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Study of chemotherapy for patients with good prognosis metastatic non-seminomatous testicular cancer

Submission date 28/02/2001	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 28/02/2001	**Overall study status** Completed	
Last Edited 11/01/2019	**Condition category** Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
TE03

Study information

Scientific Title

Study of chemotherapy for patients with good prognosis metastatic non-seminomatous testicular cancer

Study objectives
Investigate whether the toxicity of chemotherapy in patients with small volume metastasis and low blood marker concentrations can be decreased by the reduction of the dose of Bleomycin given in the BEP regimen without loss of therapeutic activity.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Primary study design
Interventional

Study design
Randomised controlled trial

Study type(s)
Treatment

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

Interventions
Arm 1: Bleomycin 30 mg x 3 per course, BEP x 4
Arm 2: Bleomycin 30 mg per course, BEP x 4

Intervention Type
Drug

Phase
Phase II

Drug/device/biological/vaccine name(s)
Bleomycin, etoposide, cisplatin (platinum)

Primary outcome(s)
Toxicity/response

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

Completion date
31/12/1986

Eligibility

Key inclusion criteria
Histologically proven non-seminomatous testicular cancer

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
Male

Key exclusion criteria
Previous radiotherapy or chemotherapy

Date of first enrolment
01/01/1986

Date of final enrolment
31/12/1986

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, Medical Research Committee and Advisory 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