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Use of mitomycin C to improve endonasal dacryocystorhinostomy (DCR) success rates

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2004	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
25/03/2020	Surgery	[ ] 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
N0176108541

Study information

Scientific Title
Use of mitomycin C to improve endonasal dacryocystorhinostomy (DCR) success rates

Study objectives
A randomized controlled trial to determine whether intraoperative use of mitomycin C can improve the success rate of endonasal dacryocystorhinostomy (DCR)

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
Surgery: Dacryocystorhinostomy

Interventions
Mitomycin C vs standard practice

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
mitomycin C

Primary outcome measure
The patient being symptom free and/or patency to saline irrigation

Secondary outcome measures
Not provided at time of registration

Overall study start date
25/01/2002

Completion date
30/06/2004

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
20 patients and 20 control patients

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
25/01/2002

Date of final enrolment
30/06/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Oxford Eye Hospital
Oxford
United Kingdom
OX2 6HE

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Oxford Radcliffe Hospitals 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