

# A pilot study to assess the effectiveness of adhesive taping in the management of patients with acute and chronic neck pain

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 31/03/2020	<b>Condition category</b> 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

### Contact name

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### Contact details

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United Kingdom  
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## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0544093490

# Study information

## Scientific Title

A pilot study to assess the effectiveness of adhesive taping in the management of patients with acute and chronic neck pain

## Study objectives

The proposed pilot study aims to determine the effectiveness of taping in providing symptomatic relief for patients with neck pain. Neck pain and stiffness is reported in 26-71% of the adult population at some point in their lifetime and thus has significant socioeconomic implications. There is little evidence to support the effectiveness of physiotherapy in the management of this patient population and whilst taping is widely used in clinical practice in patients with neck pain, an extensive literature search has failed to reveal the existence of any research to support its use in this patient group. It is proposed therefore to undertake a study to investigate the effect of taping in patients with acute and chronic neck pain with respect to pain, range of movement, neural tension, muscle activity and disability.

## 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

## Health condition(s) or problem(s) studied

Musculoskeletal Diseases: Neck pain

## Interventions

Patients will be randomly assigned to a taping or placebo taping group and will be required to attend twice a week for 2 weeks with a 1, 3 and 12 month follow-up. If successful the findings of this pilot study will lead to the undertaking of a randomised controlled trial.

## Intervention Type

Other

**Phase**

Not Specified

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

04/08/2000

**Completion date**

04/08/2003

**Eligibility****Key inclusion criteria**

20 subjects aged 18-80

**Participant type(s)**

Patient

**Age group**

Adult

**Lower age limit**

18 Years

**Upper age limit**

80 Years

**Sex**

Not Specified

**Target number of participants**

20

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

04/08/2000

**Date of final enrolment**

04/08/2003

**Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**Addenbrooke's NHS Trust**

Cambridge

United Kingdom

CB2 2QQ

## **Sponsor information**

**Organisation**

Department of Health (UK)

**Sponsor details**

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

**Sponsor type**

Government

**Website**

<http://www.doh.gov.uk>

## **Funder(s)**

**Funder type**

Government

**Funder Name**

Cambridge Consortium - Addenbrooke's (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