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A comparison of treosulfan and carboplatin in patients with ovarian cancer not suitable for treatment with cisplatin

Submission date 01/07/2001	**Recruitment status** Stopped	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data [_] Record updated in last year
Registration date 01/07/2001	**Overall study status** Stopped	
Last Edited 04/01/2012	**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
G44

Study information

Scientific Title

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
1. Carboplatin: 6 x (glomerular filtration rate (GFR)+ 25) mg intravvenous every 28 days for six cycles
2. Treosulfan: 7 g/m2 intravenous every 28 days for six cycles

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
treosulfan and carboplatin

Primary outcome(s)
Not provided at time of registration

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

Completion date
15/07/1998

Reason abandoned (if study stopped)
Objectives no longer viable

Eligibility

Key inclusion criteria

1. Not considered for cisplatin ≥75 mg/m2
2. Stages Ic-IV
3. Histologically proven epithelial ovarian cancer
4. Life expectancy .3 months
5. World Health Organisation (WHO) Performance status 0-3
6. Creatinine clearance ≥20 ml/min White Blood Cell (WBC) Count ≥3.5 x 10^9/l Platelets ≥100 x 10^9/l

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/1993

Date of final enrolment
15/07/1998

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
UKCCCR Register Co-ordinator
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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/01/2006		Yes	No