# Effects of a slight delay in cord clamping time versus milking the cord in preterm infants

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>		
29/09/2006	No longer recruiting	☐ Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
29/09/2006		[X] Results		
Last Edited	Condition category	Individual participant data		
29/03/2012	Pregnancy and Childbirth			

#### Plain English summary of protocol

Not provided at time of registration

### Contact information

#### Type(s)

Scientific

#### Contact name

Dr Heike Rabe

#### Contact details

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#### Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0051177741

## Study information

#### Scientific Title

#### **Study objectives**

To compare two types of intervention regarding the clamping of the umbilical cord after delivery of a preterm infant less than 33 weeks gestation.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

East Sussex Research Ethics Committee, REC Ref no 06/Q1905/15, 21/04/2006.

#### Study design

Randomised controlled trial

#### Primary study design

Interventional

#### Secondary study design

Randomised controlled trial

#### Study setting(s)

Not specified

#### 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:

#### **Interventions**

Clamping method 1 vs clamping method 2

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome measure

Acceptability of cord clamping methods.

#### Secondary outcome measures

#### Not provided at time of registration

#### Overall study start date

01/04/2006

#### Completion date

30/04/2008

## **Eligibility**

#### Key inclusion criteria

- 1. Singletons
- 2. Inborn
- 3. Preterm infants < 34 weeks gestation

#### Participant type(s)

**Patient** 

#### Age group

Neonate

#### Sex

**Not Specified** 

#### Target number of participants

58 preterm infants

#### Key exclusion criteria

- 1. Multiple pregnancy
- 2. Congenital malformations
- 3. Rhesus incompatibility

#### Date of first enrolment

01/04/2006

#### Date of final enrolment

30/04/2008

### Locations

#### Countries of recruitment

England

**United Kingdom** 

## Study participating centre Brighton & Sussex University Hospitals NHS Trust (RSCH) Brighton

## Sponsor information

#### Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

#### 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.dh.gov.uk/Home/fs/en

## Funder(s)

#### Funder type

Government

#### **Funder Name**

Brighton and Sussex University Hospitals NHS Trust (UK)

#### Funder Name

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

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/10/2000		Yes	No