

A randomised controlled trial to determine the efficacy of two different corticosteroid injections in the treatment of interdigital neuromas of the foot

Submission date 02/12/2004	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 27/04/2005	Overall study status Completed	☐ Protocol
Last Edited 28/07/2017	Condition category Musculoskeletal Diseases	☐ Statistical analysis plan
		☐ Results
		☐ Individual participant data
		☐ 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

CC 04 01

Study information

Scientific Title

A randomised controlled trial to determine the efficacy of two different corticosteroid injections in the treatment of interdigital neuromas of the foot

Study objectives

To determine the efficacy of two different corticosteroid injections in the treatment of interdigital neuromas of the foot.

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Interdigital neuromas of the foot

Interventions

Comparison of the efficacy of the following injections administered into the affected interdigital space of the foot:

1. Betamethasone 4 mg and mepivacaine 3%
2. Depomedrone 80 mg and mepivacaine 3%
3. Mepivacaine 3% as a control

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Betamethasone, mepivacaine, depomedrone

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/05/2005

Eligibility

Key inclusion criteria

Patients presenting with clinical signs of interdigital neuromas of the foot.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/04/2000

Date of final enrolment

31/05/2005

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Stafford Central Clinic

Stafford

United Kingdom

ST16 3AE

Sponsor information

Organisation

Cannock Chase Primary Care Trust (UK)

Funder(s)

Funder type

Government

Funder Name

Cannock Chase Primary Care Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration