

# Randomised trial of neo-adjuvant therapy in patients with stage IIIA non small cell lung cancer

<b>Submission date</b> 28/02/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 28/02/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 20/02/2020	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

**Study website**  
<https://www.ctu.mrc.ac.uk/studies/all-studies/l/lu20/>

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

## Secondary identifying numbers

LU20

# Study information

## Scientific Title

-

## Study objectives

To compare in patients with stage IIIA non small cell lung cancer a policy of standard therapy with thoracic radiotherapy versus a policy of chemotherapy followed, if feasible, by surgery

## 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

Cancer

## Interventions

1. One group receives standard therapy with thoracic radiotherapy
2. The other group receives chemotherapy followed, if feasible, by surgery

## Intervention Type

Other

## Phase

Phase III

## Primary outcome measure

1. Survival time
2. Quality of life

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/09/1995

**Completion date**

01/05/1999

## Eligibility

**Key inclusion criteria**

1. No previous treatment for the current lung cancer
2. Either sex
3. World Health Organisation (WHO) 0-2, T3N1M0 or T1-3N2M0 stage

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Both

**Target number of participants**

350

**Total final enrolment**

48

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/09/1995

**Date of final enrolment**

01/05/1999

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**  
**MRC Clinical Trials Unit**  
London  
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NW1 2DA

## **Sponsor information**

### **Organisation**

Medical Research Council (MRC) (UK)

### **Sponsor details**

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+44 (0)20 7636 5422  
[clinical.trial@headoffice.mrc.ac.uk](mailto: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

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/09/2005	20/02/2020	Yes	No