

# Randomised controlled trial of sucrose analgesia for repeated capillary blood sampling

<b>Submission date</b> 29/09/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 29/09/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 08/10/2014	<b>Condition category</b> Signs and Symptoms	<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**  
Dr Simon Mitchell

**Contact details**  
SMH Central Manchester & Manchester Children's University Hospitals  
St Mary's Hospital for Women & Children  
Oxford Road  
Manchester  
United Kingdom  
M13 0JH

## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

**Secondary identifying numbers**  
N0453162768

## Study information

## **Scientific Title**

### **Study objectives**

To compare chronic pain exposure in infants randomised to receive either sucrose analgesia or control (water) for capillary blood sampling.

### **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)**

Treatment

### **Participant information sheet**

### **Health condition(s) or problem(s) studied**

Chronic pain exposure

### **Interventions**

Infants were randomised to receive:

1. Sucrose analgesia
2. Control (water)

### **Intervention Type**

Drug

### **Phase**

Not Applicable

### **Drug/device/biological/vaccine name(s)**

Sucrose

### **Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/02/2005

**Completion date**

31/12/2006

## **Eligibility**

**Key inclusion criteria**

Not provided at time of registration

**Participant type(s)**

Patient

**Age group**

Child

**Sex**

Both

**Target number of participants**

120

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/02/2005

**Date of final enrolment**

31/12/2006

## **Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**SMH Central Manchester & Manchester Children's University Hospitals**

Manchester

United Kingdom

M13 0JH

# Sponsor information

## Organisation

Record Provided by the NHSTCT Register - 2006 Update - 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.dh.gov.uk/Home/fs/en>

# Funder(s)

## Funder type

Government

## Funder Name

Central Manchester and Manchester Children's University Hospitals NHS Trust (UK)

## Funder Name

NHS R&D Support Funding (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