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Efficacy, tolerability and safety of 0.2% Brimonidine-Tartrate for the treatment of acute non-arteritic anterior ischaemic optic neuropathy

Submission date
12/09/2003

Recruitment status
No longer recruiting

Registration date
12/09/2003

Overall study status
Completed

Last Edited
04/04/2016

Condition category
Eye Diseases

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] 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
N0123109164

Study information

Scientific Title
Efficacy, tolerability and safety of 0.2% Brimonidine-Tartrate for the treatment of acute non-arteritic anterior ischaemic optic neuropathy (NAION): a 3-month, double-masked, randomised and placebo-controlled trial

Study objectives
To investigate the effect of this medication which is free from harmful side effects in NAION.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Primary study design
Interventional

Study design
Double-masked randomised and placebo-controlled trial

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Eye Diseases: Non-arteritic anterior ischemic optic neuropathy

Interventions
Randomised controlled trial

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Brimonidine-Tartrate

Primary outcome(s)
Safety of 0.2% Brimonidine-Tartrate for the treatment of acute non-arteritic anterior ischemic optic neuropathy (NAION)

Key secondary outcome(s)
Not provided at time of registration

Completion date
30/06/2003

Eligibility

Key inclusion criteria
Patients with acute non-arteritic anterior ischemic optic neuropathy

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/07/2001

Date of final enrolment
30/06/2003

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
University Hospitals of Leicester
Leicester
United Kingdom
LE1 4PW

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
University Hospitals of Leicester NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration