ISRCTN96733516 https://doi.org/10.1186/ISRCTN96733516

Supportive treatment in non-small cell lung cancer

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

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Danielle Andrews

Contact details
MRC Clinical Trials Unit
222 Euston Road
London
United Kingdom
NW1 2DA
+44 (0)20 7636 5422
clinical.trial@headoffice.mrc.ac.uk

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 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