# Using moxibustion to turn the breech baby

| Submission date   | Recruitment status       | <ul><li>Prospectively registered</li></ul>    |
|-------------------|--------------------------|-----------------------------------------------|
| 30/09/2004        | No longer recruiting     | <pre>Protocol</pre>                           |
| Registration date | Overall study status     | <ul><li>Statistical analysis plan</li></ul>   |
| 30/09/2004        | Completed                | Results                                       |
| Last Edited       | Condition category       | Individual participant data                   |
| 28/10/2016        | Pregnancy and Childbirth | <ul><li>Record updated in last year</li></ul> |

# Plain English summary of protocol

Not provided at time of registration

# Contact information

### Type(s)

Scientific

#### Contact name

Mr T Draycott

#### Contact details

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# Additional identifiers

# Protocol serial number

N0234131719

# Study information

#### Scientific Title

Using moxibustion to turn the breech baby

# Study objectives

Does moxibustion to the classical acupuncture point bladder 67 increase the version rate to cephalic in uncomplicated breech pregnancies?

#### 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

### Study type(s)

Treatment

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

Breech pregnancy

#### **Interventions**

A double blinded randomised controlled trial.

Does moxibustion to the classical acupuncture point bladder 67 increase the version rate to cephalic in uncomplicated breech pregnancies?

### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome(s)

Presentation of the baby one week after treatment and at delivery.

#### Key secondary outcome(s))

Not provided at time of registration

#### Completion date

01/07/2005

# **Eligibility**

#### Key inclusion criteria

210 women aged 16-46 years with an uncomplicated breech pregnancy at 34-35 weeks.

#### Participant type(s)

**Patient** 

#### Healthy volunteers allowed

### Age group

Adult

#### Sex

Female

### Key exclusion criteria

Not provided at time of registration

#### Date of first enrolment

01/01/2004

#### Date of final enrolment

01/07/2005

# Locations

## Countries of recruitment

**United Kingdom** 

England

# Study participating centre Department of Obstetrics and Gynaecology

Bristol United Kingdom BS10 5NB

# Sponsor information

### Organisation

Department of Health

# Funder(s)

# Funder type

Government

#### **Funder Name**

North Bristol NHS Trust

# **Results and Publications**

Individual participant data (IPD) sharing plan

**IPD sharing plan summary**Not provided at time of registration