

# MRC Study of carcinoma in situ of the contralateral testis

<b>Submission date</b> 08/02/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 08/02/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 18/08/2009	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data

**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**  
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

**Study type(s)**

Treatment

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

Cancer

**Interventions**

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

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Determine prevalence of CIS and the development of invasive tumour.

**Key secondary outcome(s)**

Not provided at time of registration

**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

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Male

**Key exclusion criteria**

Previous chemotherapy

**Date of first enrolment**

01/01/1989

**Date of final enrolment**

31/12/1995

**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, 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

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/10/1998		Yes	No