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Quality of Vision with the Acrysof Natural Intraocular Lens

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 15/04/2015	**Condition category** Surgery	

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
N0199136008

Study information

Scientific Title
Quality of Vision with the Acrysof Natural Intraocular Lens

Study objectives
Does the new Acrysof Natural Intraocular Lens, which reduces the amount of damaging blue light reaching the retina, have any significant effect on visual outcome or colour vision post cataract surgery?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised prospective single-centre surgical study

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
Randomised, prospective single centre surgical study to determine whether the new Acrysof Natural Intraocular Lens has any significant effect on visual outcome or colour vision post cataract surgery

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
1. Colour vision
2. Visual acuity
3. Contrast sensitivity

Secondary outcome measures
Not provided at time of registration

Overall study start date
10/12/2003

Completion date
28/02/2006

Eligibility

Key inclusion criteria
20 patients and 20 controls having standard cataract surgery

Participant type(s)
Patient

Age group
Senior

Sex
Both

Target number of participants
40

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
10/12/2003

Date of final enrolment
28/02/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Royal Berkshire Hospital
Reading
United Kingdom
RG1 5AN

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 Berkshire and Battle 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