# A randomised trial of total prostatectomy versus radiotherapy versus no immediate treatment in early prostate cancer

Submission date 06/04/2000	<b>Recruitment status</b> Stopped	<ul><li>Prospectively registered</li><li>Protocol</li></ul>		
Registration date	Overall study status	Statistical analysis plan		
06/04/2000	Stopped  Condition category	Results		
Last Edited		Individual participant data		
19/09/2017	Cancer	Record updated in last year		

# Plain English summary of protocol

Not provided at time of registration

# Contact information

#### Type(s)

Scientific

#### Contact name

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#### Contact details

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## Additional identifiers

Protocol serial number

**PR06** 

# Study information

#### Scientific Title

A randomised trial of total prostatectomy versus radiotherapy versus no immediate treatment in early prostate cancer

#### Study objectives

To determine whether total prostatectomy or radiotherapy confer benefit in terms of survival or development of metastases over no immediate treatment alone.

#### 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

Prostate cancer

#### **Interventions**

- 1. The first group receives total prostatectomy
- 2. The second receive radiotherapy
- 3. The third receive no immediate treatment

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome(s)

- 1. Survival time to metastases
- 2. Morbidity
- 3. Local recurrence
- 4. Quality of life
- 5. Time to progression

### Key secondary outcome(s))

Not provided at time of registration

#### Completion date

01/06/1996

#### Reason abandoned (if study stopped)

The overall target for recruitment was 1800 patients, but the trial accrued only 35 (mean age 69 years, median PSA level at entry 12 ng/mL). Recruitment was formally halted following the recommendation of the independent Data Monitoring Committee in October 1996.

# **Eligibility**

#### Key inclusion criteria

- 1. Newly diagnosed, biopsy-proven adenocarcinoma of the prostate
- 2. Stage T1b/T1c/T2, N0, M0
- 3. Negative bone scan
- 4. No previous malignancy (except skin)

#### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

No

#### Age group

**Not Specified** 

#### Sex

Male

#### Key exclusion criteria

Opposite of inclusion criteria

#### Date of first enrolment

01/06/1994

#### Date of final enrolment

01/06/1996

# Locations

#### Countries of recruitment

**United Kingdom** 

England

Belgium

**Switzerland** 

## Study participating centre Department of Urology

London United Kingdom SW3 6JJ

# Sponsor information

#### Organisation

Medical Research Council (MRC) (UK)

# Funder(s)

## Funder type

Research council

#### **Funder Name**

Medical Research Council (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?
Other publications	Early closure in reported in 2004	01/12/2004		Yes	No