

# Clinical trial comparing sclerification and non sclerification during nail surgery techniques

<b>Submission date</b> 29/09/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 29/09/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 06/10/2014	<b>Condition category</b> 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**  
Ms Susan Walton

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Horton Park  
Bradford  
United Kingdom  
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