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# 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

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| **Submission date**<br>29/09/2006 | **Recruitment status**<br>No longer recruiting | [ ] Prospectively registered<br>[ ] Protocol |
| **Registration date**<br>29/09/2006 | **Overall study status**<br>Completed | [ ] Statistical analysis plan<br>[ ] Results |
| **Last Edited**<br>18/10/2017 | **Condition category**<br>Surgery | [ ] Individual participant data<br>[ ] 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
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
London
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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