

Title	GooD4Mum: A general practice-based quality improvement collaborative for diabetes prevention in women with previous gestational diabetes
Authors(s)	O'Reilly, Sharleen, Dunbar, James A., Best, James D., et al.
Publication date	2018-11-15
Publication information	O'Reilly, Sharleen, James A. Dunbar, James D. Best, and et al. "GooD4Mum: A General Practice-Based Quality Improvement Collaborative for Diabetes Prevention in Women with Previous Gestational Diabetes" 13, no. 2 (November 15, 2018).
Publisher	Elsevier
Item record/more information	http://hdl.handle.net/10197/10621
Publisher's statement	This is the author's version of a work that was accepted for publication in Primary Care Diabetes. Changes resulting from the publishing process, such as peer review, editing, corrections, structural formatting, and other quality control mechanisms may not be reflected in this document. Changes may have been made to this work since it was submitted for publication. A definitive version was subsequently published in Primary Care Diabetes (13, 2, (2018)) https://doi.org/10.1016/j.pcd.2018.10.006
Publisher's version (DOI)	10.1016/j.pcd.2018.10.006

Downloaded 2023-10-05T14:16:07Z

The UCD community has made this article openly available. Please share how this access benefits you. Your story matters! (@ucd_oa)



© Some rights reserved. For more information

Elsevier Editorial System(tm) for Primary

Care Diabetes

Manuscript Draft

Manuscript Number: PCD-D-17-00290R1

Title: GooD4Mum: a general practice-based quality improvement collaborative for diabetes prevention in women with previous gestational diabetes

Article Type: Original Research

Keywords: diabetes prevention; gestational diabetes; primary care; quality improvement collaboratives; general practice.

Corresponding Author: Dr. Sharleen O'Reilly, Ph.D.

Corresponding Author's Institution: University College Dublin

First Author: Sharleen O'Reilly, Ph.D.

Order of Authors: Sharleen O'Reilly, Ph.D.; James A Dunbar; James D Best; Vincent Versace; Dale Ford; Doris Young; Sophy Shih; Richard Bills; Wendy Shepherdley; Edward D Janus; the MAGDA Study Group

Abstract: Aims:

Gestational diabetes (GDM) and Type 2 diabetes pose tremendous health and economic burdens as worldwide incidence increases. Primary care-based systematic diabetes screening and prevention programs could be effective in women with previous GDM. GooD4Mum aimed to determine whether a Quality Improvement Collaborative (QIC) would improve postpartum diabetes screening and prevention planning in women with previous GDM in general practice.

Methods:

Fifteen general practices within Victoria (Australia) participated in a 12-month QIC, which consisted of baseline and four quarterly audits, guideline-led workshops and Plan-Do-Study-Act feedback cycles after each audit. The primary outcome measures were the proportion of women on local GDM registers completing a diabetes screening test and a diabetes prevention planning consultation within the previous 15 months. Results:

Diabetes screening increased with rates more than doubled from 26% to 61% and postpartum screening increased from 43% to 60%. Diabetes prevention planning consultations did not show the same level of increase (0% to 10%). The recording of body mass index improved (51% to 69%) but those with normal body mass index did not.

Conclusions:

GooD4Mum supported increased diabetes screening and the monitoring of high risk women with previous GDM in general practice.

UCD Institute of Food and Health University College Dublin Belfield Dublin 4 Ireland

Dear A/Professor Blackberry,

GooD4Mum: a general practice-based quality improvement collaborative for diabetes prevention in women with previous gestational diabetes

We would like to thank the reviewers for their time and expert feedback. We appreciate the interest in our paper and feel the observations have enhanced the manuscript's clarity. We have endeavoured to clarify the areas of weakness identified and revised the manuscript accordingly. We have taken the reviewer comments and responded to them individually in the response to review document.

My co-authors and I would be grateful for your consideration of our study as an original research paper in Primary Care Diabetes. The author for correspondence is at the address listed above, email <u>Sharleen.oreilly@ucd.ie</u>

Yours sincerely,

Sharleen O'Reilly, PhD, AdvAPD, RD

Dear A/Prof Blackberry,

We would like to thank the reviewers for their time and expert feedback. We appreciate the interest in our paper and feel the observations have enhanced the manuscript's clarity. We have endeavoured to clarify the areas of weakness identified and revised the manuscript accordingly. We have taken the reviewer comments and responded to them individually below.

Reviewer 1

General comments:

The authors have presented data from an analysis of Australian women with previous GDM and found that GooD4Mum brought significant improvements in the postnatal diabetes screening and diabetes prevention planning consultation among women with GDM in general practice. The analysis presented in this manuscript consists of two parts. First, the authors presented the influence of external factors on the planned activity over time. In the second part, the authors calculated the numbers and the rates of women for every measure included in this project at each stage. Though the aim of this paper is clearly elucidated and the topic is important, the study design and results are weak and did not support the conclusion.

Response:

Thank you for your comments. We appreciate that the quality improvement methodology through the use of Collaboratives is a quasi-experimental design and has inherent issues. The methodology is however well described in the literature and our study does meet the SQUIRE standards for the method. We have reframed our conclusions to better reflect the data.

We now state "*Conclusions:* GooD4Mum supported increased diabetes screening and the monitoring of high risk women with previous GDM in general practice. "

Major comment 1

The study lacks a control group at the same period to check whether the GooD4Mum can increase the rate of participation in the diabetes screening and in the monitor of BMI and other risk factors, due to the possible improvement in awareness over time.

Response:

Clinical guidelines for the follow-up of women who have had GDM exist in many counties but followup rates remain obstinately low.^{1,2} Awareness increasing over time is an unlikely explanation because screening rates have been low despite reminders sent to the women and their GPs from a National Gestational Diabetes Register.³

These guidelines represent the systematic review of randomised trials about **what** should be done but not **how**. e.g. NICE⁴. The Institute for Healthcare Improvement in Boston developed Collaboratives to improve the uptake of evidence. Collaboratives have been used in many countries. They were adapted by Prof Sir John Oldham in UK for use in primary care.⁵ He trained the Australian Government-funded Australian Primary Care Collaboratives team which has have been successful.^{6,7}

By using Collaboratives methodology we were following a decade of Australian Government policy on how to improve the uptake of evidence in primary care. Collaboratives do not include a control group and we acknowledge that this results in a quasi-experimental design. We have amended the text to include this aspect:

"This <u>quasi-experimental research</u> project sought to apply QIC methods to the care of women with previous GDM..."

We have also provided more clarification in the methods on the activities:

"For women with previous GDM, this was engaging in <u>annual</u> diabetes screening and having a consultation where their <u>lifestyle-related modifiable</u> diabetes risk was assessed."

References

- McGovern A, Butler L, Jones S, et al. Diabetes screening after gestational diabetes in England: a quantitative retrospective cohort study. *The British Journal of General Practice*. 2014;64(618):e17-e23. doi:10.3399/bjgp14X676410.
- Goueslard, K; Cottenet, J; Mariet, A-S; Sagot, P; Petit, J-M; Quantin, C. Early screening for type 2 diabetes following gestational diabetes mellitus in France: hardly any impact of the 2010 guidelines. Acta Diabetologica. 2017;54(7):645-651.
- 3. Boyle DIR, Versace VL, Dunbar JA et al. Results of the first recorded evaluation national gestational diabetes mellitus register: challenges in screening, registration, and follow-up for diabetes risk. PLOS ONE 13(8): e0200832. DOI.org/10.1371/journal.pone.0200832
- 4. National Institute for Health and Clinical Excellence. Diabetes in pregnancy: Management of diabetes and its complications from preconception to the postnatal period. NICE, 2008.
- 5. Sic ratio ut componitur: the small book about large system change. Oldham J. Kingsham Press. 2005.
- 6. Knight AW, Caesar C, Ford D, Coughlin A, Frick C. Improving primary care in Australia through the Australian Primary Care Collaborative program: a quality improvement report. BMJ Quality and Safety 2012;21:948-955.
- Knight AW, Ford D, Audehm R, Colagiuri S, Best JD. Australian Primary Care Collaboratives program: improving diabetes care. BMJ Quality and Safety 2012; DOI:10.1136/BMJQS-2011-000460

Major comment 2

Despite an increased rate of women screened for DM, the rate of participation in the diabetes prevention planning is not significant different. It is quite weak to believe that the increased screening of DM is attributable to diabetes prevention planning consultations.

Response:

The general practices viewed the screening as a separate activity to the diabetes prevention planning consultation, which is why we have treated them as such in our analysis. Diabetes screening requires a small amount of GP time (asking a woman to take a test request slip to the nearest phlebotomy service or doing the blood test then and there) while the other activity required a much larger amount of time. The prevention planning session was usually booked in as a separate consultation and/or it was delivered by practice nurses often on a separate day.

We have additional qualitative data that was collected as part of the study from practices but not reported in this paper due to space constraints. We have decided to include a component of this data to add context to the data already included.

The methods now includes:

"Qualitative interviews and focus groups were conducted in each practice upon completion of the QIC activity to explore barriers and enablers to the intervention."

"The qualitative data was analysed thematically by an experienced qualitative researcher (SOR) and coded transcripts were checked by the participants for accuracy of interpretation."

The results now includes:

"The qualitative data proposed several barriers to conducting the diabetes prevention planning consultation: 1) it was only emphasised in the final six months of GooD4Mum and therefore had less time to become embedded within daily practice activity; 2) organising it within normal practice workflows was challenging; 3) women were reluctant to attend it for financial and time reasons; 4) limited lifestyle modification referral options existed; and 5) some GPs and practice nurses lacked confidence to engage in a lifestyle modification consultation."

The discussion now includes:

"Diabetes screening and BMI monitoring in GooD4Mum aligns well with previous QIC diabetes prevention initiatives [15] but providing a diabetes prevention planning consultation was the more challenging component of the project - it had low uptake by both practice staff and women and practice staff reported several barriers to engaging in the activity. In looking at the QIC activities as behaviours that need to be changed and using the Theoretical Domains Framework[25] to map them, the screening activity required minor environmental restructuring to ensure GPs knew which women needed screening alongside some education and training with modelling promoted via teleconferences. These were easier behaviours to change because staff already know they will get reimbursed for doing the blood test and that HbA1c is a useful diabetes indicator. The prevention planning activity was a new process for the practices and required substantial behaviour change. Within the theoretical domains framework it called for: cognitive and interpersonal skills (training practice nurses to perform the tasks and support women to engage in lifestyle change), belief about capability and consequences (both staff and woman), environmental restructuring (needed patient and staff resources plus space to conduct consultation), education and training, persuasion and enablement to influence practice nurses and GPs optimism that the prevention planning consultations was worthwhile. It may simply be that providing a diabetes planning consultation to all women with a history of GDM is not appropriate and that providing the consultation in a more targeted fashion would yield better results. This should be explored in further work."

Major comment 3

Many major confounders in this study were not adjusted in the analysis, e.g. social economic status. Besides, it is still unrevealed whether the characters of the study population at each audit stage are different over time.

Response:

The reviewer points to an important difference between an epidemiological study or a controlled trial and improvement work. The population does change over time as more women are entered into the register and a higher proportion are recalled. Numerator and denominator are both changing.

For these reasons, the standard tool in improvement work is the *run chart* developed by the Institute of Healthcare Improvement¹. Its purpose is to help improvement teams formulate aims by

depicting how well processes are performing. It helps the practice teams determine when changes are true improvements by displaying a pattern of data that they can observe as they make changes. Run charts also indicate the direction of work on improvement and the value of particular changes.

For instance, if the run chart showed 100% follow up, there would be no need to look at SES or ethnicity affecting performance. If the run chart was obstinately at 80%, it might prompt thought about the demographic of the non-attenders.

In the absence of individual-level SES data, the best we could do is describe the area-level SES conditions where the practice is located which would not be useful for adjusting for SES. Supplementary Table 2 and Supplementary Table 3 provide a summary of outcomes by location and practice size – any analyses beyond these summaries is limited due to the sample size.

Reference

 Run chart tool. Institute for Healthcare Improvement. http://www.ihi.org/resources/Pages/Tools/RunChart.aspx

Major comment 4

The conclusion mainly stemmed from Table 2, but this table failed to give clear information, including: (a) it is better to point out which groups have statistical difference, if the difference assessed by ANOVA is significant; (b) the total number of women registered is far less than the numbers documented in text (N=481) and is also not consistent with the numbers showed in the supplementary table 2 and table 3, which are simply divided in different ways.

Response:

Thank you for this suggestion. There is an extra column added to Table 2 that highlights the pairwise differences.

The discrepancy described is between the total number of women (n=481) and the total number of practices that participated over the length of the study (n=14). The numbers presented in the Supplementary Table 2 and Supplementary Table 3 provide summary statistics of the outcome measures classified by location (rural v metro) and practice size (small, medium and large based upon the number of GPs in each practice). We considered this important to include to provide the readership with some context around these issues. Due to the small numbers, no attempt has been made to test for significance between groups and we believe the 95% Cls presented are sufficient.

Minor comments:

Some errors in text and tables need to be corrected. For example,location and size was explored descriptively (Supplementary Tables 2 and 3), rather than Supplementary Tables 1 and 2. (Line 173)

Repsonse:

Thank you for highlighting these minor errors, we have edited the text accordingly.

Reviewer 2

Concluding statements:

How did it support diabetes prevention if diabetes prevention planning was little improved? Only 1 in 10 presented for diabetes prevention planning. Even if every one of them adopted the necessary interventions, it still leaves 90% of potentially vulnerable individuals not undertaking necessary prevention strategies.

Was it perhaps effective in detecting those with T2D or those with pre-diabetes rather than preventing diabetes?

Response:

Thank you for your feedback. We agree that our abstract conclusions needed to be framed better to reflect our findings.

We have now revised it to say "GooD4Mum supported increased diabetes screening and the monitoring of high-risk women with previous GDM in general practice."

Our study was focused on delivering guideline-led diabetes prevention care, which detail regular diabetes screening and supporting women to achieve a healthy weight, diet and regular exercise through behaviour change (prevention planning) as the core activities that need to be completed to prevent diabetes. It is these activities that need to be enacted to deliver guideline-led care and what we focus on in this study. As a result, we are unable to make any conclusion as to its effectiveness in detecting women with T2DM or prediabetes versus preventing diabetes but it would be an important consideration in future research. We have edited the discussion limitations to include:

""Patient level data were not collected within GooD4Mum due to ethics approval restrictions, which limited our ability to explore the impact of factors such as age, <u>blood glucose measurement values</u>, education level or socioeconomic status as potential modifiers of engagement with the general practice and consequently the QIC activity."

We provide additional information on the diabetes prevention planning session issues to Reviewer 1 above. We have added additional qualitative data to address this concern (see above).

Typographical errors:

Line 53 Line 127

Reponse:

These have been amended, thank you.

Results:

It would be interesting to know what the pickup rate for diabetes and prediabetes was in the screened population. Is this data available? Could it be included in this publication? This might inform, for example, the reasons for the low rates of diabetes prevention consultations.

Response:

As in a Collaborative, we do not have access the actual values for the diabetes screening tests. The data collection was focused on the rate of the screening rather than the biochemical result. We do not have ethical clearance to access this information.

We have added further detail on this as a limitation to the discussion section:

"Patient level data were not collected within GooD4Mum due to ethics approval restrictions, which limited our ability to explore the impact of factors such as age, <u>blood glucose measurement values</u>, education level or socioeconomic status as potential modifiers of engagement with the general practice and consequently the QIC activity."

and additional information in the methods section:

"The standard of care provided to patients was aligned with guidelines and <u>no personal or identified</u> <u>data was shared outside the general practice.</u>"

We have additional qualitative data that was collected as part of the study from practices but not reported in this paper due to space constraints. We have decided to include a component of this data to add context to the data already included. We have detailed the exact text above in response to a Reviewer 1 query.

Discussion:

Some discussion re cost effectiveness would be interesting. Many people involved. (both practices and PHNs)- How resource intensive is this project- is it sustainable? Financial costs: Funding for the projects, and costs to the women in time and money.

Response:

We agree that this information would be useful. We have included the following additional information:

In the methods:

"Cost information was captured throughout the intervention from a QIC and intervention perspective. The cost data was collected from women and practices participating using cost diaries alongside recorded project expenses."

"The cost data were analysed using a pathway approach and only cost descriptions could be provided."

In the results:

"Total GooD4Mum intervention costs were estimated at \$AUD 52,923, comprising project coordination \$AUD 11,573, QIC Local Program Officers time cost \$AUD 1,919, GP and practice staff time cost \$AUD 14,172, materials development and production \$AUD 24,405 and website resources \$AUD 854. The average cost per practice was estimated at \$AUD 3,528 during the QIC project. However, more than one third of total costs were associated with the handbook and material development, which would not be required for future implementation. Excluding the research and development costs, it was anticipated to deliver the intervention to one general practice would cost \$AUD 2,166. Healthcare costs were collected from women with a GDM history in the participating general practices. However, the results were not representative due to a very small sample size (N=3 pre-intervention and N=10 post-intervention) and not reported as a result."

In the discussion:

"Similarly GooD4Mum represented a modest investment to improve diabetes screening and risk monitoring amongst a high-risk population. Further research using a full economic evaluation is needed to assess the value for money of this type of intervention."

Highlights

- GooD4Mum is a quality improvement study for diabetes prevention after gestational diabetes
- 15 general practices in Victoria, Australia participated in GooD4Mum collaborative study
- Diabetes screening rates doubled (30% to 60%) and 20% increase in BMI monitoring
- Improving screening and monitoring of women with previous gestational diabetes is feasible

GooD4Mum: a general practice-based quality improvement collaborative for diabetes prevention in women with previous gestational diabetes

RUNNING TITLE: Postpartum diabetes prevention in primary care

SL O'Reilly^a*, JA Dunbar^b, JD Best^c, V Versace^b, D Ford^d, D Young^e, S Shih^f, R Bills^g, W Shepherdley^g, ED Janus^{h,i}, the MAGDA Study Group.

^a Institute of Physical Activity and Nutrition, Deakin University, 221 Burwood Highway, Burwood, VIC 3125, Australia. <u>Sharleen.oreilly@deakin.edu.au</u>

^b Deakin Rural Health, School of Medicine, Faculty of Health, Deakin University, Warrnambool, VIC 3280,, Australia. <u>James.dunbar@deakin.edu.au</u>, <u>vincent.versace@deakin.edu.au</u>

^c Lee Kong Chian School of Medicine, Imperial College London and Nanyang
 Technological University, 11 Mandalay Road, 308232 Singapore. <u>jamesbest@ntu.edu.sg</u>

^d Improvement Foundation, 8/19 Grenfell St, Adelaide SA 5000, Australia.

Dale.Ford@improve.org.au

Faculty of Medicine, Dentistry and Health Sciences, Building 181, University of
 Melbourne, Grattan Street, Melbourne, VIC 3010, Australia. <u>D.young@unimelb.edu.au</u>

^fCentre for Population Health Research, Deakin University, 221 Burwood Highway, Burwood, VIC 3125, Australia. <u>sophy.shih@deakin.edu.au</u>

^g Brooke Street Medical Centre, 14 Brooke Street, Woodend VIC 3422, Australia. <u>RBills@bsmc.net.au</u>, <u>WShepherdley@bsmc.net.au</u> ^h General Internal Medicine Unit, Western Health, Sunshine Hospital, 176 Furlong Rd, St Albans, VIC 3021, Australia.

ⁱ Department of Medicine, Melbourne Medical School – Western Precinct, University of Melbourne, 176 Furlong Rd, St Albans, VIC 3021, Australia. <u>edwarddj@unimelb.edu.au</u>

* Corresponding Author: Dr Sharleen O'Reilly, UCD Institute of Food and Health,
University College Dublin, Belfield, Dublin 4, Ireland. Phone +353 1 716 2157, Fax +353
1 716 1104, <u>sharleen.oreilly@ucd.ie</u>

1	GooD4Mum: a general practice-based quality improvement collaborative for	
2	diabetes prevention in women with previous gestational diabetes	
3	RUNNING TITLE: Postpartum diabetes prevention in primary care	
4	SL O'Reilly ^{a*} , JA Dunbar ^b , JD Best ^c , V Versace ^{be} , D Ford ^{de} , D Young ^{ef} , S Shih <mark>i^b</mark> , R Bills ^g , W Shepherdley ^g ,	Formatted: Superscript
5	ED Janus ^{h,i} , the MAGDA Study Group.	
6	^a Institute of Physical Activity and Nutrition, Deakin University, 221 Burwood Highway, Burwood, VIC	
7	3125, Australia. Sharleen.oreilly@deakin.edu.au	
8	^b Deakin Rural Health, School of Medicine, Faculty of Health, Deakin University, Warrnambool, VIC	
9	3280,, Australia. James.dunbar@deakin.edu.au, vincent.versace@deakin.edu.auGentre for	
10	Population Health Research, Deakin University, 221 Burwood Highway, Burwood, VIC 3125,	
11	Australia. <u>James.dunbar@deakin.edu.au, sophy.shih@deakin.edu.au</u>	
12	^c Lee Kong Chian School of Medicine, Imperial College London and Nanyang Technological University,	
13	11 Mandalay Road, 308232 Singapore. jamesbest@ntu.edu.sg	
14	^d Improvement Foundation, 8/19 Grenfell St, Adelaide SA 5000, Australia.	
15	Dale.Ford@improve.org.au	
16	^e Faculty of Medicine, Dentistry and Health Sciences, Building 181, University of Melbourne, Grattan	Formatted: Superscript
17	<u>Street, Melbourne, VIC 3010, Australia. D.γoung@unimelb.edu.au</u>	
18	^f School of Medicine, Deakin University, Locked bag 20000, Geelong VIC 3220, Australia.	Formatted: Superscript
19	vincent.versace@deakin.edu.au Centre for Population Health Research, Deakin University, 221	
20	Burwood Highway, Burwood, VIC 3125, Australia. James.dunbar@deakin.edu.au,	
21	<u>sophy.shih@deakin.edu.au</u>	
22		

- 23 ^e-Improvement Foundation, 8/19 Grenfell St, Adelaide SA 5000, Australia.
- 24 Dale.Ford@improve.org.au
- 25 ⁴ Faculty of Medicine, Dentistry and Health Sciences, Building 181, University of Melbourne, Grattan
- 26 Street, Melbourne, VIC 3010, Australia. <u>D.young@unimelb.edu.au</u>
- 27 ^g Brooke Street Medical Centre, 14 Brooke Street, Woodend VIC 3422, Australia.
- 28 RBills@bsmc.net.au, WShepherdley@bsmc.net.au
- 29 ^h General Internal Medicine Unit, Western Health, Sunshine Hospital, 176 Furlong Rd, St Albans, VIC
- 30 3021, Australia.
- ⁱ Department of Medicine, Melbourne Medical School Western Precinct, University of Melbourne,
- 32 176 Furlong Rd, St Albans, VIC 3021, Australia. edwarddj@unimelb.edu.au
- 33 * Corresponding Author: Dr Sharleen O'Reilly, UCD Institute of Food and Health, University College
- 34 Dublin, Belfield, Dublin 4, Ireland. Phone +353 1 716 2157, Fax +353 1 716 1104,
- 35 <u>sharleen.oreilly@ucd.ie</u>
- 36
- 37

38 GooD4Mum: a general practice-based quality improvement collaborative for

39 diabetes prevention in women with previous gestational diabetes

- 40 Abstract
- 41 Aims Gestational diabetes (GDM) and Type 2 diabetes pose tremendous health and economic
- 42 burdens as worldwide incidence increases. Primary care-based systematic diabetes screening and
- 43 prevention programs could be effective in women with previous GDM. GooD4Mum aimed to
- 44 determine whether a Quality Improvement Collaborative (QIC) would improve postpartum diabetes
- 45 screening and prevention planning in women with previous GDM in general practice.
- 46 Methods Fifteen general practices within Victoria (Australia) participated in a 12-month QIC,
- 47 consisting of baseline and four quarterly audits, guideline-led workshops and Plan-Do-Study-Act
- 48 feedback cycles after each audit. The primary outcome measures were the proportion of women on
- 49 local GDM registers completing a diabetes screening test and a diabetes prevention planning
- 50 consultation within the previous 15 months.
- 51 **Results** Diabetes screening increased with rates more than doubled from 26% to 61% and
- 52 postpartum screening increased from 43% to 60%. Diabetes prevention planning consultations did
- 53 not show the same level of increase (0% to 10%). The recording of body mass index improved overall
- 54 (51% to 69%) but the number of women with normal body mass index did not.

55 Conclusions GooD4Mum supported diabetes prevention in general practice through increaseding
 56 diabetes screening and the monitoring of high risk women with previous GDM in general practice.

