Efficacy of trypan blue in posterior capsulorhexis with optic capture in pediatric cataracts

Submission date	Recruitment status	Prospectively registered
02/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
08/09/2005	Completed	[X] Results
Last Edited	Condition category	Individual participant data
03/10/2017	Eye Diseases	

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

Efficacy of trypan blue in posterior capsulorhexis with optic capture in pediatric cataracts:

Study objectives

Trypan blue staining helps to achieve good posterio capsulorhexis for optic capture in patients with pediatric cataracts: a randomised controlled trial

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

Cataracts

Interventions

Lens aspiration with posterior capsulorhexis with optic capture of intraocular lens with or without typan blue.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Trypan blue

Primary outcome measure

Achievement of optic capture of posterior chamber intraocular lens.

Secondary outcome measures

Good visual rehabilitation.

Overall study start date

01/01/2002

Completion date

31/12/2002

Eligibility

Key inclusion criteria

Children with developmental or congenital cataract with visually significant lenticular opacity (3 mm or larger).

Participant type(s)

Patient

Age group

Child

Sex

Both

Target number of participants

26

Key exclusion criteria

Eyes with traumatic cataract and associated ocular abnormalities were excluded.

Date of first enrolment

01/01/2002

Date of final enrolment

31/12/2002

Locations

Countries of recruitment

India

Study participating centre Rajendra Prasad Centre for Ophthalmic Sciences

New Delhi India 110029

Sponsor information

Organisation

Rajendra Prasad Centre for Ophthalmic Sciences (India)

Sponsor details

All India Institute of Medical Sciences (AIIMS) New Delhi India 110029

rvajpayee@rediffmail.com

Sponsor type

Research organisation

ROR

https://ror.org/02dwcqs71

Funder(s)

Funder type

Research organisation

Funder Name

Rajendra Prasad Centre for Ophthalmic Sciences, AIIMS, New Delhi (India)

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 typeDetailsDate createdDate addedPeer reviewed?Patient-facing?Results articleresults16/03/2006YesNo