

# The development of a psychometrically sound questionnaire to adequately record patients' needling sensation during acupuncture

<b>Submission date</b> 16/02/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 03/04/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 12/09/2011	<b>Condition category</b> Signs and Symptoms	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Dr Peter White

### 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

05/Q1702/157

# Study information

## Scientific Title

## Study objectives

Fourth-year medical student project.

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

Approved by the Southampton and South West Hampshire Research Ethics Committee on 27/1/06, reference number: 05/Q1702/157

## Study design

Qualitative and questionnaire-based design

## Primary study design

Observational

## Secondary study design

Other

## Study setting(s)

Not specified

## Study type(s)

Treatment

## Participant information sheet

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

Pain

## Interventions

Acupuncture

## 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**

06/02/2006

**Completion date**

01/10/2007

## Eligibility

**Key inclusion criteria**

1. Aged between 18 and 80 (inclusive)
2. To have recently received acupuncture treatment

**Participant type(s)**

Patient

**Age group**

Adult

**Lower age limit**

18 Years

**Sex**

Both

**Target number of participants**

1st phase = 50, 2nd qualitative phase = 10, 3rd phase = 100

**Key exclusion criteria**

Neurological impairment (no sensation)

**Date of first enrolment**

06/02/2006

**Date of final enrolment**

01/10/2007

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

School of Health Professions

Southampton

United Kingdom

SO17 1BJ

# Sponsor information

## Organisation

University of Southampton (UK)

## Sponsor details

Highfield

Southampton

England

United Kingdom

SO17 1BJ

jbk1@soton.ac.uk

## Sponsor type

University/education

## ROR

<https://ror.org/01ryk1543>

# Funder(s)

## Funder type

University/education

## Funder Name

University of Southampton (UK)

## Alternative Name(s)

University of Southampton UK

## Funding Body Type

Government organisation

## Funding Body Subtype

Universities (academic only)

## Location

United Kingdom

# 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

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/05/2008		Yes	No