same time interval (p-value not reported but authors state most women in both groups maintained their weight or gained weight at the 4-week follow-up visit)

Study	Pregnancy outcomes	Other outcomes
Knight 2001 ⁸³	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Anxiety scores at each treatment median (IQR): intervention (1) 8 (IQR 6–9), (2) 8.5 (IQR 6–9), (3) 7 (IQR 6–9), (4) 7 (IQR 4–9); comparator (1) 10 (IQR 7–13), (2) 8 (IQR 6–11), (3) 7 (IQR 5–10), (4) 8 (IQR 5–9). For the anxiety score there was evidence of a time effect ($p = 0.006$) but no evidence of a group effect ($p = 0.4$) or a group-time effect ($p = 0.2$) Depression scores at each treatment median (IQR): intervention (1) 9.5 (IQR 8–15), (2) 9 (IQR 7–11), (3) 8.5 (IQR 7–12), (4) 7 (IQR 5–11); comparator (1) 11 (IQR 8–14), (2) 9 (IQR 7–12), (3) 8 (IQR 7–12), (4) 8 (IQR 6–10). For the depression scores there was evidence of a time effect ($p = 0.002$) and again no group ($p = 0.9$) or group-time interaction ($p = 0.5$)
Koren 2010 ⁸⁴	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> More women in placebo group ($n = 46$, 36%) than Diclectin ($n = 31$, 23.7%) used alternate therapies for NVP ($p = 0.04$) At completion 48.9% of Diclectin group requested compassionate use of medication vs. 32.8% of placebo group ($p = 0.009$)
Koren 2013 ¹²⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Maina 2014 ⁸⁵	<ul style="list-style-type: none"> Two major pregnancy complications occurred in the follow-up: a central venous catheter-related sepsis and a postpartum haemorrhage No adverse fetal outcomes were reported (defined as miscarriage, stillbirth, preterm delivery or low birthweight) One baby was small for gestational age No major or minor birth defects were detected 	<ul style="list-style-type: none"> No statistically significant association to an increase of any adverse effects such as lassitude, drowsiness, dry mouth, headache, dizziness, fainting or skin intolerance as compared with placebo group ($p = 0.2$)
Maltepe 2013 ⁸⁶	<ul style="list-style-type: none"> Median gestation at resolution: 26 weeks in the pre-emptive group vs. 33 weeks in the control group ($p = 0.18$) 	<ul style="list-style-type: none"> More cases of NVP were reported as resolved before labour in the intervention compared with the control group [18/23 (78.2%) vs. 11/22 (50%); $p < 0.002$]
Mao 2009 ⁸⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Markose 2004 ¹²⁴	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
McParlin 2008 ⁸⁸	<ul style="list-style-type: none"> No statistically significant difference in rates of miscarriage, termination of pregnancy, gestation at delivery, birthweight, incidence of small for gestational age, or admissions to the special care baby unit 	<ul style="list-style-type: none"> No statistically significant differences reported in terms of QoL using the SF-36 v2 [physical component score intervention = 42.75 (SD 6.9) vs. comparator = 39.1 (SD 8.6); mental component score intervention = 31.0 (SD 9.8) vs. comparator = 35.1 (SD 12.5); $p > 0.05$] No statistically significant difference between groups in reported satisfaction with care [intervention = 29.2 (SD 3.3) vs. comparator = 29.5 (SD 4.8); $p > 0.05$] Lower total admission time reported for the intervention group [27.2 hours (SD 50.7) vs. the comparator 84.1 hours (SD 86.2); $p = 0.001$]
Mohammadbeigi 2011 ⁸⁹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Monias 1957 ⁹⁰	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported

Study	Pregnancy outcomes	Other outcomes
Moran 2002 ¹²⁵	<ul style="list-style-type: none"> No difference in mean gestation at delivery or birthweight for term infants Two preterm deliveries 	<ul style="list-style-type: none"> Total number of admissions: steroid group = 3 (IQR 1–11); comparator = 1 (IQR 1–7) ($p < 0.005$) Max weight loss (kg): steroid group = 6.5 (IQR 5–8); comparator = 2 (IQR 0.8–3.5) ($p < 0.005$) No serious side effects reported in steroid group
Naeimi Rad 2012 ⁹¹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Narenji 2012 ⁹²	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Satisfaction rate of treatment was higher in the ginger syrup than in the vitamin B6 group (94% vs. 54%) (p-value not available)
Nelson-Piercy 2001 ⁹³	<ul style="list-style-type: none"> No difference in birthweight or gestational age at delivery (excluding the triplet pregnancy) between groups ($p > 0.05$) No difference in the numbers of babies born with birthweights less than the 5th centile ($p > 0.05$) 	<ul style="list-style-type: none"> Length of stay post-randomisation (days): intervention = 7.0 (range 2.0–21.0), comparator = 7.0 (range 2.0–26.0), relative risk = 0.84 (no CI reported) Duration of i.v. fluids (hours): intervention = 72.0 (range 0–168.0), comparator = 96.0 (range 24.0–168.0), relative risk = 0.92 (no CI reported) Well-being rating improvement: intervention = 6.5 (range 1.0–10.0), comparator = 3.5 (range –2.0 to 8.0), relative risk = 0.021 (no CI reported) Weight change (kg): intervention = 1.25 (range –0.5 to 5.0), comparator = –1.0 (range –2.0 to 4.5), relative risk = 0.025 (no CI reported) No. readmitted for hyperemesis: intervention = 5, comparator = 8, relative risk = 1.6 (0.7–3.5)
Neri 2005 ⁹⁴	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Rate of food intake – number of cases improved: intervention (first) 7 (16.2%), (second) 15 (34.8%), (third) 21 (48.8%); comparator (first) 2 (5.2%), (second) 10 (26.3%), (third) 14 (36.8%). There was no statistically significant difference between the two groups ($p > 0.2$). but statistically significant improvement was noted at each time point measured within each treatment compared with baseline ($p < 0.05$ and more specifically $p = 0.001$ in most cases)
Oliveira 2013 ⁹⁵	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Four patients in each group reported sedation ($p > 0.05$)
Ozgoli 2009 ⁹⁶	<ul style="list-style-type: none"> States no adverse effects reported 	<ul style="list-style-type: none"> None of the participants reported any complications during the treatment period
Pasha 2012 ⁹⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Pongrojapaw 2007 ⁴²	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Statistically significant difference in the side effect of drowsiness after treatment reported: dimenhydrinate (77.6%) compared with ginger (5.9%) ($p < 0.01$)
Rosen 2003 ⁹⁸	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> 77% of the intervention group gained weight compared with 54% of the control group ($p = 0.001$) Three dehydration events reported in the study group compared with 12 events in the control group ($p = 0.013$) No statistically significant difference in the use of other medication or ketonuria between groups
Saberi 2013 ¹³	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No side effects experienced

Study	Pregnancy outcomes	Other outcomes
Safari 1998 ⁹⁹	<ul style="list-style-type: none"> No statistically significant difference in the two groups of neonates with respect to birthweight or APGAR scores at 1 and 5 minutes ($p > 0.05$) One patient in the methylprednisolone group was delivered at 35 weeks' gestation of a male infant with Smith–Lemli–Opitz syndrome; this infant subsequently died on the second day after birth 	<ul style="list-style-type: none"> No patients from the methylprednisolone group but five patients from the promethazine group were readmitted for recurrence of vomiting ($p = 0.001$)
Saha 2009 ¹²⁶	<ul style="list-style-type: none"> All pregnancies ended with term deliveries (36–40 weeks' gestation) of healthy infants Mean infant birthweight was 2995 g (2270–4000 g) 	<ul style="list-style-type: none"> Tube-related complications were limited to late tube dislodgement requiring simple replacement via the established percutaneous tract in two patients
Sahakian 1991 ¹⁰⁰	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Smith 2002 ¹⁰¹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Medical Outcomes Study 36 Short Form Health Survey: <ul style="list-style-type: none"> Social function: (1) intervention – day 1 48.3 (SD 28.6), day 14 51.9 (SD 29.0), day 28 54.0 (SD 27.1); (2) comparator – day 1 45.1 (SD 30.3), day 14 50.7 (SD 28.4), day 28 51.4 (SD 32.0); (3) sham – day 1 42.9 (SD 28.5), day 14 48.5 (SD 28.1), day 28 48.9 (SD 28.0); (4) control – day 1 45.3 (SD 28.5), day 14 45.1 (SD 28.7), day 28 37.8 (SD 22.5) ($p = 0.01$, $p = 0.01$ and $p = 0.001$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time respectively) Vitality: (1) intervention – day 1 23.4 (SD 18.0), day 14 30.0 (SD 19.7), day 28 31.9 (SD 20.4); (2) comparator – day 1 21.6 (SD 17.7), day 14 27.4 (SD 23.1), day 28 27.0 (SD 18.9); (3) sham – day 1 22.6 (SD 18.3), day 14 25.8 (SD 18.0), day 28 27.0 (SD 19.0); (4) control – day 1 24.1 (SD 15.4), day 14 22.8 (SD 15.1), day 28 26.0 (SD 17.0) ($p > 0.05$, $p = 0.001$ and $p = 0.05$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time respectively) Physical function: (1) intervention – day 1 63.7 (SD 24.6), day 14 68.5 (SD 26.1), day 28 68.0 (SD 21.0); (2) comparator – day 1 58.7 (SD 26.5), day 14 63.3 (SD 23.1), day 28 64.0 (SD 23.5); (3) sham – day 1 63.1 (SD 25.0), day 14 66.0 (SD 24.5), day 28 66.0 (SD 21.0); (4) control – day 1 63.0 (SD 25.2), day 14 59.4 (SD 27.3), day 28 63.3 (SD 25.0) ($p > 0.05$, $p = 0.01$ and $p = 0.05$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time respectively) Physical role function: (1) intervention – day 1 12.6 (SD 26.4), day 14 7.3 (SD 14.6), day 28 8.7 (SD 15.5); (2) comparator – day 1 11.7 (SD 7.0), day 14 6.3 (SD 13.5), day 28 6.6 (SD 14.1); (3) sham – day 1 11.4 (SD 26.8), day 14 6.0 (SD 11.9), day 28 5.0 (SD 11.7); (4) control – day 1 9.3 (SD 23.0), day 14 4.1 (SD 10.4), day 28 11.9 (SD 25.0) ($p > 0.05$, $p = 0.001$ and $p = 0.05$ for differences in

Study	Pregnancy outcomes	Other outcomes
		<p>SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time respectively)</p> <ul style="list-style-type: none"> ● Bodily pain: (1) intervention – day 1 60.0 (SD 24.5), day 14 65.9 (SD 22.5), day 28 65.2 (SD 22.8); (2) comparator – day 1 60.7 (SD 26.7), day 14 66.5 (SD 24.7), day 28 68.0 (SD 23.0); (3) sham – day 1 59.0 (SD 26.1), day 14 60.0 (SD 24.5), day 28 65.0 (SD 25.1); (4) control – day 1 64.3 (SD 24.7), day 14 65.9 (SD 23.0), day 28 66.6 (SD 22.7) ($p > 0.05$, $p = 0.001$ and $p > 0.05$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time respectively) ● Mental health: (1) intervention – day 1 59.2 (SD 18.0), day 14 66.2 (SD 20.2), day 28 64.7 (SD 18.8); (2) comparator – day 1 56.6 (SD 18.4), day 14 60.0 (SD 20.1), day 28 62.0 (SD 19.1); (3) sham – day 1 57.3 (SD 18.1), day 14 62.0 (SD 17.7), day 28 64.6 (SD 17.7); (4) control – day 1 58.0 (SD 19.7), day 14 58.2 (SD 19.9), day 28 58.6 (SD 20.0) ($p = 0.05$, $p = 0.001$ and $p = 0.01$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time, respectively) ● Emotional role function: (1) intervention – day 1 54.9 (SD 45.5), day 14 61.4 (SD 44.7), day 28 68.7 (SD 43.0); (2) comparator – day 1 47.4 (SD 44.5), day 14 49.1 (SD 45.8), day 28 57.2 (SD 52.0); (3) sham – day 1 54.0 (SD 46.6), day 14 59.7 (SD 45.8), day 28 60.7 (SD 44.4); (4) control – day 1 52.7 (SD 45.4), day 14 54.7 (SD 45.3), day 28 53.2 (SD 45.0) ($p > 0.05$, $p = 0.001$ and $p > 0.05$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time, respectively) ● General health perception: (1) intervention – day 1 67.9 (SD 19.1), day 14 65.3 (SD 19.4), day 28: 67.0 (SD 20.6); (2) comparator – day 1 64.1(SD 21.0), day 14 62.6 (SD 19.0), day 28 63.8 (SD 20.3); (3) sham – day 1 67.7 (SD 20.9), day 14 64.7 (SD 19.4), day 28 66.1 (SD 19.5); (4) control – day 1 67.7 (SD 7.5), day 14 63.4 (SD 18.7), day 28 65.3 (SD 21.6) ($p > 0.05$, $p = 0.001$ and $p > 0.05$ for differences in SF-36 scores across groups, time observation of the SF-36 score at days 1, 14 and 28, and differences in SF-36 scores across all groups over time, respectively)
Smith 2004 ¹⁰²	<ul style="list-style-type: none"> ● No statistically significant differences between groups reported in terms of live birth, congenital abnormalities or birthweight ($p > 0.05$) 	<ul style="list-style-type: none"> ● Not reported
Sripramote 2003 ¹⁰³	<ul style="list-style-type: none"> ● Not reported 	<ul style="list-style-type: none"> ● Minor side effects reported in both groups: sedation (ginger 26.6% vs. vitamin B6 32.8%; $p = 0.439$), and heartburn (ginger 9.4% vs. vitamin B6 6.3%; $p = 0.510$)
Steele 2001 ¹⁰⁴	<ul style="list-style-type: none"> ● Not reported 	<ul style="list-style-type: none"> ● Not reported

Study	Pregnancy outcomes	Other outcomes
Sullivan 1996 ¹⁰⁵	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Hospital stay (days), similar between groups (ondansetron 4.47 ± 2.3 vs. promethazine 4.47 ± 1.5 days; $p = 1.00$) Only reported side effect was sedation; eight women in the promethazine group vs. none in ondansetron group ($p = 0.002$)
Tan 2009 ¹⁰⁷	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference in rates of hospital re-admission ($p > 0.05$)
Tan 2010 ¹⁰⁶	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference in reported well-being score: metoclopramide = 7.6 (SD 2.2), promethazine = 7.1 (SD 2.3); $p = 0.24$ No statistically significant difference in reported hospital stay (days): metoclopramide = 1.8 (IQR 1.5–2.5), promethazine = 1.7 (IQR 1.5–2.4); $p = 0.71$ Statistically significantly more women in the promethazine group reported feeling drowsy ($p = 0.001$) and dizzy ($p < 0.001$)
Tan 2013 ¹⁰⁸	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> No statistically significant difference in hospital stays: D-Saline = 43 ± 21 compared with 48 ± 21 for N-Saline ($p = 0.14$)
Veciana 2001 ¹⁰⁹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Women in the intervention group gained statistically significantly more weight compared with control (2.9 lb vs. 1.2 lb; $p = 0.003$)
Vutyavanich 1995 ⁴¹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Vutyavanich 2001 ¹¹⁰	<ul style="list-style-type: none"> Miscarriages: one in ginger group, three in placebo group Delivery at term: 91.4% in the placebo group, 96.9% in the ginger group No congenital anomalies 	<ul style="list-style-type: none"> On day 7, 88% of ginger group reported symptom improvement vs. 29% of placebo group ($p < 0.001$)
Werntoft 2001 ¹¹¹	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported
Wibowo 2012 ¹¹²	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Change in plasma B6 high dose (78.59 ± 73.89) vs. (low dose: -35.32 ± 89.41) ($p < 0.05$)
Willets 2003 ¹¹³	<ul style="list-style-type: none"> Two miscarriages, one stillbirth and one neonatal death reported in the ginger group Reported birth defects were minor and similar to general hospital population 	<ul style="list-style-type: none"> Not reported
Yost 2003 ¹¹⁴	<ul style="list-style-type: none"> Spontaneous abortion: steroid $n = 2$ (4%), placebo $n = 3$ (6%); $p = 0.6$ Gestational diabetes: steroid $n = 3$ (5%), placebo $n = 3$ (6%); $p = 0.96$ Pregnancy hypertension: steroid $n = 4$ (7%), placebo $n = 8$ (15%); $p = 0.2$ Preterm delivery ≤ 36 weeks: steroid $n = 7$ (13%), placebo $n = 4$ (7%); $p = 0.4$ Caesarean delivery: <ul style="list-style-type: none"> Primary: steroid $n = 6$ (11%), placebo $n = 13$ (24%); $p = 0.06$ Repeat: steroid $n = 4$ (7%), placebo $n = 6$ (11%); $p = 0.5$ 	<ul style="list-style-type: none"> Number of emergency room visits: steroids 0.7 ± 1.2, placebo 0.5 ± 1.0 ($p = 0.4$) Number of admissions: steroids 1.9 ± 1.8, placebo 1.6 ± 1.0 ($p = 0.3$) Women rehospitalised, n (%): steroids 19 (34), placebo 19 (35) ($p = 0.9$) Hospital days, first admission: steroids 1.9 ± 0.9, placebo 2.2 ± 1.2 ($p = 0.5$) Total hospital days, all admissions: steroids 7.6 ± 18.0, placebo 4.3 ± 4.3 ($p = 0.2$)
Zhang 2005 ¹¹⁵	<ul style="list-style-type: none"> Not reported 	<ul style="list-style-type: none"> Not reported

Study	Pregnancy outcomes	Other outcomes
Ziaei 2004 ¹¹⁶	<ul style="list-style-type: none">Not reported	<ul style="list-style-type: none">Drowsiness during first 48 hours: intervention $n = 0$ (0%) vs. comparator $n = 6$ (15%); $p = 0.026$Drowsiness between the third and the 10th days: intervention $n = 0$ (0%) vs. comparator $n = 6$ (15%); $p = 0.026$

APGAR, American Pediatric Gross Assessment Record; D-Saline, dextrose saline; N-Saline, normal saline.

Appendix 9 Systematic review of published economic evaluations: inclusion criteria

A systematic review to identify any existing economic studies will be undertaken. The method will be similar to the review of clinical effectiveness studies except that (i) the inclusion criteria for study design will target cost analysis, cost-effectiveness, cost-utility, cost-consequence and cost-benefit studies; decision model-based analyses and outcomes will be QoL, costs and incremental cost-effectiveness ratios; (ii) quality assessment will be appropriate to the study design, for example the Consensus on Health Economic Criteria list¹³⁰ for economic evaluations and the Philips checklist¹³¹ for model-based analyses; (iii) data considered relevant will be extracted. If available, economic models will be reviewed and utilised where appropriate to inform a model-based economic evaluation based on the above proposed systematic review.

Appendix 10 Cost of drug interventions and recommended daily doses

TABLE 43 Weekly cost of all interventions

Preparation	Patient-initiated first-line intervention	Low-estimate weekly cost	High-estimate weekly cost
Tablets	Vitamin B6: pyridoxine hydrochloride (non-proprietary)	£0.12	£2.59
Tablets	Vitamin B12: cyanocobalamin (non-proprietary)	£0.87	£2.62
Solution	Ginger (FortiCare)	NE	£2.21 (assumes 125 ml per week)
Physical therapy	Acupressure/acupuncture	NE	First appointment: £50–70 ^a Subsequent appointments: £35–50 ^a
Hypnosis	Hypnotherapy	NE	£50–90 for a private hypnotherapy session ^a
Preparation	Clinician-prescribed second-line interventions	Low-estimate weekly cost	High-estimate weekly cost
Antihistamines			
Tablets	Hydroxyzine: Atarax® (Alliance)	£0.91	£1.22
Tablets	Cyclizine (non-proprietary)	£0.74	£2.22
i.v./i.m. injection	Cyclizine: Valoid	NE	£13.65
Tablets	Dimenhydrinate: Arlevert® (Hennig Arzneimittel)	NE	£5.04
Tablets	Chlorpromazine (non-proprietary)	NE	£1.55
i.v./i.m. injection	Chlorpromazine (non-proprietary)	£12.60	£33.60
Tablets	Codeine phosphate (non-proprietary)	£1.09	£2.24
i.v./i.m. injection	Codeine phosphate (non-proprietary)	£24.89	£49.77
Capsules	Benadryl: acrivastine (non-proprietary)	NE	£4.16
Tablets	Cetirizine (non-proprietary)	NE	£0.24
Dopamine antagonists			
Tablets	Promethazine: Phenergan	£0.74	£2.22
i.v./i.m. injection	Promethazine (non-proprietary)	£0.68 (daily) ^a	£1.20 (daily) ^a
Tablets	Prochlorperazine (non-proprietary)	£0.47	£1.42
i.v./i.m. injection	Prochlorperazine (non-proprietary)	NE	£0.52 (daily) ^a
Tablets	Domperidone (non-proprietary)	£0.39	£1.17
Tablets	Metoclopramide (non-proprietary)	£0.22	£0.66
i.v./i.m. injection	Metoclopramide (non-proprietary)	£2.24	£6.72
Serotonin antagonists			
Tablets	Ondansetron (non-proprietary)	NE	£8.69 (daily) ^a
i.v./i.m. injection	Ondansetron (non-proprietary)	NE	£1.00 (daily) ^a
Other drugs			
Tablets	Antiepileptic: gabapentin (non-proprietary)	£0.34	£1.03

continued

TABLE 43 Weekly cost of all interventions (continued)

Preparation	Patient-initiated first-line intervention	Low-estimate weekly cost	High-estimate weekly cost
Tablets	Diazepam	£0.64	£0.70
i.v./i.m. injection	Diazepam (non-proprietary)	£0.90 (daily) ^a	£1.80 (daily) ^a
Rectal tubes	Diazepam (non-proprietary): assume weight of 60 kg	NE	£41.10 (every 12 hours as required) ^a
Tablets	Dicycloverine (non-proprietary)	£12.64	£15.88
Preparation	Clinician-prescribed third-line interventions	Low-estimate weekly cost	High-estimate weekly cost
Corticosteroids			
Tablets	Methylprednisolone: medrone	£0.91	£10.02
i.v./i.m. injection	Methylprednisolone: Solu-Medrone	NE	£17.30 (daily) ^a
Tablets	Prednisone: Lodotra	£12.46	£24.92
Tablets	Prednisolone (non-proprietary)	£0.67	£1.33
i.v./i.m. injection	Prednisolone acetate: Deltastab [®] (Amdipharm Mercury Company Ltd)	£6.87	£54.96
Rectal foam	Prednisolone (non-proprietary)	£34	£68
Tablets	Hydrocortisone (non-proprietary)	£17.03	£29.05
i.v./i.m. injection	Hydrocortisone: Efcortisol	£3.24 (daily) ^a	£19.56 (daily) ^a
NE, not estimated.			
a All costs are weekly, unless otherwise stated.			

TABLE 44 Recommended dose and unit cost for all pharmacological interventions

Preparation	Patient-initiated first-line interventions	Recommended daily dose (low estimate)	Recommended daily dose (high estimate)	Unit cost
Tablets	Vitamin B6	20 mg × 3	50 mg × 3	10 mg, net price 500 = £8.48; 20 mg, net price 500 = £8.53; 50 mg, net price 28 = £3.46
Tablets	Vitamin B12	50 µg	150 µg	50 µg, net price 50 = £6.24
Solution	Ginger (FortiCare)	125 ml weekly ^a	NE	Bottle, 4 × 125 ml = £8.84
Antihistamines				
Tablets	Hydroxyzine: Atarax	25 mg × 3	25 mg × 4	10 mg, net price 84 = £2.18; 25 mg, net price 28 = £1.22
Tablets	Cyclizine (non-proprietary)	50 mg	50 mg × 3	50 mg, net price 100 = £10.58
i.v./i.m. injection	Cyclizine: Valoid	NE	50 mg × 3	50 mg/ml, 1-ml ampoule = £0.65
Tablets	Dimenhydrinate: Arlevert	NE	30 mg × 3	Cinnarizine 20 mg, dimenhydrinate 40 mg, net price 100 = £24.00
Tablets	Chlorpromazine (non-proprietary)	NE	25 mg × 3	25 mg, net price 28 = £2.07; 50 mg, net price 28 = £2.21; 100 mg, net price 28 = £2.21
i.v./i.m. injection	Chlorpromazine (non-proprietary)	25 mg × 3	50 mg × 4	25 mg/ml, 1-ml ampoule = £0.60

TABLE 44 Recommended dose and unit cost for all pharmacological interventions (*continued*)

