

A phase III study of neoadjuvant bicalutamide in patients with intermediate risk prostate cancer undergoing radiation therapy

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| Submission date 27/10/2004 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| | | <input type="checkbox"/> Protocol |
| Registration date 25/01/2005 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan |
| | | <input type="checkbox"/> Results |
| Last Edited 18/02/2008 | 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

99-07

Study information

Scientific Title

Study objectives

Not provided at time of registration

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

Localised prostate cancer

Interventions

Bicalutamide 150 mg daily for 3 months before and 2 months during radiation therapy versus no hormonal therapy.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Bicalutamide

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/06/2006

Eligibility**Key inclusion criteria**

1. T1/T2a prostate cancer with:
 - 1.1. Gleason 7, prostate specific antigen (PSA) less than 20 ng/ml
 - 1.2. Gleason less than 7, PSA 10 - 20 ng/ml
2. T2b prostate cancer with:

- 2.1. PSA less than 20 ng/ml
- 2.2. Gleason less than or equal to 7

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Male

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/07/1999

Date of final enrolment

30/06/2006

Locations

Countries of recruitment

Canada

Study participating centre

610 University Avenue

Toronto

Canada

M5G 2M9

Sponsor information

Organisation

Princess Margaret Hospital (Canada) - Department of Radiation Oncology

ROR

<https://ror.org/03zayce58>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Princess Margaret Hospital (Canada) - internal genitourinary (GU) group funds

Funder Name

AstraZeneca (Canada) - small unrestricted grant

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration