Antidyskinetic properties of topiramate: a double-blind, placebo-controlled trial in patients with Parkinson's disease and levodopainduced dyskinesias

Submission date	Recruitment status	Prospectively registered
20/05/2008	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
02/07/2008	Completed	Results
Last Edited	Condition category	Individual participant data
09/05/2016	Nervous System Diseases	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

2007 Neuro 12

Study information

Scientific Title

Antidyskinetic properties of topiramate: a double-blind, placebo-controlled trial in patients with Parkinson's disease and levodopa-induced dyskinesias

Study objectives

Levodopa therapy is effective for the motor symptoms of Parkinson's disease. However, around half of patients develop abnormal involuntary movements, or dyskinesia, after 4 - 6 years of treatment. Current treatment interventions for this are not satisfactory in all cases.

Hypothesis:

Topiramate administration will attenuate levodopa-induced dyskinesia in patients with Parkinson's disease (PD) without worsening parkinsonism.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Leeds (West) Research Ethics Committee, 24/01/2008, ref: 07/H1307/205

Study design

Multicentre, randomised, double-blind, placebo-controlled, crossover study

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

Parkinson's disease and levodopa-induced dyskinesia

Interventions

This is a randomised, crossover study.

Topiramate group: start dose 25 mg/day orally (p.o.), to be up-titrated by 25 mg/day weekly to target dose of 100 mg/day in two divided doses. Participants to stay on maintenance dose for two weeks prior to assessment. Participants will attend having not taken usual morning medications, and response to these medications will then be assessed.

Control: placebo capsules, identical in appearance. To be titrated on same schedule as topiramate.

Following a two-week down-titration period, there will be a further two-week washout period before crossover.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Topiramate

Primary outcome measure

Investigator-rated dyskinesia severity. To be scored by a blinded assessor from video recordings of participants, every 30 minutes for a total of 150 minutes. Dyskinesia to be rated at each time-point using a 5-point objective dyskinesia intensity rating scale, rating seven body parts (each limb, face, trunk and neck) with a maximum possible score of 28 at each time point. Timepoints of assessment: baseline (week 0), end of Arm 1 (week 6) and end of Arm 2 (week 16)

Secondary outcome measures

- 1. Investigator-rated parkinsonism. Unified Parkinson's Disease Rating Scale (UPDRS) part III to be assessed at 30-minute intervals during clinical assessment. Timepoints of assessment: week 0, 6 and 16.
- 2. Subject-rated dyskinesia severity:
- 2.1. Lang-Fahn Activities of Daily Living Dyskinesia Scale. Timepoints of assessment: week 0, 6, and 16.
- 2.2. Clinical Global Impression of change, assessed weekly during dose titration and at weeks 6 and 16
- 2.3. UPDRS Part IV. Timepoints of assessment: week 0, 6 and 16.
- 3. Effects on mood and activities of daily living:
- 3.1. UPDRS Part I, II. Timepoints of assessment: week 0, 6 and 16.
- 3.2. Geriatric Depression Scale-15. Timepoints of assessment: week 0, 2, 4, 6, 10, 12, 14 and 16.
- 4. Excessive daytime sleepiness: Epworth Sleepiness Scale. Timepoints of assessment: week 0, 2,
- 4, 6, 10, 12, 14 and 16.

Overall study start date

01/07/2008

Completion date

01/01/2010

Eligibility

Key inclusion criteria

- 1. Both males and females, no age limits
- 2. Patients with Parkinson's disease as defined by UK Parkinson's Disease Society Brain Bank criteria
- 3. Current use of levodopa, dose to be stable for one month prior to enrolment
- 4. Stable levodopa-induced dyskinesias

Participant type(s)

Patient

Age group

Other

Sex

Both

Target number of participants

30

Key exclusion criteria

- 1. Hypersensitivity to topiramate or its excipients
- 2. Prior surgery for PD
- 3. Hoehn and Yahr score of 5 when "off"
- 4. Dementia
- 5. History of nephrolithiasis, renal impairment, liver disease, glaucoma
- 6. Pregnancy and breastfeeding
- 7. Premenopausal females and males not using adequate contraception
- 8. Use of other antiepileptic drugs, carbonic anhydrase inhibitors, metformin, digoxin or illicit drugs

Date of first enrolment

01/07/2008

Date of final enrolment

01/01/2010

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Salford Royal NHS Foundation Trust
Salford
United Kingdom

M6 8HD

Sponsor information

Organisation

Salford Royal NHS Foundation Trust (UK)

Sponsor details

Clinical Sciences Building
Stott Lane
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United Kingdom
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Sponsor type

Hospital/treatment centre

Website

http://www.srht.nhs.uk

ROR

https://ror.org/019j78370

Funder(s)

Funder type

University/education

Funder Name

University of Manchester (UK)

Alternative Name(s)

The University of Manchester, University of Manchester UK, University of Manchester in United Kingdom, UoM

Funding Body Type

Government organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration