

Clinical trial comparing sclerification and non sclerification during nail surgery techniques

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 06/10/2014	Condition category Surgery	<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
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Horton Park
Bradford
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BD7 3EG

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0626176968

Study information

Scientific Title

Study objectives

Not provided at time of registration

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)

Other

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Nails

Interventions

When referrals are received by the nail surgery team leader the patient will be selected for one of the two methods of treatment. A sticker will be attached to the patient's notes, green for sclerify and red for no sclerification. An appointment will be sent to the patient including an information pack outlining the research taking place and issues regarding consent. Attached to the patient notes will be a data sheet, for the podiatrist performing the procedure to complete.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

To identify which technique if any is more beneficial and cost effective.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2005

Completion date

30/09/2007

Eligibility

Key inclusion criteria

The research will take place at Shipley Health Centre, Horton Park Podiatry Clinic and New Cross Street Podiatry Clinic.

The selection criteria will be the existing protocol for a nail surgery patient and aged 16 years and over.

The database to administrate the trial will be stored in a secure location.

Consent will be obtained prior to participation in the study.

Participant type(s)

Patient

Age group

Adult

Lower age limit

16 Years

Sex

Both

Target number of participants

200

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/09/2005

Date of final enrolment

30/09/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Podiatry Department
Bradford
United Kingdom
BD7 3EG

Sponsor information

Organisation

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

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

Government

Funder Name

Bradford South and West 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