

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	<input type="checkbox"/> Prospectively registered
Registration date 19/08/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 08/04/2015	Condition category Cancer	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Contact details

UKCCCR Register Co-ordinator
MRC Clinical Trials Unit
222 Euston Road
London
United Kingdom
NW1 2DA

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

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

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 measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2000

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

Age group

Adult

Sex

Both

Target number of participants

Not provided at time of registration

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

England

United Kingdom

Study participating centre

MRC Clinical Trials Unit

London

United Kingdom

NW1 2DA

Sponsor information

Organisation

UK Co-ordinating Committee for Cancer Research (UKCCCR)

Sponsor details

MRC Clinical Trials Unit

222 Euston Road

London

United Kingdom

NW1 2DA

Sponsor type

Government

ROR

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

Funder(s)

Funder type

Research organisation

Funder Name

Cancer organisations (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