ISRCTN84553798 https://doi.org/10.1186/ISRCTN84553798

Effect of Ketamine on post-operative intraocular inflammation following cataract surgery

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 18/02/2008	**Condition category** Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr S B Kaye

Contact details
St Pauls Eye Unit
8Z Link
Royal Liverpool University Hospital
Prescot Street
Liverpool
United Kingdom
L7 8XP
+44 (0)151 706 2134
Stephen.kaye@rlbuht.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0207129695

Study information

Scientific Title

Study objectives
Does ketamine reduce post-operative cystoid macula oedema in patients who undergo cataract surgery?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Prospective, randomised, parallel group, double blind, clinical 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
Surgery: Cataract

Interventions
Group A receive ketamine under general anaesthetic
Group B receive general anaesthetic without ketamine
Group C receive local anaesthetic without ketamine

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)

ketamine

Primary outcome measure
To define population for definitive randomised prospective trial on effects of ketamine on cystoid macula oedema in patients undergoing cataract surgery

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/02/2004

Completion date
01/07/2006

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Pilot study; 3 groups of 11.

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/02/2004

Date of final enrolment
01/07/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
St Pauls Eye Unit
Liverpool
United Kingdom
L7 8XP

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
Government

Funder Name
Royal Liverpool and Broadgreen University Hospitals 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	12/07/2007		Yes	No