

# Capsular repair for painful wrist ganglion - does it affect the outcome?

<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> 12/08/2016	<b>Condition category</b> Musculoskeletal Diseases	<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

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**  
N0155176206

# Study information

## Scientific Title

Capsular repair for painful wrist ganglion - does it affect the outcome?

## Study objectives

Does capsular repair following excision of painful wrist ganglion reduce pain and recurrence post operatively?

## 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)

Hospital

## 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

Musculoskeletal Diseases: Wrist ganglion

## Interventions

Capsular repair vs standard care

## Intervention Type

Other

## Phase

Not Specified

## Primary outcome measure

Reduction of pain and recurrence

## Secondary outcome measures

Not provided at time of registration

**Overall study start date**

01/03/2006

**Completion date**

01/03/2008

## **Eligibility**

**Key inclusion criteria**

1. Age 16-90
2. Male or female
3. Painful wrist ganglion
4. Mentally competent and able to give informed consent

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Both

**Target number of participants**

44

**Key exclusion criteria**

1. Patients with rheumatoid or osteo-arthritis
2. Pregnant women

**Date of first enrolment**

01/03/2006

**Date of final enrolment**

01/03/2008

## **Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

Orthopaedics Department  
Oldham

United Kingdom  
OL1 2JH

## Sponsor information

### Organisation

Department of Health

### Sponsor details

Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL  
+44 20 7307 2622  
dhmail@doh.gsi.org.uk

### Sponsor type

Government

### Website

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

### ROR

<https://ror.org/03sbpja79>

## Funder(s)

### Funder type

Hospital/treatment centre

### Funder Name

Pennine Acute Hospitals NHS Trust

## 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