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Telehealth adaptation of the Camperdown Program: a randomised controlled trial

Submission date	Recruitment status
30/10/2005	No longer recruiting
Registration date	**Overall study status**
03/02/2006	Completed
Last Edited	**Condition category**
27/04/2010	Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] 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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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
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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