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

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 06/01/2009	**Condition category** Signs and Symptoms	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr F Beech

Contact details
Emergency Dept
North Bristol NHS Trust
Frenchay Hospital
Frenchay
Bristol
United Kingdom
BS16 1ND
+44 (0)117 970 1212

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