

A randomised controlled trial of the effect of ambulation in the first stage of labour in terms of duration of labour of women with a previous caesarean

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 16/10/2015	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

Protocol serial number

N0170149149

Study information

Scientific Title

A randomised controlled trial of the effect of ambulation in the first stage of labour in terms of duration of labour of women with a previous caesarean

Study objectives

Will ambulation during labour shorten the first stage of labour in women who have previously undergone one previous caesarian?

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)

Not Specified

Health condition(s) or problem(s) studied

Pregnancy and Childbirth: Labour

Interventions

1. Encourage walking/mobilisation
2. Walk/mobilise as little/much as desired

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Duration of labour in each group

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/07/2006

Eligibility**Key inclusion criteria**

200 women in latter stages of pregnancy

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

1. Women who intend to have a second planned elective caesarian section
2. Women who do not understand/speak English well enough to give valid consent
3. Non-cephalic presentation; known fetal abnormality
4. More than one previous LSCS; previous classical, inverted T or J incision to the uterus
5. Multiple pregnancy
6. Pre-term labour 37 weeks
8. Induced labour

Date of first enrolment

20/07/2004

Date of final enrolment

31/07/2006

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Nottingham City Hospital

Nottingham

United Kingdom

NG5 1PB

Sponsor information**Organisation**

Department of Health

Funder(s)

Funder type
Government

Funder Name
Nottingham City Hospital NHS Trust (UK), NHS R&D Support Funding

Results and Publications

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes