

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	<input type="checkbox"/> Prospectively registered
Registration date 27/04/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 28/07/2017	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Contact name

Mr Paul Bewick

Contact details

Podiatric Surgery Department
Stafford Central Clinic
North Walls
Stafford
United Kingdom
ST16 3AE
+44 (0)1785 256323
Paul.bewick@ssh-tr.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

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 measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/2000

Completion date

31/05/2005

Eligibility

Key inclusion criteria

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

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

01/04/2000

Date of final enrolment

31/05/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Stafford Central Clinic
Stafford
United Kingdom
ST16 3AE

Sponsor information

Organisation

Cannock Chase Primary Care Trust (UK)

Sponsor details

Block D
Beecroft Court
Beecroft Road
Cannock
England
United Kingdom
WS11 1JP
+44 (0)1543 465100
Roger.whittaker@ssh-tr.nhs.uk

Sponsor type

Hospital/treatment centre

Funder(s)

Funder type

Government

Funder Name

Cannock Chase Primary Care 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