

Post-Operative Radiotherapy for Non-Small Cell Lung Cancer

Submission date 19/08/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 19/08/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 15/11/2019	Condition category Cancer	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

Protocol serial number
LU11

Study information

Scientific Title
Post-Operative Radiotherapy for Non-Small Cell 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

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Lung (non-small cell) cancer

Interventions

Patients are randomised two to four weeks postoperatively to either:

1. Schedule R: Radiotherapy 40 Gy given in daily fractions five days a week over three weeks to mediastinum starting four to six weeks post-operatively.
2. Schedule NoR: No further specific treatment including post-operative radiotherapy unless or until the disease recurrence.

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

31/12/2002

Eligibility**Key inclusion criteria**

1. Either sex, aged 75 years or less
2. World Health Organisation (WHO) performance status zero to two
3. Lung and cardiac function adequate for proposed resection
4. pT1 pN1 M0, pT2 pN1 M0, pT1 pN2 M0 or pT2 pN2 M0 non-small cell lung cancer
5. Complete resection of non-small cell lung cancer

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Total final enrolment

308

Key exclusion criteria

1. Previous specific anti-cancer treatment for current disease
2. Presence of other malignant disease, except basal cell carcinoma or in situ carcinoma of the cervix
3. Presence of other serious condition contraindicating surgery or radiotherapy

Date of first enrolment

01/01/2002

Date of final enrolment

31/12/2002

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

Sponsor information

Organisation

Medical Research Council (MRC) (UK)

Funder(s)

Funder type

Research council

Funder Name

Medical Research Council (MRC) (UK)

Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory Council, MRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

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	01/08/1996	15/11/2019	Yes	No