ISRCTN52345875 https://doi.org/10.1186/ISRCTN52345875

Peri-operative antibiotics in tonsillectomy patients

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[X] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 16/05/2018	**Condition category** Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr H Khalil

Contact details
ENT Department
North Bristol NHS Trust
Southmead Hospital
Westbury-on-Trym
Bristol
United Kingdom
BS10 5NB
+44 0117 9505050
research@southmead.swest.nhs.uk

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