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Prospective randomised controlled trial comparing 1% and 2% lignocaine as a form of anaesthesia prior to prostate biopsy

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 12/02/2018	**Condition category** Cancer	[_] Individual participant data [_] 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

N0059132320

Study information

Scientific Title
Prospective randomised controlled trial comparing 1% and 2% lignocaine as a form of anaesthesia prior to prostate biopsy

Study objectives
To identify any difference in relief of discomfort of prostate biopsy when using 1% or 2% lignocaine periprostatic nerve block

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)
Hospital

Study type(s)
Other

Participant information sheet
Not available in web format, please use the contact details to request a patient information sheet

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

Interventions
Prospective randomised controlled trial comparing 1% and 2% lignocaine as a form of anaesthesia prior to prostate biopsy

Intervention Type
Drug

Phase
Not Applicable

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

Primary outcome measure
Difference on visual analogue scale (VAS)

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2004

Completion date
30/04/2004

Eligibility

Key inclusion criteria
CaP (prostate cancer) patients

Participant type(s)
Patient

Age group
Adult

Sex
Male

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/01/2004

Date of final enrolment
30/04/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University of Sheffield
Sheffield
United Kingdom
S26 4SY

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Sheffield Teaching Hospitals NHS Foundation Trust (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