

The effect of videotaped preoperative information on parental anxiety during anesthetic for elective paediatric procedures

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 13/04/2010	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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
N0051154080

Study information

Scientific Title

Study objectives

Does videotape information about anaesthesia provided postoperatively reduce parental anxiety during anaesthetic induction in paediatric surgery?

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

Anxiety disorders

Interventions

Parents of children undergoing surgery will be randomised. One group will watch videotaped information, others will receive standardised information. Parental anxiety is assessed before and after witnessing their child's anaesthetic induction by completing a validated questionnaire.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

1. Anxiety before anaesthetic induction as measured by the state/trait anxiety inventory and the Amsterdam Preoperative anxiety and information score
2. Anxiety after anaesthetic induction as measured by the state/trait anxiety inventory

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/12/2004

Completion date

01/12/2005

Eligibility**Key inclusion criteria**

Parents of children having uneventful day case surgery under general anaesthetic.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

111

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/12/2004

Date of final enrolment

01/12/2005

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

Brighton & Sussex University Hospitals NHS Trust (RA)

Brighton
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BN1 3JN

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
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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 (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2007		Yes	No