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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

### Study design
Double-masked randomised and placebo-controlled trial

### Primary study design
Interventional

### 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

### Study outputs

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|---|---|---|---|---|---|
| Participant information sheet | Participant information sheet | 11/11/2025 | 11/11/2025 | No | Yes |