

# Effect of acupressure at P6 on nausea and vomiting during labour

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 30/09/2016	<b>Condition category</b> Pregnancy and Childbirth	<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**  
N0084078266

## Study information

**Scientific Title**

Effect of acupressure at P6 on nausea and vomiting during labour

**Study objectives**

Can the use of acupressure during labour help prevent nausea and vomiting?

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Study design**

Randomised placebo controlled parallel group trial

**Primary study design**

Interventional

**Secondary study design**

Randomised parallel trial

**Study setting(s)**

Not specified

**Study type(s)**

Prevention

**Participant information sheet****Health condition(s) or problem(s) studied**

Pregnancy and Childbirth: Vomiting, nausea

**Interventions**

Randomised controlled trial comparing (a) acupressure at P6 versus (b) acupressure at a point with no known acupressure point or pathway.

**Intervention Type**

Other

**Phase**

Not Applicable

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/07/2000

**Completion date**

30/06/2003

## Eligibility

**Key inclusion criteria**

100 women in each group.

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Female

**Target number of participants**

200

**Key exclusion criteria**

Does not match inclusion criteria

**Date of first enrolment**

01/07/2000

**Date of final enrolment**

30/06/2003

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

18 Northwood Drive

Hessle, East Yorkshire

United Kingdom

HU13 0TA

## 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**

Research organisation

**Funder Name**

The North and South Bank Research and Development Consortium (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