Preparation	Patient-initiated first-line interventions	Recommended daily dose (low estimate)	Recommended daily dose (high estimate)	Unit cost
Tablets	Codeine phosphate (non-proprietary)	30 mg × 3	60 mg × 3	15 mg, net price 28 = £1.23; 30 mg, net price 28 = £1.46; 60 mg, net price 28 = £2.99
i.v./i.m. injection	Codeine phosphate (non-proprietary)	30 mg × 3	60 mg × 3	60 mg/ml, 1-ml ampoule = £2.37
Capsules	Benadryl: Acrivastine (non-proprietary)	NE	8 mg × 3	8 mg, 12 capsule pack = £2.75; 8 mg, 24 capsule pack = £4.76
Tablets	Cetirizine (non-proprietary)	NE	10 mg	10 mg, net price 30 = £1.01
Dopamine antagonists				
Tablets	Promethazine: Phenergan	10 mg × 2	20 mg × 3	10 mg, net price 56 = £2.96; 25 mg, net price 56 = £4.65
i.v./i.m. injection	Promethazine (non-proprietary)	25 mg	50 mg	25 mg/ml, 1-ml ampoule = £0.68; 25 mg/ml, 2-ml ampoule = £1.20
Tablets	Prochlorperazine (non-proprietary)	5 mg × 2	10 mg × 3	5 mg, net price 28 = £0.95; 5 mg, net price 84 = £1.37
i.v./i.m. injection	Prochlorperazine (non-proprietary)	12.5 mg (when required) ^a	NE	12.5 mg/ml, 1-ml ampoule = £0.52
Tablets	Domperidone (non-proprietary)	10 mg	10 mg × 3	10 mg, net price 30 = £1.67; 10 mg, net price 100 = £5.57
Tablets	Metoclopramide (non-proprietary)	10 mg	10 mg × 3	10 mg, net price 28 = £0.88
i.v./i.m. injection	Metoclopramide (non-proprietary)	10 mg	10 mg × 3	5 mg/ml, 2-ml ampoule = £0.32
Serotonin antagonists				
Tablets	Ondansetron (non-proprietary)	NE	16 mg	4 mg, net price 30 = £5.37; 8 mg, net price 10 = £43.43
i.v./i.m. injection	Ondansetron (non-proprietary)	NE	4 mg	2 mg/ml, 2-ml ampoule = £1.00
Other drugs				
Tablets	Antiepileptic: gabapentin (non-proprietary)	300 mg	300 mg × 3	600 mg, net price 100 = £9.81; 800 mg, net price 100 = £32.22
Tablets	Diazepam	2 mg × 3	30 mg	2 mg, net price 28 = £0.86; 5 mg, net price 28 = £0.88; 10 mg, net price 28 = £0.94
i.v./i.m. injection	Diazepam (non-proprietary)	5 mg × 4	10 mg × 4	5 mg/ml, 2-ml ampoule = £0.45
Rectal tubes	Diazepam (non-proprietary): assume weight of 60 kg	500 µg/kg (repeated after 12 hours as required) ^a	NE	2 mg/ml, net price 1.25 ml (2.5 mg) tube = £1.13; 2.5 ml (5 mg) tube = £1.09; 4 mg/ml, 2.5 ml (10 mg) tube = £1.37
Tablets	Dicycloverine (non-proprietary)	10 mg × 3	20 mg × 3	10 mg, net price 100 = £60.19; 20 mg, net price 84 = £63.52

continued

TABLE 44 Recommended dose and unit cost for all pharmacological interventions (*continued*)

Preparation	Clinician-prescribed third-line interventions	Recommended daily dose (low estimate)	Recommended daily dose (high estimate)	Unit cost
Corticosteroids				
Tablets	Methylprednisolone: Medrone	2 mg	40 mg	2 mg, net price 30 = £3.88; 4 mg, net price 30 = £6.19; 16 mg, net price 30 = £17.17; 100 mg, net price 20 = £48.32
i.v./i.m. injection	Methylprednisolone: Solu-Medrone	NE	1 g	40-mg vial = £1.58; 125-mg vial = £4.75; 500-mg vial = £9.60; 1-g vial = £17.30; 2 g vial = £32.86
Tablets	Prednisone: Lodotra	10 mg	20 mg	1 mg, net price 30 = £26.70; 2 mg, net price 30 = £26.70; 2 mg, net price 100 = £89.00; 5 mg, net price 30 = £26.70; 5 mg, net price 100 = £89.00
Tablets	Prednisolone (non-proprietary)	10 mg	20 mg	1 mg, net price 28 = £1.25; 5 mg, net price 28 = £1.33; 25 mg, net price 56 = £50.00
i.v./i.m. injection	Prednisolone acetate: Deltastab	Varies according to size ^a	NE	25 mg/ml, 1-ml ampoule = £6.87
Rectal foam	Prednisolone (non-proprietary)	1-metred application (20 mg)	1-metred application (20 mg) × 2	20 mg, 14 application canister = £68.00
Tablets	Hydrocortisone (non-proprietary)	20 mg	30 mg	10 mg, net price 30 = £51.46; 20 mg, net price 30 = £73.00
i.v./i.m. injection	Hydrocortisone: Efcortisol	100 mg × 3	500 mg × 4	100 mg/ml, 1-ml ampoule = £1.08; 5-ml ampoule = £4.89

NE, not estimated.

^a Recommended doses are listed on a daily basis, unless otherwise stated.**TABLE 45** Cost of patient-initiated first-line interventions

Pharmacological preparation	Patient-initiated first-line intervention	Total low-estimate weekly cost (£)	Total high-estimate weekly cost (£)
Tablets	Vitamin B6: pyridoxine hydrochloride (non-proprietary)	0.12	2.59
Tablets	Vitamin B12: cyanocobalamin (non-proprietary)	0.87	2.62
Solution	Ginger (FortiCare)	NE	2.21 (assuming 125 ml per week)
Physical therapy	Acupressure/acupuncture	35	50
Hypnosis	Hypnotherapy	50	90

NE, not estimated.

TABLE 46 Cost of patient-initiated first-line interventions following a GP visit

Pharmacological preparation	Patient-initiated first-line intervention	Total low-estimate weekly cost (£)	Total high-estimate weekly cost (£)	GP clinic consultation (£)	Urine ketones strip (£)	Total (£)
Tablets	Vitamin B6: pyridoxine hydrochloride (non-proprietary)	0.12	2.59	45	0.05	45.17–47.64
Tablets	Vitamin B12: cyanocobalamin (non-proprietary)	0.87	2.62	45	0.05	45.92–47.67
Solution	Ginger (FortiCare)	NE	2.21 (assuming 125 ml per week)	45	0.05	47.26

NE, not estimated.

TABLE 47 Cost of clinician-prescribed second-line interventions following a GP visit

Pharmacological preparation	Clinician-prescribed second-line intervention	Total low-estimate weekly cost (£)	Total high-estimate weekly cost (£)	GP clinic consultation (£)	Urine ketones strip (£)	Total (£)
Antihistamines						
Tablets	Cyclizine (non-proprietary)	0.74	2.22	45	0.05	45.79–47.27
Tablets	Hydroxyzine: Atarax	0.91	1.22	45	0.05	45.96–46.27
Tablets	Dimenhydrinate: Arlevert	NE	5.04	45	0.05	50.09
Tablets	Chlorpromazine (non-proprietary)	NE	1.55	45	0.05	46.60
Tablets	Codeine phosphate (non-proprietary)	1.09	2.24	45	0.05	46.14–47.29
Capsules	Benadryl: acrivastine (non-proprietary)	NE	4.16	45	0.05	49.21
Tablets	Cetirizine (non-proprietary)	NE	0.24	45	0.05	45.29
Dopamine antagonists						
Tablets	Domperidone (non-proprietary)	0.39	1.17	45	0.05	45.44–46.22
Tablets	Metoclopramide (non-proprietary)	0.22	0.66	45	0.05	45.27–45.71
Tablets	Prochlorperazine (non-proprietary)	0.47	1.42	45	0.05	45.52–46.47
Tablets	Promethazine: Phenergan	0.74	2.22	45	0.05	45.79–47.27
Serotonin antagonists						
Tablets	Ondansetron (non-proprietary)	NE	60.83	45	0.05	105.88

NE, not estimated.

