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

Submission date	Recruitment status	[_] Prospectively registered
23/01/2004	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
23/01/2004	Completed	[_] Results
Last Edited	**Condition category**	[_] Individual participant data
31/10/2019	Urological and Genital Diseases	[_] 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

Primary study design
Interventional

Study design
Randomised controlled trial

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