

The role of mitomycin C (MMC) in Endoscopic Sinus Surgery (ESS) to the frontonasal recess: a prospective randomised trial

Submission date 21/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 02/11/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 14/09/2007	Condition category Ear, Nose and Throat	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Acronym

Mitomycin C Project

Study objectives

Not provided at time of registration

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

Chronic Rhinosinusitis (CRS) of the frontonasal recess

Interventions

Application of MMC/placebo following ESS and assessment for impact on symptoms of CRS

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2004

Completion date

01/01/2006

Eligibility

Key inclusion criteria

Patients who will be undergoing normal surgical procedure following the failure of medical treatment.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2004

Date of final enrolment

01/01/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Professorial Unit

London

United Kingdom

WC1X 8DA

Sponsor information

Organisation

Royal National Throat, Nose and Ear Hospital (UK)

Sponsor details

Professor VJ Lund
Professorial Unit
330 Gray's Inn Road
London
England
United Kingdom
WC1X 8DA

Sponsor type

Hospital/treatment centre

Website

http://www.royalfree.nhs.uk/default.aspx?top_nav_id=3&sel_left_nav=40&tab_id=121

ROR

<https://ror.org/03pf5zs64>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Royal National Throat, Nose and Ear Hospital (UK) - funded internally

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	01/11/2006		Yes	No