

A Randomised Trial Comparing External Beam Radiotherapy Alone With External Beam Radiotherapy Plus Intraluminal Irradiation for Palliation in Lung Cancer

Submission date 19/08/2002	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 19/08/2002	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 08/04/2015	Condition category Cancer	☐ Individual participant data ☐ 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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MRC Clinical Trials Unit
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Additional identifiers

Protocol serial number

ILT-2

Study information

Scientific Title

A Randomised Trial Comparing External Beam Radiotherapy Alone With External Beam Radiotherapy Plus Intraluminal Irradiation for Palliation in Lung Cancer

Study objectives

Not provided at time of registration

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Lung (non-small cell) cancer

Interventions

1. Schedule A: External beam radiotherapy, a maximum subcutaneous dose of 3250 cGy given in eight fractions over 10 days with a margin of 2 cm around the tumour.

2. Schedule B: External beam radiotherapy, a maximum subcutaneous dose of 3250 cGy given in eight fractions over 10 days with a margin of 2 cm around the tumour. Followed by intraluminal radiotherapy, a single dose of 1500 cGy to be given on the last day of external beam radiotherapy.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/12/2005

Eligibility

Key inclusion criteria

1. Biopsy proven non-small cell lung cancer
2. Fit to receive bronchoscopy
3. Fit to receive external beam radiotherapy
4. Any age
5. Shortness of breath, cough, haemoptysis, dysphagia or chest pain resulting from a primary bronchial tumour
6. No previous or concomitant malignancy, except basal cell carcinomas
7. No symptomatic metastases requiring external beam radiotherapy
8. No previous chest irradiation for lung cancer
9. No bilateral lung tumours or tracheal tumours

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2000

Date of final enrolment

31/12/2005

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

MRC Clinical Trials Unit

London

United Kingdom

NW1 2DA

Sponsor information

Organisation

UK Co-ordinating Committee for Cancer Research (UKCCCR)

ROR

<https://ror.org/054225q67>

Funder(s)**Funder type**

Research organisation

Funder Name

Cancer organisations (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration