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Glaucoma compliance aids research project

Submission date 28/09/2007	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 28/09/2007	**Overall study status** Completed	
Last Edited 01/09/2015	**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
N0034186696

Study information

Scientific Title
Glaucoma compliance aids research project

Study objectives
1. Do patients receiving glaucoma compliance aids show higher compliance rates with their mediations than those without the aids?
2. Do patients rate the treatment aid as easy to use?

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
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: Glaucoma

Interventions
Glaucoma compliance aids vs no aids.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
1. Improvement of compliance in using glaucoma medications, based on rates of using the daily treatment dose before and after using the glaucoma compliance aid.
2. Ease of using compliance aids, based on patients answers to questions related to: squeezing out drops, getting drops into eye, difficulty in squeezing bottle, difficulty in controlling number of drops, difficulty aiming drops.

Secondary outcome measures

No secondary outcome measures

Overall study start date
01/09/2006

Completion date
01/03/2007

Eligibility

Key inclusion criteria
1. Primary open angle glaucoma patients
2. Patients using prostaglandin analogues anti glaucoma medications only
3. Patients putting eye drops themselves

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
100 patients from Ophthalmology outpatients

Key exclusion criteria
1. Patients with other eye problems
2. Patients using more than one topical medication
3. Patients not able to consent to participate in the project
4. Patients not putting eye drops themselves
5. Patients under the age of 40 years of age

Date of first enrolment
01/09/2006

Date of final enrolment
01/03/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Barnsley Hospital NHS Foundation Trust
Barnsley
United Kingdom
S75 2EP

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
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dhmail@doh.gsi.org.uk

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Barnsley Hospital NHS Foundation 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