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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

Protocol serial number
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

Study type(s)
Other

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(s)
Difference on visual analogue scale (VAS)

Key secondary outcome(s))
Not provided at time of registration

Completion date
30/04/2004

Eligibility

Key inclusion criteria

CaP (prostate cancer) patients

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
Male

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
United Kingdom

England

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

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Sheffield Teaching Hospitals NHS Foundation Trust (UK)

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes