What is the cost-effectiveness of endoscopy undertaken by nurses? A Multi-Institution NUrse Endoscopy Trial (MINUET)

Submission date 25/04/2003	Recruitment status No longer recruiting	Prospectively registered		
23/04/2003		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
25/04/2003	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
13/02/2009	Digestive System			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers HTA 97/37/09

Study information

Scientific Title

Acronym

MINUET

Study objectives

The project will evaluate the acceptability, effectiveness, outcome and cost of upper and lower gastrointestinal endoscopy undertaken by nurses in hospital. We plan to compare:

- 1. The acceptability to patients of these procedures when undertaken by nurses or doctors
- 2. The quality of the process of these procedures when undertaken by nurses or doctors
- 3. The outcome for, and value to, patients of these procedures when undertaken by nurses or doctors
- 4. The resources consumed by the NHS and patients through these procedures when undertaken by nurses or doctors.

We also plan to develop an economic model to predict the effect of nurse endoscopies on the labour market and training requirements for clinical nurse specialists.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Digestive system diseases

Interventions

Endoscopy undertaken by nurses versus doctors

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Acceptability and anxiety scores, completeness of examination, incidence of technical complications, recording of clinical findings, and need for further investigation or procedure.

2. Incidence of clinical complications, quality of life (generic and disease-specific) measured preprocedure and at one and 12 months post-procedure.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2001

Completion date

31/10/2004

Eligibility

Key inclusion criteria

Patients undergoing upper and lower gastrointestinal (GI) endoscopy.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

1888

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/09/2001

Date of final enrolment

31/10/2004

Locations

Countries of recruitment

United Kingdom

Wales

Study participating centre University of Wales Swansea Swansea United Kingdom SA2 8PP

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Quarry House Quarry Hill Leeds United Kingdom LS2 7UE +44 (0)1132 545 843 Sheila.Greener@doh.gsi.gov.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/en/index.htm

ROR

https://ror.org/03sbpja79

Funder(s)

Funder type

Government

Funder Name

NIHR Health Technology Assessment Programme - HTA (UK)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Other publications	HTA monograph	01/10/2006		Yes	No
Results article	clinical effectiveness results	10/02/2009		Yes	No
Results article	cost efectiveness results	10/02/2009		Yes	No