# Quality of life assessment and patient acceptability of intermittent versus continuous sex hormone suppression as treatment for locally advanced or metastatic prostatic cancer

Submission date	Recruitment status	Prospectively registered		
30/09/2005  Registration date	No longer recruiting  Overall study status	☐ Protocol		
		Statistical analysis plan		
30/09/2005	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
17/09/2012	Cancer			

## Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

Mr Timothy Philp

#### Contact details

Consultant Urologist WXUHT London United Kingdom E11 1NR

# Additional identifiers

Protocol serial number N0080056792

# Study information

Scientific Title

#### **Study objectives**

This study will investigate patient acceptance, quality of life and survival/disease progression free interval in patients with locally advanced or metastatic prostate cancer in remission after six months treatment and disease control with androgen blockade.

#### 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

Cancer: Prostate

#### **Interventions**

Intermittent vs continuous sex hormone suppression

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome(s)

Patient acceptance, quality of life and survival/disease progression.

#### Key secondary outcome(s))

Not provided at time of registration

#### Completion date

31/12/2003

# **Eligibility**

#### Key inclusion criteria

200 patients recruited from Urological Oncology Clinical who have had six months androgen blockade.

#### 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

31/12/1997

#### Date of final enrolment

31/12/2003

## Locations

#### Countries of recruitment

**United Kingdom** 

England

# Study participating centre

**Consultant Urologist** 

London United Kingdom E11 1NR

# Sponsor information

#### Organisation

Department of Health

# Funder(s)

### Funder type

Charity

#### **Funder Name**

# **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	01/10/2004		Yes	No