ISRCTN97845378 https://doi.org/10.1186/ISRCTN97845378

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

Registration date
27/04/2005

Overall study status
Completed

Last Edited
28/07/2017

Condition category
Musculoskeletal Diseases

[ ] Prospectively registered
[ ] Protocol
[ ] 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

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
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England
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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