# Taping and stability of the shoulder girdle: a treatment for subacromial (rotator cuff) syndromes

Recruitment status	<ul><li>Prospectively registered</li></ul>
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

Mr PJ Pender

#### Contact details

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# Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

# Study information

## Scientific Title

Taping and stability of the shoulder girdle: a treatment for subacromial (rotator cuff) syndromes

## **Study objectives**

Can taping the shoulder girdle to improve its stability give pain relief from subacromial (rotator cuff) syndromes?

## 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

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

#### **Interventions**

- 1. Active scapular taping
- 2. Neutral scapular taping

## Intervention Type

Other

#### Phase

Not Applicable

## Primary outcome measure

The main outcome measures will be pain relief and patient preference for taping position. Pain will be measured using a visual analogue scale before and after tape application in the abducted shoulder position. Participant's preference to taping will also be noted.

## Secondary outcome measures

Not provided at time of registration

## Overall study start date

04/11/1996

## Completion date

30/04/1998

# Eligibility

## Key inclusion criteria

- 1. 25 patients with shoulder pain of at least 2 months duration will be studied
- 2. All will be aged 18-45 years with anterior instability, a positive impingement test and normal plain radiographs
- 3. Recruitment will be from rheumatology and orthopaedic outpatient clinics

## Participant type(s)

**Patient** 

## Age group

Adult

## Lower age limit

18 Years

## Upper age limit

45 Years

#### Sex

Both

## Target number of participants

25

## Key exclusion criteria

Not provided at time of registration

## Date of first enrolment

04/11/1996

## Date of final enrolment

30/04/1998

# Locations

# Countries of recruitment

England

**United Kingdom** 

# Study participating centre Royal Cornwall Hospitals NHS Trust

Truro United Kingdom TR1 2HZ

# Sponsor information

## Organisation

NHS R&D Regional Programme Register - 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.doh.gov.uk

# Funder(s)

# Funder type

Government

## **Funder Name**

NHS Executive South West (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