- 57
- 58
- 59
- Abbreviations: gestational diabetes, GDM; glycated haemoglobin A1c, HbA1c; quality improvement
 collaborative, QIC; general practitioner, GP; body mass index, BMI.
- 62

63 Highlights

- GooD4Mum is the first quality improvement collaborative study focused on increasing diabetes
 screening and diabetes prevention planning consultations in general practice for women with
 previous gestational diabetes.
- Using established collaborative methods in 15 general practices, we found diabetes screening
 rates doubled and a 20% increase in body mass index monitoring.
- Though challenging, this study suggests that improving screening activity and monitoring of high
 risk women with previous gestational diabetes in a primary care setting is feasible.
- 71 72
- 73 Word count
- 74 2827

75 Introduction

76 The prevalence of diabetes is growing worldwide [1] and a history of gestational diabetes (GDM) 77 confers increased risk of developing Type 2 diabetes [2]. The incidence of GDM in Australian women 78 is 6% [3] and is higher for some ethnic and socio-economic groups. For women who develop GDM, 79 their risk of developing Type 2 diabetes within 5-10 years is sevenfold higher than for women who 80 have not had GDM [2]. In 2011 Australia started a National Gestational Diabetes Register (Register) to help women manage their diabetes risk by providing them with information booklets and regular 81 82 screening reminders [4]. Women with GDM typically have their 6-8 week postpartum check-up with 83 their general practitioner (GP) [5, 6]; although the Register provides screening reminders around this 84 time, postpartum screening rates do not appear to be increasing and remain low - around 30% over 85 three years [7, 8]. Significant barriers exist for mothers and general practitioners (GPs) around 86 diabetes screening and lifestyle change [5, 6, 9]. The main screening barriers are time pressures, 87 losing laboratory request forms, and arranging transport and childcare [6, 10], while those for 88 lifestyle modification are apathy towards change, time pressures and mixed messages [6]. Screening 89 appears to be the main stumbling block in supporting this population to reduce their risk of diabetes 90 as regular screening will help identify those at higher risk of developing diabetes and enable their 91 engagement in effective diabetes prevention lifestyle interventions [11] earlier. 92 Quality improvement Collaboratives (QIC) is a methodology developed by the Boston Institute for 93 Healthcare Improvement that can be applied to achieve system change within an organisation or its 94 teams [12]. QICs differ from randomised controlled trials because they aim to implement existing 95 evidence, usually a clinical guideline based on systematic review [13, 14], and they are concerned 96 with external and internal validity; randomised controlled trials are primarily focused on internal 97 validity [13]. QIC has shown measurable health care improvements in specific areas, for example 98 diabetes treatment and diabetes prevention in older adults [15]. This guasi-experimental research 99 project sought to apply QIC methods to the care of women with previous GDM based on prior

Australian QIC success [15] and the fact that the woman and GP identify general practice as the desired location <u>f</u>or care delivery [6]. The aim of this project was to determine whether a QIC based in general practice would improve postpartum diabetes screening, weight monitoring and diabetes prevention planning in women with previous GDM.

104 Methods

105 Context

106 Approximately 300,000 women give birth each year in Australia and at least 17,000 are diagnosed 107 with GDM [3]. The broad adoption of the WHO diagnostic criteria has increased GDM prevalence to 108 10% [16]. General practices in Australia receive support from Primary Health Networks, which are 109 government-funded and independent organisations. Almost a quarter of all general practices have 110 participated in a QIC project [15] and the Primary Health Networks provide QIC support through QIC 111 Local Program Officers. 112 The basic QIC constituents are: 1) convening an Expert Reference Panel to define the quality 113 improvement aim and measures, and approving the handbook; 2) identifying change principles and 114 ideas to address underlying causes of the evidence-to-practice gaps; 3) developing the intervention, 115 action periods and learning workshops to support the quality improvement process; and 4) using 116 small local tests of change through Plan-Do-Study-Act cycles. This QIC project was called GooD4Mum 117 and the Expert Reference Panel consisted of diabetes experts, general practice health professionals 118 (practice nurse, GP, dietitian), QIC experts and guideline developers. Victorian Primary Health 119 Networks agreed to participate in GooD4Mum and identified general practices with QIC experience 120 from their catchment areas they felt were suitable for recruitment. There was no funding attached 121 to GooD4Mum participation. Out of the 26 general practices identified and approached, 15 122 consented to participate (rural N=3, urban N=12). The reasons for declining were insufficient

123 capacity (N=6) and lack of staff interest (N=5).

125 Intervention

126	The GooD4Mum project team consisted of: a project manager, who is an implementation science
127	trained research dietitian; general practice leads, who ranged from practice managers to GPs to
128	practice nurses; QIC Local Program Officers; and the advisory group with key stakeholder
129	representation including women with previous GDM. Each participating general practice initially
130	identified a small GooD4Mum project team (typically a doctor and another staff member) to drive
131	the project activities and nominated a lead to engage with the project manager. The Primary Health
132	Networks nominated their QIC local program officers to engage with the project manager during
133	GooD4Mum. The project manager provided each QIC Local Program Officer and general practice
134	leads with one-to-one project training prior to the project starting. GooD4Mum registered with the
135	Royal Australian College of General Practitioners for Category A Continuous Professional
136	Development, which was important for GP participation. The QIC methods were unfamiliar to three
137	practices, these practices required additional support from the project manager and QIC Local
138	Program Officer.
139	GooD4Mum was divided into four three-monthly activity periods. During each activity period,
140	general practice teams used the Model of Improvement (three improvement questions and mini
141	quality improvement cycles using the Plan-Do-Study-Act approach). A minimum of one Plan-Do-
142	Study-Act cycle report was required for each activity period. Women with a previous or current GDM
143	diagnosis were identified through a combination of practice software and manual patient record
144	searches to form local general practice GDM registers, which were audited using the quality
145	improvement measures prior to each learning workshop. QIC Local Program Officers assisted general
146	practices with conducting audits, creating and maintaining local practice registers, reinforcing
147	learning workshop messages and providing guidance on completing Plan-Do-Study-Act cycles. The

124

149	and Primary Health Networks through emailed quarterly report cards. Similarly, Plan-Do-Study-Act
150	cycles were formally collated and shared among general practice teams at six monthly intervals.
151	General practice teams and QIC Local Program Officers attended four 90-minute online learning
152	workshops (webinars), facilitated by the project manager. The webinars provided interactive
153	learning on the change principles, quality improvement process and guidelines. The audit data from
154	each practice was shared during the webinar and a core component was sharing ideas and
155	collaborative problem solving. Webinars had a prescribed format (welcome, learning outcomes
156	outlined, reflection and discussion of audit data, learning topic with guest presenter/s, sharing ideas,
157	question time and review of learning outcomes, reminders) and the topics were progressive
158	(webinar one: creating and cleaning a local GDM register, webinar two: the practicalities of
159	postpartum screening, webinar three: lifestyle modification for diabetes prevention, webinar four:
160	sharing success through case studies). Each webinar recording was made accessible via the project
161	website for all participants; the website also hosted a discussion board, non-identified Plan-Do-
162	Study-Act reports and quarterly newsletters.
163	Study of Intervention
164	The approach chosen for assessing the impact of the intervention was assessing the guidelines and
165	determining what objective actions would reflect them being put into practice. For women with
166	previous GDM, this was engaging in annual diabetes screening and having a consultation where their

determining what objective actions would reflect them being put into practice. For women with previous GDM, this was engaging in <u>annual</u> diabetes screening and having a consultation where their <u>lifestyle-related modifiable</u> diabetes risk wasere assessed. For diabetes screening, the issue of whether the test type (arduous oral glucose tolerance test versus quicker fasting blood glucose or Haemoglobin A1c (HbA1c) influenced observed outcomes was questioned but the Expert Reference Panel deemed any change in screening activity would be sufficient evidence that the intervention was driving the behaviour because of the previously low level of engagement of women in screening over time [7, 8]. Also any changes in the first 3 months postpartum diabetes screening were specific to changes in oral glucose tolerance testing, which would differentiate the effect of different

174	diabetes screeing tests. For the diabetes prevention consultation, a specific project form was
175	required to be printed for each womaen and this enabled general practices to differentiate
176	intervention consultations with from standard ones. Cost information was captured throughout the
177	intervention from a QIC and intervention perspective. The cost data was collected from women and
178	practices participating using cost diaries alongside recorded project expenses. Qualitative interviews
179	and focus groups were conducted in each practice upon completion of the QIC activity to explore
180	barriers and enablers to the intervention.
181	Measures
182	The outcome measures were decided by the Expert Reference Panel based on guidelines [17],
183	previously used QIC diabetes prevention measures and measures that were readily extractable from
184	clinical software within a busy clinical setting. All data were aggregated at the practice level and non-
185	identifiable. The general practice lead conducted the manual data extraction every three months.
186	Audit data were manually checked against patient records to ensure counts were accurate and
187	complete. The primary outcome measures were the proportions of women on individual general
188	practice audits: 1) who completed a diabetes-screening test; and 2) who engaged in a diabetes
189	prevention planning consultation within the previous 15 months. The 15-month timeframe was
190	chosen to allow for local variation in appointment scheduling and return of screening results.
191	Additional secondary outcome measures included oral glucose tolerance test screening rates by
192	three months postpartum and distribution of normal body mass index (BMI) within the practice
193	audit. BMI measurement was identified as a critical measure to identify high-risk women within the
194	register. The change in measures were calculated as average percentage change over time.
195	Analysis

196 Run charts were used to report the results of changes in measures over the 12 month intervention197 (Supplementary Table 1 details each measure). Repeated-measure ANOVA was used to determine if

198	measures differed significantly between audits. A Greenhouse-Geisser correction was applied where
199	the assumption of sphericity was violated and post-hoc tests were corrected for multiple
200	comparisons using the Bonferroni method. The cost data were analysed using a pathway approach
201	and only cost descriptions could be provided. The qualitative data was analysed thematically by an
202	experienced qualitative researcher (SOR) and coded transcripts were checked by the participants for
203	accuracy of interpretation.

204 Ethical considerations

GooD4Mum had ethical approval provided by Deakin University (HEAG-H 167_2014). The project was managed and data were analysed by an external person to remove the influence of power relationships. The standard of care provided to patients was aligned with guidelines and no personal or identified data was shared outside the general practice. Each general practice consented to inclusion and there were no funding incentives provided to participate.

210 Results

211 Fifteen general practices participated and fourteen completed the project work. One practice was 212 acquired by a larger provider during the project and subsequently withdrew, they were excluded 213 from the analysis as a result. The Expert Reference Panel determined that a three-monthly audit 214 frequency was appropriate due to the relatively low prevalence of GDM in general practice 215 populations and the period being sufficient to allow women time to engage in diabetes screening, or 216 attend an appointment for a diabetes prevention planning consultation, or both. The submission of 217 audits ranged from 100% to 93% each quarter and the number of women on registers with screening 218 within three months of delivery grew from 43% to 60%. Approximately 481 women with a history of 219 GDM were involved in the GooD4Mum project. Thirty-eight Plan-Do-Study-Act cycles were reported 220 over the 12-month project and an average of three cycles were reported per general practice.

221	There was a general trend of improvement in variables measured over the duration of the project,
222	reflected in the main by shifts in screening practices and BMI monitoring (Table 1). The average
223	number of women per practice with a diagnosis of GDM was 26 (Table 2). At baseline, the average
224	level of screening occurring was 26%, rising to 61% at 12 months (P=0.002). BMI monitoring
225	increased from 51% at baseline to 69% at 12 months (P=0.003). The postpartum diabetes screening
226	and diabetes prevention action planning consultations rose over the course of the project (from 43%
227	to 60% for screening, P=0.066; from 1% to 10% for consultations, P=0.183). The impact of practice
228	location and size was explored descriptively (Supplementary Table $\frac{24}{24}$ and $\frac{32}{2}$). The average rate of
229	conversion to Type 2 Diabetes was 6% (\pm 7 SD) over the 12 months but 2 general practices have
230	missing data for this variable.
231	The qualitative data proposed reare several potential reasons barriers to conducting the diabetes
232	prevention planning for the low uptake of the consultation: 1) it was only emphasised in the final six
233	months of GooD4Mum and therefore had less time to become embedded within daily practice
234	activity; 2) organising it within normal practice workflows was challenging; 3) women were reluctant
235	to attend it for financial and time reasons; 4) limited lifestyle modification referral options existed;
236	and 5) some GPs and practice nurses lacked confidence to engage in a lifestyle modification
237	consultation
238	The planned intervention activity was influenced by several external factors (Table 1). Briefly, during
239	the project's first quarter, glycated haemoglobin (HbA1c) became a government funded (Medicare)
240	screening test for high-risk individuals and women with a history of gestational diabetes were
241	eligible. At the time, HbA1c was not present in any GDM-specific guidelines and the expert reference
242	panel recommended that HbA1c screening was not recommended for first postpartum screening
243	test but suitable thereafter. Primary Health Networks were restructured by the Australian
244	government at the halfway point of the project, impacting the capacity of Local Program Officers to
245	be involved and the project manager assumed responsibility for this activity during the final part of

246	the project. The general practice software had initial limitations extracting data for some measures,
247	but this was resolved within the first 2 quarters. The lifestyle modification program used for diabetes
248	prevention in Victoria (State-funded and run by Diabetes Australia, Victoria) had a period of funding
249	uncertainty in quarter three. During that time, GPs were unable to refer their patients into the
250	program and alternative referral plans were developed. When the program was funded again in the
251	final quarter, these alternative plans were rescinded.
252	Total GooD4Mum intervention costs were estimated at \$AUD 52,923, comprising project
253	coordination \$AUD 11,573, QIC Local Program Officers time cost \$AUD 1,919, GP and practice staff
254	time cost \$AUD 14,172, materials development and production \$AUD 24,405 and website resources
255	\$AUD 854. The average cost per practice was estimated at \$AUD 3,528 during the QIC project.
256	However, more than one third of total costs were associated with the handbook and material
257	development, which would not be required for future implementation. Excluding the research and
258	development costs, it was anticipated to deliver the intervention to one general practice would cost
259	\$AUD 2,166. Healthcare costs were collected from women with a GDM history in the participating
260	general practices. However, the results were not representative due to a very small sample size (N=3
261	pre-intervention and N=10 post-intervention) and not reported as a result.
262	Discussion
263	The GooD4Mum QIC was able to demonstrate improved diabetes screening and BMI monitoring in
264	women with previous GDM – the rate of screening doubled and a twenty percent increase in BMI
265	monitoring occurred. Diabetes prevention QICs work by screening the practice population aged over
266	40 and largely identify people aged 50-69 [15] but the average age for GDM diagnosis is 30 years
267	[18], which means these women will generally be overlooked by diabetes prevention efforts and go
268	unnoticed in general practice. The creation and regular maintenance of a local GDM register enables
269	practices to promote awareness of this growing population and embed diabetes prevention within
270	routine care.

271	Although several studies have reported the outcomes of interventions to improve postpartum
272	screening rates or lifestyle modification programs to reduce T2DM risk in women with previous GDM
273	[19-22], only two screening reminder studies have been located in general practice [20, 23] and
274	none has addressed both outcomes together. Most have limited generalisability due to being
275	conducted in a single organisation and few used a multimodal approach, which is known to be a
276	critical aspect for supporting change in health behaviour [24]. Participating GooD4Mum general
277	practices had varying levels of experience with QIC methods, had different practice sizes and were
278	located in urban and rural areas – all of which adds to the external validity of the findings.
279	QICs are multifaceted interventions that bring together many of the successful approaches identified
280	in systematic reviews for professional behaviour change (educational meetings, educational
281	outreach, local opinion leaders, audit and feedback, computerised reminders and tailored
282	interventions), which can yield changes in the order of 50% of participants [14]. Diabetes screening
283	and BMI monitoring in GooD4Mum aligns well with previous QIC diabetes prevention initiatives [15]
284	but providing a diabetes prevention planning consultation was the more challenging component of
285	the project and it had low uptake by both practice staff and women and practice staff reported
286	several barriers to engaging in the activity. In looking at the QIC activities as behaviours that need to
287	be changed and using the Theoretical Domains Framework[25] to map them, the screening activity
288	required minor environmental restructuring to ensure GPs knew which women needed screening
289	alongside some education and training with modelling promoted via teleconferences. These were
290	easier behaviours to change because staff already know they will get reimbursed for doing the blood
291	test and that HbA1c is a useful diabetes indicator. The prevention planning activity was a new
292	process for the practices and required substantial behaviour change. Within the theoretical domains
293	framework it called for: cognitive and interpersonal skills (training practice nurses to perform the
294	tasks and support women to engage in lifestyle change), belief about capability and consequences
295	(both staff and woman), environmental restructuring (needed patient and staff resources plus space
296	to conduct consultation), education and training, persuasion and enablement to influence practice

297	nurses and GPs optimism that the prevention planning consultations was worthwhile. There are
298	several potential reasons for the low uptake of the consultation: 1) it was only emphasised in the
299	final six months of GooD4Mum and therefore had less time to become embedded within daily
300	practice activity; 2) organising it within normal practice workflows was challenging; 3) women were
301	reluctant to attend it for financial and time reasons; 4) limited lifestyle modification referral options
302	existed; and 5) some GPs and practice nurses lacked confidence to engage in a lifestyle modification
303	consultation [25]. It may simply be that providing a diabetes planning consultation to all women with
304	a history of GDM is not appropriate and that providing the consultation in a more targeted fashion
305	would yield better results. This should be explored in further work. <u>Similarly GooD4Mum</u>
306	represented a modest investment to improve diabetes screening and risk monitoring amongst a
307	high-risk population. Further research using a full economic evaluation is needed to assess the value
308	for money of this type of intervention.
309	Limitations
310	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the
310	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the
310 311	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the
310 311 312	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates
310 311 312 313	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates [21]. Similarly the changes in diabetes screening could be attributed solely to the availability of
310 311 312 313 314	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates [21]. Similarly the changes in diabetes screening could be attributed solely to the availability of HbA1c as a Medicare funded item and easier test to undertake, yet the change seen in postpartum
 310 311 312 313 314 315 	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates [21]. Similarly the changes in diabetes screening could be attributed solely to the availability of HbA1c as a Medicare funded item and easier test to undertake, yet the change seen in postpartum screening was only due to increased oral glucose tolerance testing, pointing to GooD4Mum
 310 311 312 313 314 315 316 	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates [21]. Similarly the changes in diabetes screening could be attributed solely to the availability of HbA1c as a Medicare funded item and easier test to undertake, yet the change seen in postpartum screening was only due to increased oral glucose tolerance testing, pointing to GooD4Mum stimulating a increase in screening activity across the board. The 6-8 week postpartum screening
 310 311 312 313 314 315 316 317 	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates [21]. Similarly the changes in diabetes screening could be attributed solely to the availability of HbA1c as a Medicare funded item and easier test to undertake, yet the change seen in postpartum screening was only due to increased oral glucose tolerance testing, pointing to GooD4Mum stimulating a increase in screening activity across the board. The 6-8 week postpartum screening audit lacks complete data for audit one and two so a full picture cannot be seen for the whole
 310 311 312 313 314 315 316 317 318 	GooD4Mum was a small-scale, uncontrolled QIC conducted in a single State in Australia. While the changes in the measures could be attributed to epiphenomena, we know that usual care during the same timeframe was not producing change in diabetes screening [7, 8] or lifestyle modification rates [21]. Similarly the changes in diabetes screening could be attributed solely to the availability of HbA1c as a Medicare funded item and easier test to undertake, yet the change seen in postpartum screening was only due to increased oral glucose tolerance testing, pointing to GooD4Mum stimulating a increase in screening activity across the board. The 6-8 week postpartum screening audit lacks complete data for audit one and two so a full picture cannot be seen for the whole project, a clear limitation as this information would have provided a more nuanced picture of the

322	of engagement with the general practice and consequently the QIC activity. The small sample size
323	limited the level of insight that could be gained from the QIC project. Only three Plan-Do-Study-Act
324	cycles were reported on average per practice, which is low for QIC projects and a possible limitation.
325	Staff turnover was the main reason for practices missing Plan-Do-Study-Act cycles and some
326	practices wrote up several Plan-Do-Study-Act activities within a single report, which reduced the
327	number of Plan-Do-Study-Act reports they submitted. It is possible that a cluster randomised
328	controlled trial approach with more refined measures would address the majority of the limitations
329	identified.
330	Conclusions
331	This QIC project demonstrates significant improvements in type 2 diabetes screening and BMI
332	monitoring but further improvements are possible, particularly around diabetes prevention planning
333	consultations. Future practice needs to build upon the learnings of this project and ensure that a
334	systems approach is taken to improve outcomes for women with previous GDM.
334 335	systems approach is taken to improve outcomes for women with previous GDM. Funding
335	Funding
335 336	Funding This work was supported by the Greater Green Triangle University Department of Rural Health,
335 336 337	Funding This work was supported by the Greater Green Triangle University Department of Rural Health, Flinders University and Deakin University and the NHMRC Translating Research Into Practice
335 336 337 338	Funding This work was supported by the Greater Green Triangle University Department of Rural Health, Flinders University and Deakin University and the NHMRC Translating Research Into Practice Fellowship (1069254 to S.L.O'R.). The funders had no role in study design, data collection and
335 336 337 338 339	Funding This work was supported by the Greater Green Triangle University Department of Rural Health, Flinders University and Deakin University and the NHMRC Translating Research Into Practice Fellowship (1069254 to S.L.O'R.). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.
335 336 337 338 339 340	Funding This work was supported by the Greater Green Triangle University Department of Rural Health, Flinders University and Deakin University and the NHMRC Translating Research Into Practice Fellowship (1069254 to S.L.O'R.). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript. Conflict of interest
 335 336 337 338 339 340 341 	Funding This work was supported by the Greater Green Triangle University Department of Rural Health, Flinders University and Deakin University and the NHMRC Translating Research Into Practice Fellowship (1069254 to S.L.O'R.). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript. Conflict of interest The authors declare that they have no conflict of interest.

345	Paddy Phillips (South Australia Health), Alison Nankervis (The Womens Hospital), Greg Johnson
346	(Diabetes Australia), John Catford (Epworth Healthcare), Bill Jeffries (Lyell McEwin Hospital), John
347	Rasa (Networking Health Victoria), Liza Kelsall (Department of Health and Human Services Victoria),
348	Douglas Boyle (University of Melbourne), Bill Hague (University of Adelaide), Ken Sikaris (Melbourne
349	Pathology), Wendy Scheil (South Australia Health), Craig Bennett (Diabetes Australia), Peter
350	Baghurst (University of Adelaide).
351	We sincerely thank all GooD4Mum participants and organisations especially the general practices,
352	Medicare Locals and Primary Health Networks which participated in the study; Dino Asproloupos for
353	senior project management; Expert Reference Group; Dr Sue Phillips, CEO Therapeutics Guidelines;
354	and Dr Siew Lim for manuscript review. The concept of the study was contained within a National
355	Health and Medical Research Council (NHMRC) Partnership project grant (AppID: 533956) called

- 356 Mothers After Gestational Diabetes In Australia (MAGDA).
- The contents of this publication are solely the responsibility of the individual authors and do notreflect the views of the NHMRC.

359	References
360 361	[1] International Diabetes Federation, IDF Diabetes Atlas, 7th ed., International Diabetes Federation,
362	Brussels, Belgium, 2015.
363	[2] L. Bellamy, JP. Casas, A.D. Hingorani, D. Williams, Type 2 diabetes mellitus after gestational
364	diabetes: a systematic review and meta-analysis, Lancet, 373 (2009) 1773-1779.
365	[3] Australian Institute of Health and Welfare, Diabetes in pregnancy: its impact on Australian
366	women and their babies in: Diabetes Series: Issue 14, Australian Institute of Health and Welfare,,
367	Canberra, Australia, 2010.
368	[4] National Diabetes Services Scheme, National Gestational Diabetes Register, in, Diabetes
369	Australia, 2016.
370	[5] S.A. Wilkinson, S.S. Lim, S. Upham, A. Pennington, S.L. O'Reilly, D. Asproloupos, D.H. Mc Intyre,
371	J.A. Dunbar, Who's responsible for the care of women during and after a pregnancy affected by
372	gestational diabetes?, The Medical journal of Australia, 201 (2014) S78-S81.
373	[6] A. Pennington, S.L. O'Reilly, D. Young, J. Dunbar, Improving follow-up care for women with a
374	history of gestational diabetes: perspectives of GPs and patients, Aust J Prim Health, 23 (2016) 66-
375	74.
376	[7] C. Chamberlain, A. McLean, J. Oats, B. Oldenburg, S. Eades, A. Sinha, R. Wolfe, Low Rates of
377	Postpartum Glucose Screening Among Indigenous and non-Indigenous Women in Australia with
378	Gestational Diabetes, Matern Child Health J, (2014) 1-13.
379	[8] D.I.R. Boyle, V.L. Versace, J.A. Dunbar, W. Scheil, E. Janus, J.J.N. Oats, T. Skinner, S. Shih, S.
380	O'Reilly, K. Sikaris, L. Kelsall, P.A. Phillips, J.D. Best, M.S.G. on behalf of, Results of the first recorded
381	evaluation of a national gestational diabetes mellitus register: Challenges in screening, registration,
382	and follow-up for diabetes risk, PloS one, 13 (2018) e0200832.
383	[9] K.K. Nielsen, A. Kapur, P. Damm, M. de Courten, I.C. Bygbjerg, From screening to postpartum
384	follow-up - the determinants and barriers for gestational diabetes mellitus (GDM) services, a
385	systematic review, BMC Pregnancy Childbirth, 14 (2014) 41.

- 386 [10] E. Keely, H. Clark, A. Karovitch, I. Graham, Screening for type 2 diabetes following gestational
- diabetes: Family physician and patient perspectives, Canadian Family Physician, 56 (2010) 558-563.
- 388 [11] R. Ratner, C. Christophi, B. Metzger, D. Dabelea, P. Bennett, X. Pi-Sunyer, S. Fowler, S. Kahn,
- 389 Prevention of diabetes in women with a history of gestational diabetes: effects of metformin and
- 390 lifestyle interventions, J Clin Endocrinol Metab, 93 (2008) 4774 4779.
- 391 [12] Institute of Healthcare Improvement, The Breakthrough Series: IHI's Collaborative Model for
- 392 Achieving Breakthrough Improvement, in: IHI Innovation Series white paper, Institute of Healthcare
- 393 Improvement, , Boston, MA, 2003.
- 394 [13] D. Oliver, David Oliver: Should practical quality improvement have parity of esteem with
- evidence based medicine?, BMJ, 357 (2017).
- 396 [14] L.M.T. Schouten, M.E.J.L. Hulscher, J.J.E. van Everdingen, R. Huijsman, R.P.T.M. Grol, Evidence
- 397 for the impact of quality improvement collaboratives: systematic review, BMJ, 336 (2008) 1491-
- 398 1494.
- 399 [15] A.W. Knight, D. Ford, R. Audehm, S. Colagiuri, J. Best, The Australian Primary Care Collaboratives
 400 Program: improving diabetes care, BMJ Quality & Safety, 21 (2012) 956-963.
- 401 [16] R.G. Moses, I. Goluza, J.P. Borchard, A. Harman, A. Dunning, M. Milosavljevic, The prevalence of
- 402 diabetes after gestational diabetes An Australian perspective, Australian and New Zealand Journal
- 403 of Obstetrics and Gynaecology, 57 (2017) 157-161.
- 404 [17] Endocrinology Expert Group, Therapeutic Guidelines: endocrinology, Therapeutic Guidelines
- Limited, Melbourne, 2014.
- 406 [18] A.J. Lee, R.J. Hiscock, P. Wein, S.P. Walker, M. Permezel, Gestational Diabetes Mellitus: Clinical
- 407 Predictors and Long-Term Risk of Developing Type 2 Diabetes: A retrospective cohort study using
- 408 survival analysis, Diabetes Care, 30 (2007) 878-883.
- 409 [19] A. Ferrara, M.M. Hedderson, S.D. Brown, C.L. Albright, S.F. Ehrlich, A.-L. Tsai, B.J. Caan, B.
- 410 Sternfeld, N.P. Gordon, J.A. Schmittdiel, E.P. Gunderson, A.A. Mevi, W.H. Herman, J. Ching, Y. Crites,
- 411 C.P. Quesenberry, The Comparative Effectiveness of Diabetes Prevention Strategies to Reduce

- 412 Postpartum Weight Retention in Women With Gestational Diabetes Mellitus: The Gestational
- 413 Diabetes' Effects on Moms (GEM) Cluster Randomized Controlled Trial, Diabetes Care, 39 (2016) 65-
- 414 74.
- 415 [20] A.K. Shea, B.R. Shah, H.D. Clark, J. Malcolm, M. Walker, A. Karovitch, E.J. Keely, The
- 416 effectiveness of implementing a reminder system into routine clinical practice: does it increase
- 417 postpartum screening in women with gestational diabetes?, Chronic diseases in Canada, 31 (2011)
- 418 58-64.
- 419 [21] S.L. O'Reilly, J.A. Dunbar, V. Versace, E. Janus, J.D. Best, R. Carter, J.J.N. Oats, T. Skinner, M.
- 420 Ackland, P.A. Phillips, P.R. Ebeling, J. Reynolds, S.T.F. Shih, V. Hagger, M. Coates, C. Wildey, M.S.
- 421 Group, Mothers after Gestational Diabetes in Australia (MAGDA): A Randomised Controlled Trial of a
- 422 Postnatal Diabetes Prevention Program, PLOS Med, 13 (2016) e1002092.
- 423 [22] J. Guo, J.-L. Chen, R. Whittemore, E. Whitaker, Postpartum Lifestyle Interventions to Prevent
- 424 Type 2 Diabetes Among Women with History of Gestational Diabetes: A Systematic Review of
- 425 Randomized Clinical Trials, Journal of Women's Health, 25 (2015) 38-49.
- 426 [23] H.D. Clark, I.D. Graham, A. Karovitch, E.J. Keely, Do postal reminders increase postpartum
- 427 screening of diabetes mellitus in women with gestational diabetes mellitus? A randomized
- 428 controlled trial, American Journal of Obstetrics and Gynecology, 200 (2009) 634.e631-634.e637.
- 429 [24] C. Yarrington, C. Zera, Health Systems Approaches to Diabetes Screening and Prevention in
- 430 Women with a History of Gestational Diabetes, Current Diabetes Reports, 15 (2015) 1-6.
- 431 [25] J. Cane, D. O'Connor, S. Michie, Validation of the theoretical domains framework for use in
- 432 behaviour change and implementation research, Implementation Science, 7 (2012) 37.

433

 Table 1. Diabetes prevention collaborative for women with previous gestational diabetes in general

practice and its evolution over time.

Aim	• 100% women with providus GDM with	in participating general practices to have a					
A		in participating general practices to have a					
	diabetes screening test within the past	diabetes screening test within the past 15 months					
	 100% women with previous GDM with 	100% women with previous GDM within participating general practices to be					
		provided with the opportunity to receive a consultation discussing a diabetes					
		prevention action plan within the past 15 months					
Timeline	Planned activity	External influences					
Jul-Sep	Handbook preparation						
2014	 Expert Reference Panel meeting – 						
2011	change principles and measures						
	 Handbook finalised and endorsed by 						
	Expert Reference Panel, provided to						
	all general practices						
	 General practice recruitment to 						
	quality improvement collaborative						
Oct-Dec	 Audit 1 where the program manager 	Australian government announced					
2014	and Local Program Officer supported	HbA1c as a screening test under					
-017	practices to perform initial audit and	Medicare funding (Late Nov)					
	form the baseline register of women	 Handbook materials, audit support 					
	with previous GDM	and education materials updated to					
	 Learning workshop 1 occurred 	reflect change and delivered to					
	 Plan-Do-Study-Act cycle/s 	general practice					
	undertaken in general practice and	General practice audit software data					
	records provided to program	extraction coding issue, negotiation					
	manager	with several software providers to					
	manager	resolve issue					
Jan-Apr	• Audit 2 performed with continued	General practice software patch rolled					
2015	support	out to fix coding issue (Jan/Feb)					
	 Learning workshop 2 occurred 						
	Plan-Do-Study-Act cycle/s						
	undertaken in general practice and						
	records provided to program						
	manager						
May-	Audit 3 performed with continued	State-wide diabetes prevention					
Aug	support	program places freeze on new					
2015	Learning workshop 3 occurred	participants due to funding					
	Plan-Do-Study-Act cycle/s	renegotiation, practices unable to					
	undertaken in general practice and	refer women with previous GDM					
	records provided to program	American Diabetes Association and					
	manager	NICE revise guidelines to include					
		HbA1c as screening test for women					
		with previous gestational diabetes					
		Australian government restructure of					
		Primary Health Networks reduces					
		Local Program Officer capacity					
Sep-Dec	Audit 4 performed with continued	State-wide diabetes prevention					

2015		support	program refunded and new
	٠	Learning workshop 4 occurred	participants accepted
	٠	Plan-Do-Study-Act cycle/s	
		undertaken in general practice and records provided to program	
		manager	
	•	Final audit performed	

Outcome	Baseline	Audit 1	Audit 2	Audit 3	Audit 4	Significant
	Nov 2014	Feb 2015	May 2015	Aug 2015	Nov 2015	<u>comparisons</u>
Women registered (N)	26.7 (8.6, 44.8)	23.4 (6.3, 40.5)	29.6 (12.6, 46.6)	31.1 (14.5, 47.8)	34.4 (17.2, 51.5)	<u>Audit 1 v Audit 4;</u> <u>Audit 2 v Audit 4</u>
Women screened for diabetes (N)	9.1 (1.4, 16.9)	10.8 (2.3, 19.3)	14.9 (5.4, 24.4)	16.4 (6.0, 26.8)	20.0 (9.1, 30.9)	<u>Baseline v Audit 3;</u> <u>Baseline v Audit 4;</u> <u>Audit 1 v Audit 4</u>
Women screened for diabetes (%)	26.1 (11.6, 40.7)	32.9 (16.1, 49.7)	53.4 (38.9, 68.0)	54.2 (39.4, 69.0)	61.0 (48.6, 73.4)	<u>Baseline v Audit 5</u>
Women with BMI recorded (%)	50.7 (28.1, 73.3)	55.9 (34.7, 77.1)	65.6 (48.9, 82.2)	69.6 (55.1, 84.0)	68.8 (53.4, 84.2)	<u>Baseline v Audit 3;</u> Baseline v Audit 4
Women with normal BMI recorded (%)	19.1 (4.7, 33.4)	22.1 (9.4, 34.9)	31.3 (15.0, 47.6)	32.8 (16.5, 49.1)	33.2 (16.6, 49.9)	<u>No significant</u> <u>comparisons</u>
Women with diabetes prevention action planning consultation (%)	0.9 (-1.0, 2.7)	5.6 (-5.7, 16.8)	7.3 (-3.8, 18.4)	9.1 (-3.4, 21.7)	10.3 (-3.0, 23.5)	<u>No significant</u> <u>comparisons</u>
Women screened for diabetes within first three months postpartum (%)	Data incomplete	Data incomplete	42.3 (22.4, 64.2)	46.6 (23.8, 69.5)	59.9 (39.5, 80.3)	<u>No significant</u> comparisons

Table 2. Summary of means and 95% confidence intervals from the general practices completing Good4Mum (N=14).

Supplementary Table 1. GooD4Mum quality improvement collaborative measures

Measure	Description
GDM Register	The number of women within the clinical database that are coded with a diagnosis matching the GDM definition
T2DM Screening	The number of women on the GDM Register who have had an OGTT/FPG measurement recorded within the previous 15 months
T2DM diagnosis	The number of women on the GDM Register who have had a diagnosis of T2DM recorded
T2DM Prevention Care	The number of women on the GDM Register who had the GooD4Mum diabetes prevention action plan printed out
Postpartum Follow Up of Gestational Diabetes	The number of women on the GDM Register who gave birth within the previous year <u>and</u> had an OGTT measurement recorded within 3 months of delivery
Body Mass Index (BMI) - Recorded	The number of women on the GDM Register with recorded weight <u>and</u> height OR BMI
Normal Body Mass Index (BMI)	The number of women on the GDM Register where BMI is < 25

Supplementary Table 2. Summary of means and 95% confidence intervals from the general practices completing Good4Mum divided into metropolitan and rural areas using Accessibility/Remoteness Index of Australia (ARIA).

	Metro (n=11)	Rural (n=3)			
Total register (n, 95%	Total register (n, 95%CI)				
Audit 1	25.5 (4.1, 46.8)	31.3 (-57.9, 120.5)			
Audit 2	25.9 (3.6, 48.2)	14.3 (-3.6, 32.3)			
Audit 3	32.2 (10.3, 54.1)	20.0 (-10.2, 50.2)			
Audit 4	32.8 (11.2, 54.4)	25.0 (-8.42, 58.4)			
Audit 5	36.3 (14.2, 58.2)	27.3 (-10.3, 65.0)			
Diabetes screened (%	6 register, 95%CI)				
Audit 1	24.5 (8.8, 40.2)	32.3 (-58.5, 123.1)			
Audit 2	28.1 (9.1, 47.1)	50.7 (-23.5, 124.9)			
Audit 3	55.4 (38.0, 72.8)	46.3 (-19.1, 111.8)			
Audit 4	53.7 (35.2, 72.3)	56.0 (0.1, 109.9)			
Audit 5	61.0 (44.9, 77.1)	61.0 (35.8, 86.2)			
Diabetes prevention	consultation (% register, 95%CI)				
Audit 1	1.1 (-1.3, 3.5)	0 (0)			
Audit 2	7.1 (-7.6, 21.8)	0 (0)			
Audit 3	8.9 (-5.6, 23.5)	1.3 (-4.4, 7.1)			
Audit 4	10.7 (-5.7, 27.1)	3.3 (-5.4, 12.1)			
Audit 5	12.3 (-5.0, 30.0)	3.0 (-4.5, 10.5)			

Supplementary Table 3. Summary of means and 95% confidence intervals from the general practices completing Good4Mum divided into practice size, using number of effective full-time general practitioners employed at baseline.

	Small practice (n=2)	Medium practice (n=4)	Large practice (n=8)		
Total register (n, 95%CI)					
Audit 1	5.5 (-39.0, 50.0)	17.5 (-0.9, 35.9)	36.6 (4.5, 68.8)		
Audit 2	6.5 (-50.7, 63.7)	16.8 (-0.8, 34.3)	31.0 (-0.3, 62.3)		
Audit 3	7.0 (-31.1, 45.1)	22.0 (19.1, 24.9)	39.0 (8.6, 69.4)		
Audit 4	8.0 (-42.8, 58.8)	22.0 (19.1, 24.9)	41.5 (12.3, 70.7)		
Audit 5	9.5 (-60.4, 79.4)	26.5 (17.7, 35.3)	44.5 (14.6, 74.4)		
Diabetes screened (% r	egister, 95%CI)				
Audit 1	22.0 (-257.5, 301.5)	18.5 (-6.4, 43.4)	31.0 (6.3, 55.7)		
Audit 2	41.0 (-480.0, 562.0)	19.5 (7.3, 46.3)	37.6 (13.6, 61.6)		
Audit 3	75.0 (-242.7, 392.7)	48.0 (17.0, 79.0)	50.8 (28.9, 72.6)		
Audit 4	66.5 (-143.2, 276.2)	36.0 (-5.1, 77.1)	60.3 (40.1, 80.5)		
Audit 5	68.5 (-166.6, 303.6)	46.8 (12.0, 81.6)	66.3 (49.9, 82.7)		
Diabetes prevention consultation (% register, 95%CI)					
Audit 1	0 (0)	3.0 (-6.6, 12.6)	0 (0)		
Audit 2	36.5 (-427.3, 500.3)	1.3 (-2.7, 5.2)	0 (0)		
Audit 3	36.5 (-427.3, 500.3)	3.5 (-3.4, 10.4)	1.9 (-1.4, 5.2)		
Audit 4	41.5 (-485.8, 568.8)	3.5 (-3.4, 10.4)	3.9 (-0.1, 7.9)		
Audit 5	43.5 (-509.2, 596.2)	3.5 (-3.4, 10.4)	5.4 (-1.0 11.8)		

SQUIRE 2.0 Rep	orting Checklist	
Section	Description	Location
Title	Indicate that the manuscript concerns an initiative to improve	1-2
	healthcare (broadly defined to include the quality, safety,	
	effectiveness, patientcenteredness, timeliness, cost, efficiency, and	
	equity of healthcare)	
Abstract	a. Provide adequate information to aid in searching and indexing	31-47
	b. Summarize all key information from various sections of the text	
	using the abstract format of the intended publication or a structured	
	summary such as: background, local problem, methods,	
	interventions, results, conclusions	
Introduction	Why did you start?	
Problem	Nature and significance of local problem	64-68
description		70.70
Available	Summary of what is currently known about the problem, including	70-79
knowledge	relevant previous studies	00.00
Rationale	Informal or formal frameworks, models, concepts, and/or theories	80-88
	used to explain the problem, any reasons or assumptions that were used to develop the intervention(s), and reasons why the	
	intervention(s) was expected to work	
Specific aims	Purpose of the project and of this report	89-90
Methods	What did you do?	05 50
Context	Contextual elements considered important at the outset of	93-110
Context	introducing the intervention(s)	55 110
Intervention	a. Description of the intervention(s) in sufficient detail that others	113-149
	could reproduce it	110 110
	b. Specifics of the team involved in the work	
Study of	a. Approach chosen for assessing the impact of the intervention(s)	151-162
intervention	b. Approach used to establish whether the observed outcomes were	
	due to the intervention(s)	
Measures	a. Measures chosen for studying processes and outcomes of the	164-175
	intervention(s), including rationale for choosing them, their	
	operational definitions, and their validity and reliability	
	b. Description of the approach to the ongoing assessment of	
	contextual elements that contributed to the success, failure,	
	efficiency, and cost	
	c. Methods employed for assessing completeness and accuracy of	
Australia	data	477 400
Analysis	a. Qualitative and quantitative methods used to draw inferences from the data	177-182
	b. Methods for understanding variation within the data, including	
	the effects of time as a variable	
Ethical	Ethical aspects of implementing and studying the intervention(s) and	184-188
considerations	how they were addressed, including, but not limited to, formal	104 100
	ethics review and potential conflict(s) of interest	
Results	What did you find?	
Results	a. Initial steps of the intervention(s) and their evolution over time	190-221
	(e.g., time-line diagram, flow chart, or table), including modifications	
	made to the intervention during the project	
	b. Details of the process measures and outcome	
	c. Contextual elements that interacted with the intervention(s)	

	d. Observed associations between outcomes, interventions, and	
	relevant contextual elements	
	e. Unintended consequences such as unexpected benefits,	
	problems, failures, or costs associated with the intervention(s).	
	f. Details about missing data	
Discussion	What does it mean?	
Summary	a. Key findings, including relevance to the rationale and specific aims b. Particular strengths of the project	223-231
Interpretation	a. Nature of the association between the intervention(s) and the	232-254
	outcomes	
	b. Comparison of results with findings from other publications	
	c. Impact of the project on people and systems	
	d. Reasons for any differences between observed and anticipated	
	outcomes, including the influence of context	
	e. Costs and strategic trade-offs, including opportunity costs	
Limitations	a. Limits to the generalizability of the work b. Factors that might	256-273
	have limited internal validity such as confounding, bias, or	
	imprecision in the design, methods, measurement, or analysis	
	c. Efforts made to minimize and adjust for limitations	
Conclusions	a. Usefulness of the work	275-281
	b. Sustainability	
	c. Potential for spread to other contexts	
	d. Implications for practice and for further study in the field	
	e. Suggested next steps	
Funding	Sources of funding that supported this work. Role, if any, of the	283-286
	funding organization in the design, implementation, interpretation,	
	and reporting	

Data Statement Click here to download Data Statement: dataprofile.xml