

Pharmacological Treatment of Congenital Nystagmus

Submission date 12/09/2003	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 19/04/2007	Condition category Eye Diseases	<input type="checkbox"/> Individual participant data

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
N0123112717

Study information

Scientific Title**Acronym**

PTCN

Study objectives

To investigate whether memantine and gabapentin improve nystagmus.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Leicestershire Ethics Committee

Study design

Placebo controlled 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**Health condition(s) or problem(s) studied**

Nystagmus

Interventions

Administration of gabapentin, memantine or a placebo

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Memantine and gabapentin

Primary outcome measure

Visual acuity

Secondary outcome measures

Nystagmus amplitude

Overall study start date

01/08/2004

Completion date

01/03/2006

Eligibility**Key inclusion criteria**

1. Congenital nystagmus
2. Over 18 years of age

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

48

Key exclusion criteria

1. Other neurological condition
2. Pregnancy or breastfeeding
3. Unable or unwilling to give consent

Date of first enrolment

01/08/2004

Date of final enrolment

01/03/2006

Locations**Countries of recruitment**

England

United Kingdom

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

Sponsor information

Organisation
University Hospitals Leicester (UK)

Sponsor details
Gwendolen House
Gwendolen Road
Leicester
England
United Kingdom
LE5 4QF

Sponsor type
University/education

ROR
<https://ror.org/02fha3693>

Funder(s)

Funder type
Charity

Funder Name
Nystagmus Network UK

Funder Name
Ulverscroft Foundation

Alternative Name(s)

Funding Body Type
Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Funder Name

University Hospitals of Leicester NHS Trust

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	01/02/2007		Yes	No