

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

Submission date 23/01/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 23/01/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 30/10/2019	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Protocol serial number
R/42/3.96/Spra

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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

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(s)

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.

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Upper age limit

45 Years

Sex

All

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

United Kingdom

England

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)

Funder(s)**Funder type**

Government

Funder Name

NHS Executive South West (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration