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# 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

**Registration date**
30/09/2005

**Overall study status**
Completed

**Last Edited**
16/10/2015

**Condition category**
Pregnancy and Childbirth

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] 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**
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

### Secondary study design
Randomised controlled trial

### Study setting(s)
Hospital

### Study type(s)
Not Specified

### Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

### 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 measure
Duration of labour in each group

### Secondary outcome measures
Not provided at time of registration

### Overall study start date
20/07/2004

### Completion date
31/07/2006

## Eligibility

### Key inclusion criteria
200 women in latter stages of pregnancy

### Participant type(s)
Patient

### Age group
Adult

### Sex
Female

### Target number of participants
200

### 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
England

United Kingdom

**Study participating centre**
**Nottingham City Hospital**
Nottingham
United Kingdom
NG5 1PB

# Sponsor information

**Organisation**
Department of Health

**Sponsor details**
Richmond House
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**Sponsor type**
Government

**Website**
http://www.dh.gov.uk/Home/fs/en

# Funder(s)

**Funder type**
Government

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

# 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