

Analgesia During Arterial Puncture Trial in the Emergency Department: a comparison of lignocaine, cryogestic spray and nothing.

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 06/01/2009	Condition category Signs and Symptoms	<input type="checkbox"/> Individual participant data

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

N0234123538

Study information

Scientific Title

Acronym

ADAPTED

Study objectives

Does cryogesic spray provide adequate local anaesthesia for patients undergoing arterial puncture compared with lignocaine and our current practice of nothing.

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)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Signs and Symptoms: Pain

Interventions

Lignocaine, cryogesic spray or nothing.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Lignocaine, cryogesic spray

Primary outcome measure

Pain scores for arterial puncture with and without anaesthesia

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2003

Completion date

01/12/2003

Eligibility

Key inclusion criteria

Patients presenting to Emergency Department requiring arterial puncture.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/06/2003

Date of final enrolment

01/12/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Emergency Dept
Bristol
United Kingdom
BS16 1ND

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
Government

Funder Name
North Bristol NHS 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/08/2008		Yes	No