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

Recruitment status	Prospectively registered
No longer recruiting	☐ Protocol
Overall study status	Statistical analysis plan
Completed	Results
Condition category	Individual participant data
Nervous System Diseases	Record updated in last year
	No longer recruiting Overall study status Completed Condition category

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Laurence Jerome

Contact details

90 Wharncliffe Rd South London, Ontario Canada N6J 2K1 +1 519 432 3818 ljerome@rogers.com

Additional identifiers

Protocol serial number 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

Study type(s)

Not Specified

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(s)

Not provided at time of registration.

Key secondary outcome(s))

Not provided at time of registration.

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

Healthy volunteers allowed

No

Age group

Child

Sex

All

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)

ROR

https://ror.org/02grkyz14

Funder(s)

Funder type Other

Funder Name Self funded

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration