

Interactive Video Disk and Patient Treatment Choices for benign prostatic hyperplasia (BPH)

Submission date 23/01/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 23/01/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 31/10/2019	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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
PSI04-01

Study information

Scientific Title
Interactive Video Disk and Patient Treatment Choices for benign prostatic hyperplasia (BPH)

Study objectives

The objective was to determine whether the provision of detailed information about BPH treatment options and outcomes via an interactive video disk affected patient decision making.

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

Urological and genital diseases: benign prostatic hyperplasia (BPH)

Interventions

1. Video viewing
2. Standard care

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Providing detailed information about the risks and benefits of treatment via the interactive video did not significantly appear to affect treatment choice. The interactive video was considered acceptable and easy to understand by patients, and despite the lack of a statistically significant difference in treatment choice, we cannot exclude the possibility that important shifts in the decision making process may occur, when a video such as this is introduced into clinical practice.

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/02/1999

Eligibility**Key inclusion criteria**

Patients from the Urology Department at Ashford Hospital, Middlesex.

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

01/10/1994

Date of final enrolment

01/02/1999

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Health Promotion Sciences Unit

London

United Kingdom

WC1E 7HT

Sponsor information**Organisation**

Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Primary and Secondary Care Interface National Research and Development Programme (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration