ISRCTN44967321 https://doi.org/10.1186/ISRCTN44967321

A Study to Compare the Effect of Two Blinded Doses of Casodex (ICI 176,334 100 mg and 150 mg daily) and Castration in the Treatment of Advanced Carcinoma of the Prostate

Submission date
19/08/2002

Recruitment status
No longer recruiting

Registration date
19/08/2002

Overall study status
Completed

Last Edited
21/01/2019

Condition category
Cancer

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr - -

Contact details
UKCCCR Register Co-ordinator
MRC Clinical Trials Unit
222 Euston Road
London
United Kingdom
NW1 2DA

Additional identifiers

Protocol serial number
CASODEX 0307

Study information

Scientific Title

A Study to Compare the Effect of Two Blinded Doses of Casodex (ICI 176,334 100 mg and 150 mg daily) and Castration in the Treatment of Advanced Carcinoma of the Prostate

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)
Not Specified

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

Interventions
STAGE I PATIENTS: Patients are randomised to either:
1. Casodex 100 mg
2. Casodex 150 mg
3. Castration

STAGE II PATIENTS: Patients are randomised to either:
1. Casodex 150 mg
2. Castration

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Casodex

Primary outcome(s)
Not provided at time of registration

Key secondary outcome(s))
Not provided at time of registration

Completion date

30/06/1993

Eligibility

Key inclusion criteria
1. Locally advanced stage T3 or T4 carcinoma of the prostate with prostate-specific antigen times five the upper limit of normal or metastatic disease
2. Gleason grade 2-10
3. Evaluable disease
4. Fit to receive treatment

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Male

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/1990

Date of final enrolment
30/06/1993

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
AstraZeneca Clinical Research Group (UK)

ROR
https://ror.org/04r9x1a08

Funder(s)

Funder type
Industry

Funder Name
AstraZeneca Pharmaceuticals (UK)

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?
Results article	results	01/03/1998	21/01/2019	Yes	No