# Comparison of two methods of steroid for trigger finger

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
29/09/2006	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
29/09/2006	Completed	Results
Last Edited	Condition category	Individual participant data
31/03/2020	Signs and Symptoms	<ul><li>Record updated in last year</li></ul>

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Mr Grey Giddins

### Contact details

Orthopaedics Royal United Hospital B&NES Bath United Kingdom BA1 3NG

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abc@email.com

# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers

N0212168002

# Study information

### Scientific Title

Comparison of two methods of steroid for trigger finger

### Study objectives

There are two recognised methods of injection for trigger finger used by the hand consultants in Bath. We wish to determine whether one method is more acceptable to the patient than the other in terms of pain.

# 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

Signs and Symptoms: Pain

### **Interventions**

- 1. Lateral approach
- 2. Palmare approach

# Intervention Type

Other

#### Phase

**Not Specified** 

### Primary outcome measure

Patient perception of discomfort of injection (using Visual Analogue Scale)

### Secondary outcome measures

Not provided at time of registration

# Overall study start date

18/07/2005

# Completion date

18/07/2006

# **Eligibility**

# Key inclusion criteria

All patients seen in the hand clinic with trigger finger on whom an injection of steroid is due to be given.

# Participant type(s)

**Patient** 

### Age group

**Not Specified** 

#### Sex

**Not Specified** 

# Target number of participants

50

# Key exclusion criteria

Patients with rheumatoid arthritis.

### Date of first enrolment

18/07/2005

### Date of final enrolment

18/07/2006

# Locations

# Countries of recruitment

England

United Kingdom

# Study participating centre Orthopaedics

Bath United Kingdom BA1 3NG

# 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

Government

#### **Funder Name**

Royal United Hospital Bath NHS Trust (UK) NHS R&D Support Funding

# **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