The use of topical glyceryl trinitrate (GTN) for the treatment of plantar fasciitis. A prospective double-blinded randomised trial

Recruitment status	Prospectively registered
No longer recruiting	☐ Protocol
Overall study status	Statistical analysis plan
Completed	Results
Condition category	Individual participant data
Musculoskeletal Diseases	Record updated in last year
	No longer recruiting Overall study status Completed Condition category

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Ms Gail Ferrier

Contact details

Consultant Orthopaedic Surgeon North Cumbria Acute Hospitals NHS Trust Cumberland Infirmary Carlisle United Kingdom CA2 7HY

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0055166069

Study information

Scientific Title

The use of topical glyceryl trinitrate (GTN) for the treatment of plantar fasciitis: a prospective double-blinded randomised trial

Study objectives

Does the use of a topical GTN patch used with pressure relieving insoles and stretching exercises improve the symptoms of plantar fasciitis (heel pain) compared with current treatments?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double blinded randomised trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Muskuloskeletal Diseases: Plantar fasciitis

Interventions

One group will receive pressure relieving insoles and GTN patches, the other will receive pressure relieving insoles and placebo patches. Both groups will be instructed as to the use of the patches.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Topical glyceryl trinitrate

Primary outcome measure

Subjective, using questionnaires with visual analogue pain score

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2005

Completion date

30/07/2008

Eligibility

Key inclusion criteria

2 groups of 40 patients with chronic heel pain.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

80

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/08/2005

Date of final enrolment

30/07/2008

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

North Cumbria Acute Hospitals NHS Trust

Carlisle United Kingdom CA2 7HY

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details

The Department of Health Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

North Cumbria Acute Hospitals 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 summaryNot provided at time of registration