

# A randomised study of Mitomycin, Vinblastine and Platinum (cisplatin) (MVP) three versus six cycles in advanced Non-Small Cell Lung Cancer (NSCLC)

<b>Submission date</b> 19/08/2002	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 19/08/2002	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 16/04/2019	<b>Condition category</b> 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

Dr - -

### Contact details

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MRC Clinical Trials Unit  
222 Euston Road  
London  
United Kingdom  
NW1 2DA

## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

NCT00004209

## Secondary identifying numbers

95.24

# Study information

## Scientific Title

A randomised study of Mitomycin, Vinblastine and Platinum (cisplatin) (MVP) three versus six cycles in advanced Non-Small Cell Lung Cancer (NSCLC)

## 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

## Secondary study design

Randomised controlled trial

## Study setting(s)

Hospital

## Study type(s)

Treatment

## Participant information sheet

## Health condition(s) or problem(s) studied

Lung (non-small cell) cancer

## Interventions

Cisplatin 50 mg/m<sup>2</sup> Mitomycin-C 8 mg/m<sup>2</sup> intravenous (iv) day one  
Vinblastine 6 mg/m<sup>2</sup> iv day one

## Intervention Type

Drug

## Phase

Not Applicable

## Drug/device/biological/vaccine name(s)

Mitomycin, vinblastine and platinum (cisplatin)

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/01/1993

**Completion date**

30/11/1998

## Eligibility

**Key inclusion criteria**

1. Histological evidence of NSCLC
2. Stage IIIb or IV disease not eligible for surgery or radical radiotherapy
3. Performance status zero, one or two
4. Ability to give signed, informed consent
5. <sup>51</sup>Cr Ethylene Diamine Tetraacetic Acid (EDTA) more than 60 ml/min, creatinine clearance more than 60 ml/min, Haemoglobin (Hb) more than 10 g/dl, White Cell Count (WCC) more than  $3 \times 10^9$ /l, platelets  $100 \times 10^{12}$ /l, Liver Function Tests (LFTs) not more than two times normal unless due to metastatic disease

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

Not provided at time of registration

**Total final enrolment**

308

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/1993

**Date of final enrolment**

30/11/1998

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**MRC Clinical Trials Unit**

London

United Kingdom

NW1 2DA

## **Sponsor information**

**Organisation**

Cancer Research UK (CRUK) (UK)

**Sponsor details**

PO Box 123

Lincoln's Inn Fields

London

United Kingdom

WC2A 3PX

+44 (0)207 317 5186

kate.law@cancer.org.uk

**Sponsor type**

Charity

**Website**

<http://www.cancer.org.uk>

**ROR**

<https://ror.org/054225q67>

## **Funder(s)**

**Funder type**

Charity

**Funder Name**

Cancer Research UK

**Alternative Name(s)**

CR\_UK, Cancer Research UK - London, CRUK

**Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Other non-profit organizations

**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	01/03/2001	16/04/2019	Yes	No