

# Quality of Vision with the Acrysof Natural Intraocular Lens

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 15/04/2015	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

**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**  
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

**Study type(s)**

Treatment

**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(s)**

1. Colour vision
2. Visual acuity
3. Contrast sensitivity

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

28/02/2006

**Eligibility**

**Key inclusion criteria**

20 patients and 20 controls having standard cataract surgery

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Senior

**Sex**

All

**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**

United Kingdom

England

**Study participating centre**

**Royal Berkshire Hospital**

Reading

United Kingdom

RG1 5AN

**Sponsor information****Organisation**

Department of Health

**Funder(s)****Funder type**

Government

**Funder Name**

Royal Berkshire and Battle Hospitals NHS Trust (UK)

**Results and Publications**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**

Not provided at time of registration