

# Internal jugular vein compression to assess the correct placement of an epidural catheter in post-partum women.

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 13/10/2014	<b>Condition category</b> Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

**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**  
N0047126465

# Study information

## Scientific Title

### Study objectives

Validity of the internal jugular vein compression test to confirm correct placement of the epidural catheter.

### 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)

Not Specified

## Participant information sheet

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

Pregnancy and Childbirth: Anaesthesia

### Interventions

Patient volunteers, Single Centre Prospective, Observational Single-blind, Randomised. Chi Squared test constructing a two by two table to input the data.

### Intervention Type

Other

### Phase

Not Specified

### Primary outcome measure

Time interval after completion of the third stage of labour

### Secondary outcome measures

Total duration of the epidural Infusion. Patient weight. Depth of epidural space. Length of catheter in the epidural space.

**Overall study start date**

01/01/2003

**Completion date**

01/10/2003

## **Eligibility**

**Key inclusion criteria**

20 volunteers, post-partum women who have received an epidural for pain relief in labour.

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Female

**Target number of participants**

20

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/2003

**Date of final enrolment**

01/10/2003

## **Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**Anaesthetic Department: Labour Ward**

Birmingham

United Kingdom

B15 2TH

# Sponsor information

## Organisation

Department of Health

## Sponsor details

Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL

## Sponsor type

Government

## Website

<http://www.dh.gov.uk/Home/fs/en>

# Funder(s)

## Funder type

Government

## Funder Name

Birmingham Women's Healthcare NHS Trust (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

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/04/2007		Yes	No

