

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

Contact name

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Contact details

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Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

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(s)

1. Survival time
2. Progression-free survival

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Female

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**

United Kingdom

England

Study participating centre

MRC Clinical Trials Unit

London

United Kingdom

NW1 2DA

Sponsor information**Organisation**

Medical Research Council (MRC) (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

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
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes