

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

Contact name

Mr Santbir Mehta

Contact details

University of Sheffield

Urology

Royal Hallamshire Hospital

Sheffield

United Kingdom

S26 4SY

+44 (0)788 956 9670/271 2154

sampimehta@aol.com

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

Primary study design

Interventional

Study design

Randomised controlled trial

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