TABLE 48 Cost of clinician-prescribed second-line interventions if attending hospital as a 'day case'

Pharmacological preparation	Clinician-prescribed second-line interventions	Total low-estimate daily cost x 2 (£)	Total high-estimate daily cost x 2 (£)	Inpatient excess bed-days x 2 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) x 1 (£)	Normal saline + a proportional amount of potassium chloride x 2 + cost of administering the fluid over 2 days (£)	Total (£)
Antihistamines															
Tablets	Hydroxyzine: Atarax	0.13	0.17	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.95–285.99
Tablets	Cyclizine (non-proprietary)	0.11	0.32	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.93–286.14
i.v./i.m. injection	Cyclizine: Valoid	NE	1.95	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	287.77
Tablets	Chlorpromazine (non-proprietary)	NE	0.22	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.04
i.v./i.m. injection	Chlorpromazine (non-proprietary)	1.80	4.80	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	287.62–290.62
Tablets	Dimenhydrinate: Arlevert	NE	0.72	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.54
Tablets	Codene phosphate (non-proprietary)	0.16	0.32	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.98–286.14
i.v./i.m. injection	Doxylamine: codeine phosphate (non-proprietary)	3.56	7.11	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	289.38–292.93
Capsules	Benadryl: Acrivastine (non-proprietary)	NE	0.59	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.41
Tablets	Cetirizine (non-proprietary)	NE	0.03	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.85

TABLE 48 Cost of clinician-prescribed second-line interventions if attending hospital as a 'day case' (continued)

Pharmacological preparation	Clinician-prescribed second-line interventions	Total low-estimate daily cost x 2 (£)	Total high-estimate daily cost x 2 (£)	Inpatient excess bed-days x 2 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) x 1 (£)	Normal saline + a proportional amount of potassium chloride x 2 + cost of administering the fluid over 2 days (£)	Total (£)
Dopamine antagonists															
Tablets	Promethazine: Phenergan	0.11	0.32	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.93–286.14
i.v./i.m. injection	Promethazine (non-proprietary)	0.68	1.20	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.50–287.02
Tablets	Prochlorperazine (non-proprietary)	0.07	0.20	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.89–286.02
i.v./i.m. injection	Prochlorperazine NE (non-proprietary)	NE	0.52	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.34
Tablets	Domperidone (non-proprietary)	0.06	0.17	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.88–285.99
Tablets	Metoclopramide (non-proprietary)	0.03	0.09	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	285.85–285.91
i.v./i.m. injection	Metoclopramide (non-proprietary)	0.32	0.96	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.14–286.78
Serotonin antagonists															
Tablets	Ondansetron NE (non-proprietary)	NE	8.69	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	294.51
i.v./i.m. injection	Ondansetron NE (non-proprietary)	NE	1.00	122	38	8.51	0.05	6.80	13.55	2.96	5.84	4.94	0.08	83.09	286.82
NE, not estimated.															

TABLE 49 Cost of clinician-prescribed second-line interventions if admitted as an inpatient

Pharmacological preparation	Clinician-prescribed second-line interventions	Total low-estimate daily cost x 2 (£)	Total high-estimate daily cost x 2 (£)	Inpatient excess bed-days x 2 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) x 1 (£)	Normal saline + a proportional amount of potassium chloride x 2 + cost of administering the fluid over 2 days (£)	Total (£)
Antihistamines															
i.v./i.m. injection	Cyclizine: Valoid	NE	3.90	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	796.78
Tablets	Chlorpromazine (non-proprietary)	NE	0.44	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.32
i.v./i.m. injection	Chlorpromazine (non-proprietary)	3.60	9.60	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	796.48–802.48
Dopamine antagonists															
i.v./i.m. injection	Metoclopramide (non-proprietary)	0.64	1.92	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.52–794.80
Tablets	Prochlorperazine (non-proprietary)	0.13	0.41	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.01–793.29
i.v./i.m. injection	Prochlorperazine (non-proprietary)	NE	1.04	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.92
Tablets	Domperidone (non-proprietary)	0.11	0.33	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	792.99–793.21
Serotonin antagonists															
i.v./i.m. injection	Ondansetron (non-proprietary)	NE	2.00	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	794.88
Other drugs															
Tablets	Antiepileptic: gabapentin (non-proprietary)	0.10	0.29	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	792.98–793.17
Tablets	Benzodiazepines: diazepam (non-proprietary)	0.18	0.20	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.06–793.08

TABLE 49 Cost of clinician-prescribed second-line interventions if admitted as an inpatient (continued)

Pharmacological preparation	Clinician-prescribed second-line interventions	Total low-estimate daily cost x 2 (£)	Total high-estimate daily cost x 2 (£)	Inpatient excess bed-days x 2 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) x 1 (£)	Normal saline + a proportional amount of potassium chloride x 2 + cost of administering the fluid over 2 days (£)	Total (£)
i.v./i.m. injection	Diazepam (non-proprietary)	1.80	3.60	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	794.68–796.48
Rectal tubes	Diazepam (non-proprietary)	NE	41.10	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	833.98
Tablets	Dicycloverine (non-proprietary)	3.61	4.54	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	796.49–797.42

NE, not estimated.

TABLE 50 Cost of clinician-prescribed second-line interventions × 2 if admitted as an inpatient

Clinician-prescribed second-line intervention	Total daily cost (two antiemetics) × 2 (£)	Inpatient excess bed-days × 2 (£)	Urinary test (urine culture) × 2 (£)	Urine ketones strip × 2 (£)	Liver function test × 2 (£)	Thyroid function test × 2 (£)	Glucose × 2 (£)	Urea and electrolytes × 2 (£)	Full blood count × 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) × 1 (£)	Normal saline + a proportional amount of potassium chloride × 2 + cost of administering the fluid over 2 days (£)	Total (£)
Cheapest combination	0.50	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	793.38
Most expensive combination	50.70	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	843.58

TABLE 51 Cost of clinician-prescribed third-line interventions if admitted as an inpatient

Clinician-prescribed third-line intervention	Total daily cost (three antiemetic) × 3 (£)	Total daily cost (corticosteroid) × 2 (£)	Inpatient excess bed-days × 5 (£)	Urinary test (urine culture) × 5 (£)	Urine ketones strip × 5 (£)	Liver function test × 5 (£)	Thyroid function test × 5 (£)	Glucose × 5 (£)	Urea and electrolytes × 5 (£)	Full blood count × 5 (£)	Treatment for thromboembolism (4-night stay) (£)	Thiamine supplement (Pabrinex) × 1 (£)	Normal saline + a proportional amount of potassium chloride × 5 + cost of administering the fluid over 5 days (£)	Total (£)
Cheapest combination	1.26	0.38 (prednisolone tablets)	1325	42.55	0.25	34.00	67.75	14.80	29.20	24.70	36.60	2.25	415.45	1994.19
Most expensive combination	82.86	39.12 (hydrocortisone injection)	1325	42.55	0.25	34.00	67.75	14.80	29.20	24.70	36.60	2.25	415.45	2114.53

TABLE 52 Cost of day case management compared with inpatient management

Day case	244	38	17.02	0.10	13.60	27.10	5.92	11.68	9.88	0.16	166.18	533.64	
		Obstetrics unit x 2 (£)	Ultrasound Scan (<20 minutes) x 1 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Full blood count x 2 (£)	Thiamine supplement x 2 (£)	Normal saline and a proportional amount of potassium chloride and cost of administering the fluid on 2 days (£)	Total (£)
Inpatient	530	17.02	0.10	13.60	27.10	5.92	11.68	9.88	9.15	2.25	166.18	792.88	
		Inpatient excess bed-days x 2 (£)	Urinary test (urine culture) x 2 (£)	Urine ketones strip x 2 (£)	Liver function test x 2 (£)	Thyroid function test x 2 (£)	Glucose x 2 (£)	Urea and electrolytes x 2 (£)	Treatment for thromboembolism (1-night stay) (£)	Thiamine supplement (Pabrinex) (£)	Normal saline and a proportional amount of potassium chloride and cost of administering the fluid over 2 days (£)	Total (£)	

TABLE 53 Cost comparisons of patient-initiated first-line interventions

Comparison	Implied valuation	Effect size	Evidence on effect
Vitamin B6 : ginger	1.2 : 1	Moderate	Little evidence on effect
Vitamin B12 : vitamin B6	1.01 : 1	Small	Unknown
Vitamin B12 : ginger	1.2 : 1	Moderate	Unknown
Acupressure/acupuncture : vitamin B12	19.1 : 1	Large	Unknown
Acupressure/acupuncture : vitamin B6	19.3 : 1	Large	In the comparison between acupressure and vitamin B6, there was an improvement in both groups but no difference between groups
Acupressure/acupuncture : ginger	22.6 : 1	Large	Ginger looks promising in reducing symptoms when compared with acupressure, but findings are not conclusive
Hypnotherapy : acupressure/acupuncture	1.8 : 1	Large	Unknown
Hypnotherapy : vitamin B12	34.4 : 1	Large	Unknown
Hypnotherapy : vitamin B6	34.7 : 1	Large	Unknown
Hypnotherapy : ginger	40.7 : 1	Large	Unknown
Ginger : placebo	Not assessable	Not assessable	Ginger looks promising in reducing symptoms when compared with placebo, but findings are not conclusive
Acupressure/acupuncture : placebo	Not assessable	Not assessable	Acupressure looks promising in reducing symptoms when compared with placebo in a small number of studies while the rest show no difference between the groups
Vitamin B6 : placebo	Not assessable	Not assessable	Vitamin B6 looks promising in reducing symptoms when compared with placebo but findings are not conclusive

TABLE 54 Cost comparisons of patient-initiated first-line interventions following a GP visit

Comparison	Implied valuation	Effect size	Evidence on effect
Vitamin B6 : ginger	1.008 : 1	Small	Little evidence on effect
Vitamin B12 : vitamin B6	1.0006 : 1	Small	Unknown
Vitamin B12 : ginger	1.009 : 1	Small	Unknown
Ginger : placebo	Not assessable	Not assessable	Ginger looks promising in reducing symptoms when compared with placebo, but findings are not conclusive
Vitamin B6 : placebo	Not assessable	Not assessable	Vitamin B6 looks promising in reducing symptoms when compared with placebo, but findings are not conclusive

TABLE 55 Cost comparisons of clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit

Comparison	Implied valuation	Effect size	Evidence on effect
Metoclopramide : cetirizine	1.009 : 1	Small	Unknown
Domperidone : metoclopramide	1.01 : 1	Small	Unknown
Domperidone : cetirizine	1.02 : 1	Small	Unknown
Hydroxyzine : domperidone	1.001 : 1	Small	Unknown
Hydroxyzine : metoclopramide	1.01 : 1	Small	Unknown
Hydroxyzine : cetirizine	1.02 : 1	Small	Unknown
Prochlorperazine : hydroxyzine	1.004 : 1	Small	Unknown
Prochlorperazine : domperidone	1.005 : 1	Small	Unknown
Prochlorperazine : metoclopramide	1.02 : 1	Small	Unknown
Prochlorperazine : cetirizine	1.03 : 1	Small	Unknown
Chlorpromazine : prochlorperazine	1.003 : 1	Small	Unknown
Chlorpromazine : hydroxyzine	1.007 : 1	Small	Unknown
Chlorpromazine : domperidone	1.008 : 1	Small	Unknown
Chlorpromazine : metoclopramide	1.02 : 1	Small	Unknown
Chlorpromazine : cetirizine	1.03 : 1	Small	Unknown
Cyclizine : chlorpromazine	1.01 : 1	Small	Unknown
Cyclizine : prochlorperazine	1.02 : 1	Small	Unknown
Cyclizine : hydroxyzine	1.02 : 1	Small	Unknown
Cyclizine : domperidone	1.02 : 1	Small	Unknown
Cyclizine : metoclopramide	1.03 : 1	Small	Unknown
Cyclizine : cetirizine	1.04 : 1	Small	Unknown
Promethazine : cyclizine	1 : 1	Small	Unknown
Promethazine : chlorpromazine	1.01 : 1	Small	Unknown
Promethazine : prochlorperazine	1.02 : 1	Small	Unknown
Promethazine : hydroxyzine	1.02 : 1	Small	Unknown
Promethazine : domperidone	1.02 : 1	Small	Unknown
Promethazine : metoclopramide	1.03 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Promethazine : cetirizine	1.04 : 1	Small	Unknown
Codeine phosphate : promethazine	1.0004 : 1	Small	Unknown
Codeine phosphate : cyclizine	1.0004 : 1	Small	Unknown
Codeine phosphate : chlorpromazine	1.01 : 1	Small	Unknown
Codeine phosphate : prochlorperazine	1.02 : 1	Small	Unknown
Codeine phosphate : hydroxyzine	1.02 : 1	Small	Unknown
Codeine phosphate : domperidone	1.02 : 1	Small	Unknown
Codeine phosphate : metoclopramide	1.03 : 1	Small	Unknown

continued

TABLE 55 Cost comparisons of clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Codeine phosphate : cetirizine	1.04 : 1	Small	Unknown
Benadryl : codeine phosphate	1.04 : 1	Small	Unknown
Benadryl : promethazine	1.04 : 1	Small	Unknown
Benadryl : cyclizine	1.04 : 1	Small	Unknown
Benadryl : chlorpromazine	1.06 : 1	Modest	Unknown
Benadryl : prochlorperazine	1.06 : 1	Modest	Unknown
Benadryl : hydroxyzine	1.06 : 1	Modest	Unknown
Benadryl : domperidone	1.06 : 1	Modest	Unknown
Benadryl : metoclopramide	1.08 : 1	Modest	Unknown
Benadryl : cetirizine	1.09 : 1	Modest	Unknown
Dimenhydrinate : benadryl	1.02 : 1	Small	Unknown
Dimenhydrinate : codeine phosphate	1.06 : 1	Modest	Unknown
Dimenhydrinate : promethazine	1.06 : 1	Modest	Unknown
Dimenhydrinate : cyclizine	1.06 : 1	Modest	Unknown
Dimenhydrinate : chlorpromazine	1.07 : 1	Modest	Unknown
Dimenhydrinate : prochlorperazine	1.08 : 1	Modest	Unknown
Dimenhydrinate : hydroxyzine	1.08 : 1	Modest	Unknown
Dimenhydrinate : domperidone	1.08 : 1	Modest	Unknown
Dimenhydrinate : metoclopramide	1.1 : 1	Modest	Unknown
Dimenhydrinate : cetirizine	1.1 : 1	Modest	Unknown
Ondansetron : dimenhydrinate	2.1 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron : benadryl	2.2 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron : codeine phosphate	2.2 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron : promethazine	2.2 : 1	Large	Unknown
Ondansetron : cyclizine	2.2 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron : chlorpromazine	2.3 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron : prochlorperazine	2.3 : 1	Large	Unknown
Ondansetron : hydroxyzine	2.3 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron : domperidone	2.3 : 1	Large	Unknown

TABLE 55 Cost comparisons of clinician-prescribed second-line interventions (oral antiemetics only) following a GP visit (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Ondansetron : metoclopramide	2.3 : 1	Large	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron : cetirizine	2.3 : 1	Large	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Antihistamines : placebo	Not assessable	Not assessable	Antihistamines appear to be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case'

Comparison	Implied valuation	Effect size	Evidence on effect
Metoclopramide (tablets) : cetirizine (tablets)	1.0002 : 1	Small	Unknown
Domperidone (tablets) : metoclopramide (tablets)	1.0003 : 1	Small	Unknown
Domperidone (tablets) : cetirizine (tablets)	1.0005 : 1	Small	Unknown
Hydroxyzine (tablets) : domperidone (tablets)	1 : 1	Small	Unknown
Hydroxyzine (tablets) : metoclopramide (tablets)	1.0003 : 1	Small	Unknown
Hydroxyzine (tablets) : cetirizine (tablets)	1.0005 : 1	Small	Unknown
Prochlorperazine (tablets) : hydroxyzine (tablets)	1.0001 : 1	Small	Unknown
Prochlorperazine (tablets) : domperidone (tablets)	1.0001 : 1	Small	Unknown
Prochlorperazine (tablets) : metoclopramide (tablets)	1.0004 : 1	Small	Unknown
Prochlorperazine (tablets) : cetirizine (tablets)	1.0006 : 1	Small	Unknown
Chlorpromazine (tablets) : prochlorperazine (tablets)	1.00007 : 1	Small	Unknown
Chlorpromazine (tablets) : hydroxyzine (tablets)	1.0002 : 1	Small	Unknown
Chlorpromazine (tablets) : domperidone (tablets)	1.0002 : 1	Small	Unknown
Chlorpromazine (tablets) : metoclopramide (tablets)	1.0005 : 1	Small	Unknown
Chlorpromazine (tablets) : cetirizine (tablets)	1.0007 : 1	Small	Unknown
Cyclizine (tablets) : chlorpromazine (tablets)	1.0003 : 1	Small	Unknown

continued

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Cyclizine (tablets) : prochlorperazine (tablets)	1.0004 : 1	Small	Unknown
Cyclizine (tablets) : hydroxyzine (tablets)	1.0005 : 1	Small	Unknown
Cyclizine (tablets) : domperidone (tablets)	1.0005 : 1	Small	Unknown
Cyclizine (tablets) : metoclopramide (tablets)	1.0008 : 1	Small	Unknown
Cyclizine (tablets) : cetirizine (tablets)	1.001 : 1	Small	Unknown
Codeine phosphate (tablets) : cyclizine (tablets)	1 : 1	Small	Unknown
Codeine phosphate (tablets) : chlorpromazine (tablets)	1.0003 : 1	Small	Unknown
Codeine phosphate (tablets) : prochlorperazine (tablets)	1.0004 : 1	Small	Unknown
Codeine phosphate (tablets) : hydroxyzine (tablets)	1.0005 : 1	Small	Unknown
Codeine phosphate (tablets) : domperidone (tablets)	1.0005 : 1	Small	Unknown
Codeine phosphate (tablets) : metoclopramide (tablets)	1.0008 : 1	Small	Unknown
Codeine phosphate (tablets) : cetirizine (tablets)	1.001 : 1	Small	Unknown
Promethazine (tablets) : codeine Phosphate (tablets)	1 : 1	Small	Unknown
Promethazine (tablets) : cyclizine (tablets)	1 : 1	Small	Unknown
Promethazine (tablets) : chlorpromazine (tablets)	1.0003 : 1	Small	Unknown
Promethazine (tablets) : prochlorperazine (tablets)	1.0004 : 1	Small	Unknown
Promethazine (tablets) : hydroxyzine (tablets)	1.0005 : 1	Small	Unknown
Promethazine (tablets) : domperidone (tablets)	1.0005 : 1	Small	Unknown
Promethazine (tablets) : metoclopramide (tablets)	1.0008 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Promethazine (tablets) : cetirizine (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : promethazine (tablets)	1.0007 : 1	Small	Unknown
Prochlorperazine (injection) : codeine phosphate (tablets)	1.0007 : 1	Small	Unknown
Prochlorperazine (injection) : cyclizine (tablets)	1.0007 : 1	Small	Unknown
Prochlorperazine (injection) : chlorpromazine (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : prochlorperazine (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : hydroxyzine (tablets)	1.001 : 1	Small	Unknown

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Prochlorperazine (injection) : domperidone (tablets)	1.001 : 1	Small	Unknown
Prochlorperazine (injection) : metoclopramide (tablets)	1.002 : 1	Small	Unknown
Prochlorperazine (injection) : cetirizine (tablets)	1.002 : 1	Small	Unknown
Benadryl (capsules) : prochlorperazine (injection)	1.0002 : 1	Small	Unknown
Benadryl (capsules) : promethazine (tablets)	1.0009 : 1	Small	Unknown
Benadryl (capsules) : codeine phosphate (tablets)	1.0009 : 1	Small	Unknown
Benadryl (capsules) : cyclizine (tablets)	1.0009 : 1	Small	Unknown
Benadryl (capsules) : chlorpromazine (tablets)	1.001 : 1	Small	Unknown
Benadryl (capsules) : prochlorperazine (tablets)	1.001 : 1	Small	Unknown
Benadryl (capsules) : hydroxyzine (tablets)	1.001 : 1	Small	Unknown
Benadryl (capsules) : domperidone (tablets)	1.001 : 1	Small	Unknown
Benadryl (capsules) : metoclopramide (tablets)	1.002 : 1	Small	Unknown
Benadryl (capsules) : cetirizine (tablets)	1.002 : 1	Small	Unknown
Dimenhydrinate (tablets) : benadryl (capsules)	1.0005 : 1	Small	Unknown
Dimenhydrinate (tablets) : prochlorperazine (injection)	1.0007 : 1	Small	Unknown
Dimenhydrinate (tablets) : promethazine (tablets)	1.001 : 1	Small	Unknown
Dimenhydrinate (tablets) : codeine phosphate (tablets)	1.001 : 1	Small	Unknown
Dimenhydrinate (tablets) : cyclizine (tablets)	1.001 : 1	Small	Unknown
Dimenhydrinate (tablets) : chlorpromazine (tablets)	1.002 : 1	Small	Unknown
Dimenhydrinate (tablets) : prochlorperazine (tablets)	1.002 : 1	Small	Unknown
Dimenhydrinate (tablets) : hydroxyzine (tablets)	1.002 : 1	Small	Unknown
Dimenhydrinate (tablets) : domperidone (tablets)	1.002 : 1	Small	Unknown
Dimenhydrinate (tablets) : metoclopramide (tablets)	1.002 : 1	Small	Unknown
Dimenhydrinate (tablets) : cetirizine (tablets)	1.002 : 1	Small	Unknown

continued

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Metoclopramide (injection) : dimenhydrinate (tablets)	1.0008 : 1	Small	Unknown
Metoclopramide (injection) : benadryl (capsules)	1.001 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (injection)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : promethazine (tablets)	1.002 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Metoclopramide (injection) : codeine phosphate (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : cyclizine (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : chlorpromazine (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : hydroxyzine (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : domperidone (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : metoclopramide (tablets)	1.003 : 1	Small	Unknown
Metoclopramide (injection) : cetirizine (tablets)	1.003 : 1	Small	Unknown
Ondansetron (injection) : metoclopramide (injection)	1.0001 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (injection) : dimenhydrinate (tablets)	1.001 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : benadryl (capsules)	1.001 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : prochlorperazine (injection)	1.002 : 1	Small	Unknown
Ondansetron (injection) : promethazine (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : codeine phosphate (tablets)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : cyclizine (tablets)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Ondansetron (injection) : chlorpromazine (tablets)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : prochlorperazine (tablets)	1.003 : 1	Small	Unknown
Ondansetron (injection) : hydroxyzine (tablets)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : domperidone (tablets)	1.003 : 1	Small	Unknown
Ondansetron (injection) : metoclopramide (tablets)	1.003 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (injection) : cetirizine (tablets)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Promethazine (injection) : ondansetron (injection)	1.0007 : 1	Small	Unknown
Promethazine (injection) : metoclopramide (injection)	1.0008 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Promethazine (injection) : dimenhydrinate (tablets)	1.002 : 1	Small	Unknown
Promethazine (injection) : benadryl (capsules)	1.002 : 1	Small	Unknown
Promethazine (injection) : prochlorperazine (injection)	1.002 : 1	Small	Unknown
Promethazine (injection) : promethazine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : codeine phosphate (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : cyclizine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : chlorpromazine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : prochlorperazine (tablets)	1.003 : 1	Small	Unknown
Promethazine (injection) : hydroxyzine (tablets)	1.004 : 1	Small	Unknown
Promethazine (injection) : domperidone (tablets)	1.004 : 1	Small	Unknown
Promethazine (injection) : metoclopramide (tablets)	1.004 : 1	Small	Limited data suggest that promethazine is as effective as metoclopramide in reducing the symptoms of NVP
Promethazine (injection) : cetirizine (tablets)	1.004 : 1	Small	Unknown

continued

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Cyclizine (injection) : promethazine (injection)	1.003 : 1	Small	Unknown
Cyclizine (injection) : ondansetron (injection)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Cyclizine (injection) : metoclopramide (injection)	1.003 : 1	Small	Unknown
Cyclizine (injection) : dimenhydrinate (tablets)	1.004 : 1	Small	Unknown
Cyclizine (injection) : benadryl (capsules)	1.005 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (injection)	1.005 : 1	Small	Unknown
Cyclizine (injection) : promethazine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : codeine phosphate (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : cyclizine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : chlorpromazine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : hydroxyzine (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : domperidone (tablets)	1.006 : 1	Small	Unknown
Cyclizine (injection) : metoclopramide (tablets)	1.007 : 1	Small	Unknown
Cyclizine (injection) : cetirizine (tablets)	1.007 : 1	Small	Unknown
Chlorpromazine (injection) : cyclizine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : promethazine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : ondansetron (injection)	1.01 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Chlorpromazine (injection) : metoclopramide (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : dimenhydrinate (tablets)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : benadryl (capsules)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : promethazine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : codeine phosphate (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : cyclizine (tablets)	1.02 : 1	Small	Unknown

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Chlorpromazine (injection) : chlorpromazine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : hydroxyzine (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : domperidone (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : metoclopramide (tablets)	1.02 : 1	Small	Unknown
Chlorpromazine (injection) : cetirizine (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : chlorpromazine (injection)	1.008 : 1	Small	Unknown
Doxylamine (injection) : cyclizine (injection)	1.02 : 1	Small	Unknown
Doxylamine (injection) : promethazine (injection)	1.02 : 1	Small	Unknown
Doxylamine (injection) : ondansetron (injection)	1.02 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Doxylamine (injection) : metoclopramide (injection)	1.02 : 1	Small	Unknown
Doxylamine (injection) : dimenhydrinate (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : benadryl (capsules)	1.02 : 1	Small	Unknown
Doxylamine (injection) : prochlorperazine (injection)	1.02 : 1	Small	Unknown
Doxylamine (injection) : promethazine (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : codeine phosphate (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : cyclizine (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : chlorpromazine (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : prochlorperazine (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : hydroxyzine (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : domperidone (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : metoclopramide (tablets)	1.02 : 1	Small	Unknown
Doxylamine (injection) : cetirizine (tablets)	1.02 : 1	Small	Unknown
Ondansetron (tablets) : doxylamine (injection)	1.005 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects

continued

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Ondansetron (tablets) : chlorpromazine (injection)	1.01 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : cyclizine (injection)	1.02 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : promethazine (injection)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : ondansetron (injection)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : metoclopramide (injection)	1.03 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (tablets) : dimenhydrinate (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : benadryl (capsules)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : prochlorperazine (injection)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : promethazine (tablets)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : codeine phosphate (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : cyclizine (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : chlorpromazine (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : prochlorperazine (tablets)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : hydroxyzine (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (tablets) : domperidone (tablets)	1.03 : 1	Small	Unknown
Ondansetron (tablets) : metoclopramide (tablets)	1.03 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (tablets) : cetirizine (tablets)	1.03 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects

TABLE 56 Cost comparisons of clinician-prescribed second-line interventions if attending hospital as a 'day case' (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Antihistamines : placebo	Not assessable	Not assessable	Antihistamines appear to be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required

TABLE 57 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient

Comparison	Implied valuation	Effect size	Evidence on effect
Gabapentin (tablets) : diazepam (tablets)	1.0001 : 1	Small	Unknown
Domperidone (tablets) : gabapentin (tablets)	1.00005 : 1	Small	Unknown
Domperidone (tablets) : diazepam (tablets)	1.0002 : 1	Small	Unknown
Prochlorperazine (tablets) : domperidone (tablets)	1.0001 : 1	Small	Unknown
Prochlorperazine (tablets) : gabapentin (tablets)	1.0002 : 1	Small	Unknown
Prochlorperazine (tablets) : diazepam (tablets)	1.0003 : 1	Small	Unknown
Chlorpromazine (tablets) : prochlorperazine (tablets)	1.00004 : 1	Small	Unknown
Chlorpromazine (tablets) : domperidone (tablets)	1.0001 : 1	Small	Unknown
Chlorpromazine (tablets) : gabapentin (tablets)	1.0002 : 1	Small	Unknown
Chlorpromazine (tablets) : diazepam (tablets)	1.0003 : 1	Small	Unknown
Prochlorperazine (injection) : chlorpromazine (tablets)	1.0008 : 1	Small	Unknown
Prochlorperazine (injection) : prochlorperazine (tablets)	1.0008 : 1	Small	Unknown
Prochlorperazine (injection) : domperidone (tablets)	1.0009 : 1	Small	Unknown
Prochlorperazine (injection) : gabapentin (tablets)	1.0009 : 1	Small	Unknown
Prochlorperazine (injection) : diazepam (tablets)	1.001 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (injection)	1.001 : 1	Small	Unknown
Metoclopramide (injection) : chlorpromazine (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : prochlorperazine (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : domperidone (tablets)	1.002 : 1	Small	Unknown

continued

TABLE 57 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient (continued)

Comparison	Implied valuation	Effect size	Evidence on effect
Metoclopramide (injection) : gabapentin (tablets)	1.002 : 1	Small	Unknown
Metoclopramide (injection) : diazepam (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : metoclopramide (injection)	1.0001 : 1	Small	Evidence comparing ondansetron with metoclopramide showed mixed results, with both improving symptoms. Ondansetron was found to be more effective at reducing symptoms of vomiting than metoclopramide after 4 days
Ondansetron (injection) : prochlorperazine (injection)	1.001 : 1	Small	Unknown
Ondansetron (injection) : chlorpromazine (tablets)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Ondansetron (injection) : prochlorperazine (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : domperidone (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : gabapentin (tablets)	1.002 : 1	Small	Unknown
Ondansetron (injection) : diazepam (tablets)	1.002 : 1	Small	Unknown
Diazepam (injection) : ondansetron (injection)	1.002 : 1	Small	Unknown
Diazepam (injection) : metoclopramide (injection)	1.002 : 1	Small	Unknown
Diazepam (injection) : prochlorperazine (injection)	1.003 : 1	Small	Unknown
Diazepam (injection) : chlorpromazine (tablets)	1.004 : 1	Small	Unknown
Diazepam (injection) : prochlorperazine (tablets)	1.004 : 1	Small	Unknown
Diazepam (injection) : domperidone (tablets)	1.004 : 1	Small	Unknown
Diazepam (injection) : gabapentin (tablets)	1.004 : 1	Small	Unknown
Diazepam (injection) : diazepam (tablets)	1.004 : 1	Small	Unknown
Cyclizine (injection) : diazepam (injection)	1.0004 : 1	Small	Unknown
Cyclizine (injection) : ondansetron (injection)	1.002 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Cyclizine (injection) : metoclopramide (injection)	1.002 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (injection)	1.004 : 1	Small	Unknown
Cyclizine (injection) : chlorpromazine (tablets)	1.004 : 1	Small	Unknown
Cyclizine (injection) : prochlorperazine (tablets)	1.004 : 1	Small	Unknown
Cyclizine (injection) : domperidone (tablets)	1.005 : 1	Small	Unknown
Cyclizine (injection) : gabapentin (tablets)	1.005 : 1	Small	Unknown

TABLE 57 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Cyclizine (injection) : diazepam (tablets)	1.005 : 1	Small	Unknown
Dicycloverine (tablets) : cyclizine (injection)	1.0008 : 1	Small	Unknown
Dicycloverine (tablets) : diazepam (injection)	1.001 : 1	Small	Unknown
Dicycloverine (tablets) : ondansetron (injection)	1.003 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Dicycloverine (tablets) : metoclopramide (injection)	1.003 : 1	Small	Unknown
Dicycloverine (tablets) : prochlorperazine (injection)	1.004 : 1	Small	Unknown
Dicycloverine (tablets) : chlorpromazine (tablets)	1.005 : 1	Small	Unknown
Dicycloverine (tablets) : prochlorperazine (tablets)	1.005 : 1	Small	Unknown
Dicycloverine (tablets) : domperidone (tablets)	1.005 : 1	Small	Unknown
Dicycloverine (tablets) : gabapentin (tablets)	1.005 : 1	Small	Unknown
Dicycloverine (tablets) : diazepam (tablets)	1.005 : 1	Small	Unknown
Chlorpromazine (injection) : dicycloverine (tablets)	1.006 : 1	Small	Unknown
Chlorpromazine (injection) : cyclizine (injection)	1.007 : 1	Small	Unknown
Chlorpromazine (injection) : diazepam (injection)	1.008 : 1	Small	Unknown
Chlorpromazine (injection) : ondansetron (injection)	1.01 : 1	Small	Both ondansetron and antihistamines improve symptoms, with no significant difference in effects
Chlorpromazine (injection) : metoclopramide (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (injection)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : chlorpromazine (tablets)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : prochlorperazine (tablets)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : domperidone (tablets)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : gabapentin (tablets)	1.01 : 1	Small	Unknown
Chlorpromazine (injection) : diazepam (tablets)	1.01 : 1	Small	Unknown
Diazepam (rectal tubes) : chlorpromazine (injection)	1.04 : 1	Small	Unknown
Diazepam (rectal tubes) : dicycloverine (tablets)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : cyclizine (injection)	1.05 : 1	Small	Unknown

continued

TABLE 57 Cost comparisons of clinician-prescribed second-line interventions if admitted as an inpatient (*continued*)

Comparison	Implied valuation	Effect size	Evidence on effect
Diazepam (rectal tubes) : diazepam (injection)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : ondansetron (injection)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : metoclopramide (injection)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : prochlorperazine (injection)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : chlorpromazine (tablets)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : prochlorperazine (tablets)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : domperidone (tablets)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : gabapentin (tablets)	1.05 : 1	Small	Unknown
Diazepam (rectal tubes) : diazepam (tablets)	1.05 : 1	Small	Unknown
Antihistamines : placebo	Not assessable	Not assessable	Antihistamines appear to be better than placebo in reducing the severity of symptoms, but more larger, better-quality studies are required

TABLE 58 Cost comparisons of clinician-prescribed second-line interventions × 2 if admitted as an inpatient

Comparison	Implied valuation	Effect size	Evidence on effect
Most expensive : least expensive	1.06 : 1	Modest	Unknown

TABLE 59 Cost comparisons of clinician-prescribed third-line interventions if admitted as an inpatient

Comparison	Implied valuation	Effect size	Evidence on effect
Most expensive : least expensive	1.06 : 1	Modest	Unknown

TABLE 60 Cost comparison of 2-day day case management with 2-day inpatient management

Comparison	Implied valuation	Effect size	Evidence on effect
Inpatient : day case	1.5 : 1	Large	Results indicate that day case management is as effective at improving severity scores as inpatient management for some women. However, more, larger studies are required to provide definitive results

A decorative graphic consisting of numerous thin, parallel green lines that curve from the left side of the page towards the right, creating a sense of movement and depth.

**EME
HS&DR
HTA
PGfAR
PHR**

Part of the NIHR Journals Library
www.journalslibrary.nihr.ac.uk

This report presents independent research funded by the National Institute for Health Research (NIHR). The views expressed are those of the author(s) and not necessarily those of the NHS, the NIHR or the Department of Health

Published by the NIHR Journals Library