

Broad, Rebecca (2010) Infection prevention and control – quantitative study. [Dissertation (University of Nottingham only)] (Unpublished)

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Direct line/e-mail +44 (0) 115 8231063

Louise.Sabir@nottingham.ac.uk

30th September 2009

Ms Jacqueline Randle
Adult Branch Co-ordinator of MNSci programme
B Floor, School of Nursing
QMC Campus
Nottingham University Hospitals
NG7 2UH



Medical School Research Ethics Committee Division of Therapeutics & Molecular Medicine D Floor, South Block Queen's Medical Centre Nottingham NG7 2UH

Tel: +44 (0) 115 8231063 Fax: +44 (0) 115 8231059

Dear Ms Randle

Ethics Reference No: D/7/2009 - Please quote this number on all correspondence

Study Title: Hand Hygiene Practices: perceptions of Nurses and Health care workers working in private Nursing homes.

Lead Investigator: Ms Jacqueline Randle, Adult Branch Co-ordinator of MNSci

programme

Co Investigators: Rebecca Broad, Student Nurse, School of Nursing.

Thank you for your letter dated 18th September 2009 responding to the issues raised by the Committee and enclosing revised version of:

- Application form dated 9/18/2009
- Volunteer Consent Form dated 18/09/09

These have been reviewed and are satisfactory and the study is approved.

These have been reviewed and are satisfactory and the study amendments are approved.

Approval is given on the understanding that the Conditions of Approval set out below are followed.

Conditions of Approval

You must follow the protocol agreed and any changes to the protocol will require prior Ethic's Committee approval.

This study is approved for the period of active recruitment requested. The Committee also provides a further 5 year approval for any necessary work to be performed on the study which may arise in the process of publication and peer review.

You promptly inform the Chairman of the Ethic's Committee of

(i) Deviations from or changes to the protocol which are made to eliminate immediate hazards to the research subjects.

- (ii) Any changes that increase the risk to subjects and/or affect significantly the conduct of the research.
- (iii) All adverse drug reactions that are both serious and unexpected.
- (iv) New information that may affect adversely the safety of the subjects or the conduct of the study.
- (v) The attached End of Project Progress Report is completed and returned when the study has finished.

Yours sincerely

Professor R C Spiller

Chairman, Nottingham University Medical School Research Ethics

Committee



IMPORTANT

Medical Research Ethics Committee End of Project Progress Report

	•	ct Title: s Ref No:
<u>Se</u>	<u>ctio</u>	<u>n A</u>
-		project has been delayed, terminated or will not start please complete question 1 only. If roject has finished, complete question 2 onwards
1.	If y	rour project has not started please complete the most appropriate option below: Is planned to start on the following date:/
		Project terminated due to:
		Other:
2.		pject completion date:/ (dd/mmm/yyyy) - i.e. date last piece research data was collected
3.	Ple a.	ease enter final subject/sample numbers collected or recruited: Volunteers screened:

4.		ne project has ended, are you intending to destroy any human tissue obtained as a result he project? Yes/No		
	dep	o, please note you will be required by law to inform the person designate in your partment to ensure that you comply with standard operating procedures in your partment relating to the Human Tissue Act.		
5.	Ple	ase list your main findings from the study:		
6.	Ple	ase identify address where study will be archived:		
7.	Ple	Please state who is the custodian of the data:		
	a.	Name:		
	b.	Post Held:		
	C.	Address and Department:		
	d.	Contact details:		

Please note that is the responsibility of the Chief/Principal Investigator to ensure that all study data is archived in a secure location with restricted access to authorised personnel only and that data is held in accordance with the Data Protection Act 1998 and where appropriate with the Clinical Trials Regulations 2004. It is not good practice to store data at a personnel address or on a personal computer; it is recommended that all data be stored in a secure environment within an organisational establishment under the care of an appointed custodian.

I confirm that the information contained in this report is correct

Chief/Principal Investigator: (Signature)

Date:

Print Name:

Please return to:

Louise Sabir
UoN Medical Research Ethics Committee Secretary
c/o Division of Therapeutics & MM
D Floor, South Block
QMC Campus
Nottingham University Hospitals
Nottingham
NG7 2UH



Medical School Research Ethics Committee Membership 2008/ 2009

Chairman Professor R C Spiller, Consultant Gastroenterologist & Professor

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Dr Liz Simpson, Chief Experimental Officer.

Molecular Medical

Sciences

Dr David Turner, Senior Clinical Associate Professor in

Microbiology

Community Health

Sciences

Professor Chris Bradshaw, Consultant Psychiatrist & Professor of

Psychopharmacology.

Clinical Sciences Dr Abdol Nateri, Lecturer, Pre-Clinical Cancer Studies

Division of GI Surgery

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Nottingham.

Primary Care Dr Richard Knox, General Practitioner/ Part-time Lecturer

Division of Primary Care, QMC Campus

School of Nursing, Midwifery and

Physiotherapy

Ms Stacy Johnson, Senior Lecturer in Nursing

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School of Law, University of Nottingham.

Medical Students

nominated by ISC

Anish Sharma, 3rd Year Medical Student James Lingard, 3rd Year Medical Student

Postgradutate Student

Member

Ms Jodie Finlayson-Burden, PhD student, Division of Psychiatry

Administrator Mrs Louise Sabir, Division of Therapeutics & MM, School of

Clinical Sciences