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Post-Operative Radiotherapy for Non-Small Cell Lung Cancer

Submission date	Recruitment status	[_] Prospectively registered
19/08/2002	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
19/08/2002	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
15/11/2019	Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr - -

Contact details
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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

Primary study design
Interventional

Study design
Randomised controlled trial

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