Supportive treatment in non-small cell lung cancer

Submission date	Recruitment status	Prospectively registered
28/02/2001	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
28/02/2001	Completed	Results
Last Edited	Condition category	Individual participant data
19/02/2018	Cancer	[] 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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

LU17

Study information

Scientific Title

Supportive treatment in non-small cell lung cancer

Study objectives

To compare supportive treatment with or without immediate thoracic radiotherapy in the prevention & palliation of symptoms in patients with inoperable non small cell lung cancer.

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)

Quality of life

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied

Non small cell lung cancer

Interventions

- 1. One group receives immediate thoracic radiotherapy
- 2. The other will have radiotherapy delayed until clinically indicated

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1. Palliation
- 2. Quality of life

- 3. Other
- 4. Survival time

Secondary outcome measures

Not provided at time of registration

Overall study start date

09/10/1992

Completion date

31/05/1999

Eligibility

Key inclusion criteria

- 1. No compelling indication for immediate thoracic radiotherapy
- 2. No previous treatment for the current lung cancer, either sex, and age

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

230 - all patients followed up until death

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

09/10/1992

Date of final enrolment

31/05/1999

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

MRC Clinical Trials Unit

London United Kingdom NW1 2DA

Sponsor information

Organisation

Medical Research Council (MRC) (UK)

Sponsor details

20 Park Crescent London United Kingdom W1B 1AL +44 (0)20 7636 5422 clinical.trial@headoffice.mrc.ac.uk

Sponsor type

Research council

Website

http://www.mrc.ac.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, MRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration