

Peri-operative antibiotics in tonsillectomy patients

Submission date 12/09/2003	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 16/05/2018	Condition category 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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0234120874

Study information

Scientific Title

Peri-operative antibiotics in tonsillectomy patients

Study objectives

Does the peri-operative administration of antibiotics improve pain control in adults undergoing tonsillectomy?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised placebo controlled parallel group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Pain

Interventions

Double blind randomised placebo controlled trial.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

1. Visual analogue scale for pain at 24 h, 1 week, 2 weeks
2. Analgesia requirements at end of 2 weeks

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2004

Completion date

01/03/2005

Eligibility

Key inclusion criteria

Adult patients listed for tonsillectomy.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

186

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/03/2004

Date of final enrolment

01/03/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

ENT Department

Bristol

United Kingdom

BS10 5NB

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

North Bristol NHS Trust (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