ISRCTN82765705 https://doi.org/10.1186/ISRCTN82765705

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 [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 25/04/2003	**Overall study status** Completed	
Last Edited 13/02/2009	**Condition category** Digestive System	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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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 pre-procedure 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
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Sponsor information

Organisation
Department of Health (UK)

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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