

Telehealth adaptation of the Camperdown Program: a randomised controlled trial

Submission date 30/10/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 03/02/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 27/04/2010	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data

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

N/A

Study information

Scientific Title

Study objectives

How many participants will respond to the telehealth adaptation of the Camperdown Program?

Ethics approval required

Old ethics approval format

Ethics approval(s)

La Trobe University (Australia), granted on 23rd May 2005

Study design

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

Chronic stuttering

Interventions

Standard format Camperdown Program versus Telehealth-adapted Camperdown Program

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Percentage of syllables stuttered measured at one day pre-treatment and at 1 day, 6 months, and 12 months post-treatment, as well as 12 months post-randomisation
2. Calculation of mean stuttering
3. Severity scores (SEV scale), reported by the participant in the week prior to collection of each data point

Secondary outcome measures

Time taken to attain minimal stuttering will be evaluated as well as speech rate and speech naturalness of participants post therapy.

Overall study start date

01/04/2005

Completion date

01/03/2008

Eligibility**Key inclusion criteria**

1. Diagnosis of stuttering
2. Stuttering over 2% Syllables Stuttered (SS)
3. Over 18 years of age

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

40 (20 in each program)

Key exclusion criteria

1. Less than 2% SS
2. Unable to attend weekly clinic for three months
3. Received therapy for stuttering in previous 12 months

Date of first enrolment

01/04/2005

Date of final enrolment

01/03/2008

Locations**Countries of recruitment**

Australia

Study participating centre
Australian Stuttering Research Centre (ASRC)
Sydney
Australia
1825

Sponsor information

Organisation
University of Sydney (Australia)

Sponsor details
Australian Stuttering Research Centre
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Sponsor type
University/education

Website
<http://www.fhs.usyd.edu.au/asrc>

ROR
<https://ror.org/0384j8v12>

Funder(s)

Funder type
Research council

Funder Name
Australian National Health and Medical Research Council (NHMRC)

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	01/01/2010		Yes	No