

Using moxibustion to turn the breech baby

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 28/10/2016	Condition category Pregnancy and Childbirth	<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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

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 measure

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

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2004

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

Age group

Adult

Sex

Female

Target number of participants

210

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

England

United Kingdom

Study participating centre

Department of Obstetrics and Gynaecology

Bristol

United Kingdom

BS10 5NB

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

North Bristol NHS Trust

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