

# The efficiency of Nurse Specialist led preoperative education for children and their parents in a day surgery unit who are going to surgical operations in reducing the perioperative time

<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> 18/10/2017	<b>Condition category</b> Surgery	<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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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

## Secondary identifying numbers

N0045168791

# Study information

## Scientific Title

The efficiency of Nurse Specialist led preoperative education for children and their parents in a day surgery unit who are going to surgical operations in reducing the perioperative time

## Study objectives

If children and their parents receive preoperative education from the Specialist Theatre Nurse then the perioperative time for those patients will differ from those who don't.

## 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)

Other

## 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

Surgery

## Interventions

Randomised controlled trial - quantitative data collection.

## Intervention Type

Procedure/Surgery

## Phase

Not Specified

**Primary outcome measure**

Education children and parents pre-operative will help to minimise perioperative time. Outcome measure is arrival time in theatre.

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/06/2005

**Completion date**

01/12/2005

**Eligibility****Key inclusion criteria**

Patients aged from 0-15 years old from Day Surgery Units (Ward 16) at Birmingham Children's Hospital.

**Participant type(s)**

Patient

**Age group**

Child

**Upper age limit**

15 Years

**Sex**

Not Specified

**Target number of participants**

84

**Key exclusion criteria**

No informed consent.

**Date of first enrolment**

01/06/2005

**Date of final enrolment**

01/12/2005

**Locations****Countries of recruitment**

England

United Kingdom

**Study participating centre****Main Theatres**

Birmingham

United Kingdom

B4 6NH

## **Sponsor information**

**Organisation**

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

**Sponsor details**

The Department of Health, Richmond House, 79 Whitehall

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**Sponsor type**

Government

**Website**

<http://www.dh.gov.uk/Home/fs/en>

## **Funder(s)**

**Funder type**

Hospital/treatment centre

**Funder Name**

Birmingham Children's Hospital NHS Trust (UK)

**Funder Name**

Self-funding

**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