

Study of chemotherapy for patients with good prognosis metastatic non-seminomatous testicular 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 type="checkbox"/> Results
Last Edited 11/01/2019	Condition category Cancer	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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
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

Study design

Randomised controlled trial

Primary study design

Interventional

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