

A randomised double-blind trial to investigate the efficacy of intra-articular bupivacaine for pain relief following ankle arthroscopy

Submission date 12/09/2003	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 12/09/2003	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 18/11/2009	Condition category Signs and Symptoms	☐ 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

Protocol serial number

N0283122707

Study information

Scientific Title

Study objectives

1. To investigate whether intra-articular bupivacaine provides pain relief following ankle arthroscopy
2. To investigate whether intra-articular bupivacaine reduces supplemental analgesia requirements in the post-operative period

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised double blind trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Signs and Symptoms: Pain

Interventions

1. Normal saline
2. 0.5% Bupivacaine

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

bupivacaine

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/03/2006

Eligibility

Key inclusion criteria

48 patients aged ≥ 18 years old undergoing day case ankle arthroscopy who have given informed written consent.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2003

Date of final enrolment

31/03/2006

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Worthing & Southlands Hospitals NHS Trust

Worthing

United Kingdom

BN11 2DH

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

Sussex NHS Research Consortium (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?
Results article	results	01/12/2006		Yes	No