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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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

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 measure
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.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/1994

Completion date
01/02/1999

Eligibility

Key inclusion criteria
Patients from the Urology Department at Ashford Hospital, Middlesex.

Participant type(s)
Patient

Age group
Not Specified

Sex
Male

Target number of participants
Not provided at time of registration

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
England

United Kingdom

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)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

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

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration