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The role of radical surgery in advanced epithelial ovarian cancer

Submission date 01/07/2001	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 01/07/2001	**Overall study status** Completed	
Last Edited 30/10/2019	**Condition category** Cancer	

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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Additional identifiers

Protocol serial number
OV3006

Study information

Scientific Title
The role of radical surgery in advanced epithelial 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
Randomised controlled trial

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Ovarian cancer

Interventions
Patients are randomised to one of two treatment arms:
1. Arm A: High activity platinum based chemotherapy. The recommended chemotherapy regimen is cyclophosphamide and cisplatin given for a maximum of eight cycles.
2. Arm B: Radical debulking surgery followed by chemotherapy as described above.

Intervention Type
Mixed

Primary outcome(s)
Not provided at time of registration

Key secondary outcome(s))
Not provided at time of registration

Completion date
31/12/1995

Eligibility

Key inclusion criteria
1. Histologically confirmed epithelial ovarian cancer
2. Macroscopic residual disease unlikely in surgeon's opinion to be completely debulked without bowel resection or other procedures
3. No attempt to radically debulk
4. No prior chemotherapy or radiotherapy
5. No previous or concurrent malignancy other than non melanomatous skin cancer
6. No haematological or biochemical contra-indication to platinum chemotherapy
7. No contraindication to second laparotomy
8. Available to commence chemotherapy or undergo further surgery within 6 weeks of diagnostic laparotomy

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Female

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/1990

Date of final enrolment
31/12/1995

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
MRC Clinical Trials Unit
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
Cancer Research UK (CRUK) (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, Cancer Research UK (CRUK), CRUK

Funding Body Type
Private sector organisation

Funding Body Subtype
Other non-profit organizations

Location
United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary