ISRCTN74892692 https://doi.org/10.1186/ISRCTN74892692

A randomised controlled trial of interventions to promote external cephalic version (ECV) at term

Submission date
23/01/2004

Recruitment status
No longer recruiting

Registration date
23/01/2004

Overall study status
Completed

Last Edited
29/01/2010

Condition category
Pregnancy and Childbirth

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] 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
547

Study information

Scientific Title

Study objectives
Audit the number of uncomplicated breech pregnancies at term being offered External Cephalic Version (ECV) and evaluate the impact of providing a multifaceted intervention package on patient management in terms of eligible women being offered ECV.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Cluster randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Breech presentation birth

Interventions
1. ECV
2. No ECV

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Added 28/01/10:
The percentage of women with breech presentation at term who were offered ECV in the antenatal clinic, before and after the intervention

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/08/1995

Completion date
01/08/1997

Eligibility

Key inclusion criteria
Pregnant females with uncomplicated breech birth

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
20 consultant-based maternity units (added 28/01/10; see publication)

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/08/1995

Date of final enrolment
01/08/1997

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
North Staffs Maternity Hospital
Stoke-on-Trent
United Kingdom
ST4 6SD

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Executive West Midlands (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?
Results article	results	10/12/2001		Yes	No