ISRCTN52778666 https://doi.org/10.1186/ISRCTN52778666

India Glaucoma Outcomes and Treatment Trial

Submission date	Recruitment status	[_] Prospectively registered
11/10/2005	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
30/03/2006	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
11/01/2021	Eye Diseases	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N/A

Study information

Scientific Title
India Glaucoma Outcomes and Treatment Trial

Acronym
INGOTT

Study objectives
Surgery will be more effective than medicines in controlling glaucoma in India

Ethics approval required
Old ethics approval format

Ethics approval(s)
Approved by the Internal Review Boards at the Johns Hopkins University, Baltimore, USA and the Aravind Eye Hospital in India

Study design
Randomised controlled trial

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Glaucoma

Interventions
Medicine (timolol, latanaprost and brimonidine) versus surgery

Intervention Type
Procedure/Surgery

Primary outcome(s)
1. Intraocular pressure
2. Visual acuity
3. Quality of life

Key secondary outcome(s))
Cataract grade

Completion date
01/01/2007

Eligibility

Key inclusion criteria
Glaucoma patients over the age of 30 years in India

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
All

Total final enrolment
398

Key exclusion criteria
1. Previous surgery
2. Intraocular pressure (IOP) >35

Date of first enrolment
01/01/2003

Date of final enrolment
01/01/2007

Locations

Countries of recruitment
India

United States of America

Study participating centre
Wilmer 120
Baltimore
United States of America
21287

Sponsor information

Organisation
Pfizer Inc. (USA)

ROR
https://ror.org/01xdqrp08

Funder(s)

Funder type

Industry

Funder Name
Pfizer Inc.

Results and Publications

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	01/06/2012	11/01/2021	Yes	No