

Endoscopic ablation of Barrett's oesophagus: a randomised controlled trial comparing photodynamic therapy (PDT) versus argon beam plasma coagulation (ABPC)

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 13/07/2009	Condition category Digestive System	<input type="checkbox"/> Individual participant data

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

Study information

Scientific Title

Study objectives

To compare argon beam plasma coagulation (ABPC) and photodynamic therapy (PDT) in the ablation of metaplastic and low grade dysplastic Barrett's oesophagus, in a randomised controlled prospective clinical trial.

Ethics approval required

Old ethics approval format

Ethics approval(s)

South Sheffield research ethics committee (REC) approval received (ref: 00/031)

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

Barrett's oesophagus

Interventions

Compare ABPC and PDT in the ablation of metaplastic and low grade dysplastic Barrett's oesophagus.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Tolerability of treatment

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/2000

Completion date

01/11/2004

Eligibility

Key inclusion criteria

1. Currently in an endoscopic screening programme
2. Aged 18 years or older, either sex
3. Biopsy proven Barretts epithelium with a median length of 4 cm

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

56

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/04/2000

Date of final enrolment

01/11/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

University of Sheffield
Sheffield
United Kingdom
S10 2JF

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

Sheffield Teaching Hospitals NHS Foundation Trust (UK) - Central Campus

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?
Results article	results	01/12/2004		Yes	No