

Comparison of efficacy and safety of two forms of cyclodestructive therapy. Transscleral cyclophotocoagulation and endoscopic cyclophotocoagulation in the treatment of patients with refractory glaucoma.

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 06/01/2009	Condition category Eye Diseases	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr Philip Bloom

Contact details
Ophthalmology
Western Eye Hospital
Marylebone Road
London
United Kingdom
NW1 5YE
+44 (0)20 7886 6666

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0241032308

Study information

Scientific Title

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Not provided at time of registration

Health condition(s) or problem(s) studied

Eye Diseases: Glaucoma

Interventions

Transscleral cyclophotocoagulation vs endoscopic cyclophotocoagulation

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

01/11/1998

Completion date

01/07/2005

Eligibility

Key inclusion criteria

100 Patients will be recruited, randomised into two groups.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

100 patients will be recruited, randomised into two groups.

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/1998

Date of final enrolment

01/07/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Ophthalmology
London
United Kingdom
NW1 5YE

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
Government

Funder Name
St Mary's 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2006		Yes	No