ISRCTN95850244 https://doi.org/10.1186/ISRCTN95850244

Inoperable oesophageal cancer - stent or percutaneous endoscopically placed gastrostomy (PEG). Randomised trial to compare the risks and benefits of expanding metallic stent and PEG in inoperable oesophageal cancer

Submission date 12/09/2003	**Recruitment status** Stopped	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Stopped	
Last Edited 22/10/2012	**Condition category** Cancer	

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
N0084096607

Study information

Scientific Title

Study objectives
1. Does the early placement of a PEG feeding tube prolong life and/or improve quality of life?
2. Does PEG feeding offer advantages over stent placement?

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Cancer: Oesophageal

Interventions
1. Oesophageal stent
2. PEG

Please note that this trial was stopped due to participant recruitment issues.

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
Survival and quality of life.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/2000

Completion date
30/06/2003

Reason abandoned (if study stopped)
Participant recruitment issue

Eligibility

Key inclusion criteria
20 patients for each study arm.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
20 patients for each study arm.

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/10/2000

Date of final enrolment
30/06/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Hull Royal Infirmary
Hull
United Kingdom
HU3 2JZ

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
The North and South Bank Research and Development Consortium (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