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Treatment of high risk testicular tumour with high dose chemotherapy with peripheral blood stem cells

Submission date	Recruitment status	[ ] Prospectively registered
12/09/2003	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
12/09/2003	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
20/04/2016	Cancer	[ ] Record updated in last year

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
N0063044257

Study information

Scientific Title
Treatment of high risk testicular tumour with high dose chemotherapy with peripheral blood stem cells

Study objectives
To assess the efficacy of high dose chemotherapy with peripheral blood stem cells to see if the response rate is improved compared to conventional therapy.

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)
Not Specified

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

Interventions
High dose chemotherapy with peripheral blood stem cells vs no high dose

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
1. Time to white cell and platelet recovery
2. Confirmation of response
3. Assessment of toxicity by standard World Health Organisation (WHO) grades

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

Completion date
31/12/2008

Eligibility

Key inclusion criteria

1. Patients with histological proven non-seminomatous germ cell tumours
2. High risk factors as defined by MRC prognostic group

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Male

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/05/1994

Date of final enrolment
31/12/2008

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Clinical Oncology
Manchester
United Kingdom
M20 4BX

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
Christie Hospital NHS Trust (UK)

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