

# A placebo-controlled trial of nimodipine with cyclophosphamide/cisplatin for the treatment of advanced ovarian cancer

<b>Submission date</b> 01/07/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 01/07/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 15/01/2019	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

## 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

Secondary identifying numbers

G37

# Study information

## Scientific Title

A placebo-controlled trial of nimodipine with cyclophosphamide/cisplatin for the treatment of advanced ovarian 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

## Study design

Placebo-controlled trial

## Primary study design

Interventional

## Secondary study design

Randomised controlled trial

## Study setting(s)

Hospital

## Study type(s)

Treatment

## Participant information sheet

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

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

Ovarian cancer

## Interventions

1. Arm A: Placebo tablet
2. Arm B: Nimodipine tablet

## Intervention Type

Drug

## Phase

Not Applicable

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

Nimodipine

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/01/1989

**Completion date**

01/01/1993

## Eligibility

**Key inclusion criteria**

1. Confirmed diagnosis of invasive epithelial ovarian cancer
2. International Federation of Gynaecology and Obstetrics (FIGO) stages Tc - Tv
3. 18-70 years old
4. Performance Status less than or equal to 2
5. White Blood Cells at least  $4 \times 10^9/l$
6. Platelets at least  $120 \times 10^9/l$
7. Creatinine or Ethylene Diamine Tetraacetic Acid (EDTA) clearance greater than or equal to 60 ml/min
8. Normal Bilirubin
9. Serum glutamic-oxaloacetic transaminase (SGOT)/Serum glutamic pyruvic transaminase (SGPT) less than or equal to twice normal limit
10. No previous chemotherapy or radiotherapy

**Participant type(s)**

Patient

**Age group**

Adult

**Lower age limit**

18 Years

**Upper age limit**

70 Years

**Sex**

Female

**Target number of participants**

Not provided at time of registration

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/1989

**Date of final enrolment**

01/01/1993

## **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