

Evaluation of Lacrimal Stent (mini monoka) for treatment of punctal stenosis: a prospective randomised comparative study

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 18/10/2017	Condition category Eye Diseases	<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
N0214126228

Study information

Scientific Title

Evaluation of Lacrimal Stent (mini monoka) for treatment of punctal stenosis: a prospective randomised comparative study

Study objectives

What is the use of lacrimal stent to relieve clinically significant Epiphora caused by punctal stenosis.

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Eye Diseases: Epiphora

Interventions

Randomised comparative study of single snip procedure vs mini monoka (monocanalicular lacrimal stent)

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Decrease in epiphora
 - 1.1. subjectively using the questionnaire
 - 1.2. objectively using fluorescein dye disappearance test

Secondary outcome measures

Not provided at time of registration

Overall study start date

30/05/2003

Completion date

30/09/2005

Eligibility

Key inclusion criteria

40 patients with punctal stenosis

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

40

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

30/05/2003

Date of final enrolment

30/09/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Wolverhampton & Midland Counties Eye Infirmary
Wolverhampton Road
Wolverhampton
United Kingdom
WV3 9QR

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

Hospital/treatment centre

Funder Name

The Royal Wolverhampton 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