A randomised double blind controlled study of the effects of methylphenidate on Central Audition Deficits

Submission date	Recruitment status	Prospectively registered
06/11/2002	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
06/11/2002	Completed	Results
Last Edited	Condition category	Individual participant data
13/08/2009	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

Dr Laurence Jerome

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Acronym

CAD

Study objectives

Not provided at time of registration.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised placebo controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Central Audition Deficit (Central Auditory Processing Disorder [CAPD])

Interventions

Retesting the central audition on methylphenidate or placebo

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Methylphenidate (Ritalin®)

Primary outcome measure

Not provided at time of registration.

Secondary outcome measures

Not provided at time of registration.

Overall study start date

01/04/2002

Completion date

30/04/2003

Eligibility

Key inclusion criteria

Sixty-two consecutive attendees at a university clinic for communication disorders referred for evaluation of learning and listening problems at school by parents, teachers or family physicians and shown to have deficits of Central Audition on a Willeford standardised test battery

Participant type(s)

Patient

Age group

Child

Sex

Both

Target number of participants

62

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/04/2002

Date of final enrolment

30/04/2003

Locations

Countries of recruitment

Canada

Study participating centre

90 Wharncliffe Rd South

London, Ontario Canada N6J 2K1

Sponsor information

Organisation

University of Western Ontario (Canada)

Sponsor details

1151 Richmond Street, Suite 2 London, Ontario Canada N6A 4B8 +1 519 661 2111 wnonline@uwo.ca

Sponsor type

University/education

Website

http://www.med.uwo.ca/

ROR

https://ror.org/02grkyz14

Funder(s)

Funder type

Other

Funder Name

Self funded

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 summaryNot provided at time of registration