Randomised trial of intrapartum tractocile versus placebo in the management of fetal distress in women undergoing emergency Caesarean section

Submission date	Recruitment status	[X] Prospectively registered
12/09/2003	Stopped	Protocol
Registration date	Overall study status	Statistical analysis plan
12/09/2003	Stopped	Results
Last Edited 05/10/2011	Condition category Pregnancy and Childbirth	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

Protocol serial number N0059103677

Study information

Scientific Title

Study objectives

Not provided at time of registration

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: Fetal distress

Interventions

Intrapartum tractocile versus placebo.

Added June 2008: trial abandoned and never recruited patients.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

28/02/2006

Reason abandoned (if study stopped)

Lack of resources

Eligibility

Key inclusion criteria

Women undergoing emergency Caesarean section.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2004

Date of final enrolment

28/02/2006

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

STH NHS Trust

Sheffield United Kingdom S10 2SF

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

Sheffield Teaching Hospitals (Central Campus) - UK

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration