

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

Submission date 06/11/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 06/11/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 13/08/2009	Condition category Nervous System Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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
90 Wharnccliffe Rd South
London, Ontario
Canada
N6J 2K1
+1 519 432 3818
ljerome@rogers.com

Additional identifiers

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 summary

Not provided at time of registration