ISRCTN98448646 https://doi.org/10.1186/ISRCTN98448646

Using moxibustion to turn the breech baby

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2004	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
28/10/2016	Pregnancy and Childbirth	[ ] 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
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