

Combining surface and infiltration anaesthesia for Transrectal Ultrasonography of the Prostate (TRUS) biopsy: is it better?

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 11/04/2014	Condition category Surgery	<input type="checkbox"/> Individual participant data <input type="checkbox"/> 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
N0155153512

Study information

Scientific Title

Study objectives

Is combining surface and infiltration anaesthesia for Transrectal Ultrasonography of the Prostate (TRUS) biopsy more effective in relieving the pain of the procedure?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double blind randomised controlled trial

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Surgery: Anaesthesia

Interventions

The intervention is infiltration of local anaesthesia in periprostatic tissue with 1% lignocaine solution: and the ano-rectal mucosa anaesthetized by application of 2% lignocaine jelly.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

Pain relief by visual analogue score

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/09/2004

Eligibility

Key inclusion criteria

80 patients undergoing Transrectal Ultrasonography of the Prostate (TRUS) biopsy

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Known allergy to lignocaine.

Date of first enrolment

07/07/2004

Date of final enrolment

30/09/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre**Urology Unit**

Oldham

United Kingdom

OL1 2PN

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

Pennine Acute Hospitals NHS Trust (UK), NHS R&D Support Funding Own Account

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration