

International Collaborative Ovarian Neoplasm studies (2): a trial of CAP versus carboplatin in advanced ovarian cancer

Submission date 28/02/2001	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 28/02/2001	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 09/07/2014	Condition category Cancer	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
ICON2

Study information

Scientific Title

Study objectives

To compare cyclophosphamide, doxorubicin and cisplatin (CAP) with single-agent carboplatin in patients with advanced ovarian 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)

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. Cyclophosphamide, doxorubicin and cisplatin (CAP)
2. Single-agent carboplatin

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Cyclophosphamide, doxorubicin, cisplatin, carboplatin

Primary outcome measure

1. Survival time
2. Progression-free survival

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/1991

Completion date

31/12/1996

Eligibility

Key inclusion criteria

1. Chemotherapy indicated
2. No previous malignancy
3. No prior radiotherapy or chemotherapy
4. No contraindication to chemotherapy

Participant type(s)

Patient

Age group

Not Specified

Sex

Female

Target number of participants

2000 in 9 countries

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/01/1991

Date of final enrolment

31/12/1996

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
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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

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?
Results article	results	14/11/1998		Yes	No