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Randomised study of pre-operative radio-chemotherapy versus surgery alone in thoracic oesophageal cancer deemed to be resectable

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 28/03/2017	**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

N0188116113

Study information

Scientific Title
Randomised study of pre-operative radio-chemotherapy versus surgery alone in thoracic oesophageal cancer deemed to be resectable

Study objectives
Attempt to prove that preoperative radio-chemotherapy will improve disease-free survival for this patient group.

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
Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied
Cancer: Thoracic oesophageal

Interventions
Arm A: radio-chemotherapy followed by surgery
Arm B: surgery alone

Intervention Type
Mixed

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/2002

Completion date
01/03/2004

Eligibility

Key inclusion criteria
Patients with thoracic oesophageal cancer deemed to be resectable

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration. Added July 2008: No UK participants were recruited in this EORTC trial.

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/06/2002

Date of final enrolment
01/03/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Royal Preston Hospital
Preston
United Kingdom
PR2 9HT

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
Lancashire Teaching Hospitals NHS Trust (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