ISRCTN37957513 https://doi.org/10.1186/ISRCTN37957513

MRC Study of carcinoma in situ of the contralateral testis

Submission date	Recruitment status	[_] Prospectively registered
08/02/2001	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
08/02/2001	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
18/08/2009	Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
TE11

Study information

Scientific Title

Study objectives
To ascertain the prevalence of CIS of testis in this group of patients.

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

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

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

Interventions
All subjects biopsied. Positive biopsy: optional randomisation between radiotherapy or surveillance. Negative biospy: surveillance.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Determine prevalence of CIS and the development of invasive tumour.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/1989

Completion date
31/12/1995

Eligibility

Key inclusion criteria
1. Germ cell tumour of testis or history of one having been excised in the previous five years
2. History of maldescent
3. Volume of 12 ml or less in remaining testis

Participant type(s)
Patient

Age group
Not Specified

Sex
Male

Target number of participants
208

Key exclusion criteria
Previous chemotherapy

Date of first enrolment
01/01/1989

Date of final enrolment
31/12/1995

Locations

Countries of recruitment
England

United Kingdom

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

Sponsor information

Organisation
Medical Research Council (MRC) (UK)

Sponsor details
20 Park Crescent
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W1B 1AL
+44 (0)20 7636 5422
clinical.trial@headoffice.mrc.ac.uk

Sponsor type
Research council

Website
http://www.mrc.ac.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, MRC

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
United Kingdom

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

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?
Results article	results	01/10/1998		Yes	No