Walk in nasal endoscopy (WINES) study: a pilot evaluation of the safety and feasibility, and cost savings of introducing a radically new approach to upper gastrointestinal (GI) endoscopy

Submission date	Recruitment status	Prospectively registered
30/09/2004	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2004	Completed	Results
Last Edited	Condition category	[] Individual participant data
05/12/2014	Cancer	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number N0077132412

Study information

Scientific Title

Walk in nasal endoscopy (WINES) study: a pilot evaluation of the safety and feasibility, and cost savings of introducing a radically new approach to upper gastrointestinal (GI) endoscopy

Study objectives

Evaluation of safety and feasibility of a walk in two week cancer waiting list upper GI endoscopy using nasal endoscope and only one assistant. A single consultant ATC doing the procedures.

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

Study type(s)

Diagnostic

Health condition(s) or problem(s) studied

Cancer: Gastrointestinal

Interventions

Nasal endoscopy is currently performed in the endoscopy unit at Derby City General Hospital but in a procedure analogous to conventional endoscopy using two nurse assistants. The study will be a pilot randomised controlled study comparing conventional with nasal (ultraslim endoscopy) to assess primarily the safety and feasibility of performing nasal endoscopy with just one assistant and the patient in a seated position.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Primary endpoints will be the proportion of cases in which this procedure could be performed and the number of early to late (within 1 week) complications. Assessment of safety will be by pro forma records of complications during endoscopy and prior to discharge and by using a patient questionnaire administered 1 week later.

The endoscopist will assess the feasibility of the nasal procedure. Completeness of endoscopy and visualisation of the duodenum, antrum, body and fundus of the stomach and oesophagus will be recorded.

Key secondary outcome(s))

Secondary endpoints will be a questionnaire of the patients acceptability of the procedure and a preliminary assessment of the savings in staff costs of this procedure.

Completion date

01/12/2004

Eligibility

Key inclusion criteria

Patients attending for endoscopy through two-week cancer waiting list.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

21/10/2003

Date of final enrolment

01/12/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre
Southern Derbyshire Acute Hospitals NHS Trust
Derby

United Kingdom DE22 3NE

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

Southern Derbyshire Acute Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration