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Randomised controlled trial comparing two multifocal and a standard monofocal intraocular lens in cataract surgery

Submission date 29/09/2006	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 29/09/2006	**Overall study status** Completed	
Last Edited 28/03/2012	**Condition category** Eye Diseases	

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

N0388170265

Study information

Scientific Title

Study objectives
Is there a difference in functional outcome, principally reading acuity, between patients who have received the Tecnis multifocal, the ReSTORE multifocal or the standard monofocal intraocular lens in both eyes after cataract surgery?

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)
Not specified

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Eye Diseases: Cataract

Interventions
Randomised controlled trial

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

Reading acuity

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2005

Completion date
01/11/2008

Eligibility

Key inclusion criteria
All patients over the age of 18 undergoing cataract surgery who are assessed as suitable for lens implantation.

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/11/2005

Date of final enrolment
01/11/2008

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
The Hillingdon Hospital NHS Trust
Uxbridge
United Kingdom
UB8 3NN

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Hillingdon Hospital 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/07/2002		Yes	No