

Feasibility of the Dutch version of the AB-York Crescent of Sound; a new apparatus for assessing spatial hearing skills in the Dutch population

Published: 11-04-2012

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This study aims to assess the feasibility of the Dutch version of the AB-York Crescent of Sound in normal hearing adults and unilaterally implanted adult CI patients.

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Hearing disorders
Study type	Observational non invasive

Summary

ID

NL-OMON37881

Source

ToetsingOnline

Brief title

FeasCoS2012

Condition

- Hearing disorders

Synonym

deafness, sensory hearing loss

Research involving

Human

Sponsors and support

Primary sponsor: Universitair Medisch Centrum Utrecht

Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: Cochlear Implant users, Normal hearing subjects, Spatial hearing, Speech intelligibility in noise

Outcome measures

Primary outcome

Speech intelligibility in noise (signal noise ratio in dB (SNR)).

Secondary outcome

Spatial hearing (sound localisation)

Study description

Background summary

Normal-hearing listeners gain important benefits from having two ears. Users of a Cochlear Implant (CI) achieve high levels of spoken word recognition when speech is presented in quiet. However, they experience difficulty in the presence of competing sounds and are poor at identifying where sounds come from. At the moment, there is a lack of material to adequately test spatial hearing and speech intelligibility in noise in CI users.

Recently, a new apparatus has been developed by a research group from the University of York. This apparatus is called the 'AB-York Crescent of Sound'. To make the apparatus applicable for the Dutch population of cochlear implant users, the test material has been replaced by Dutch sentences

Study objective

This study aims to assess the feasibility of the Dutch version of the AB-York Crescent of Sound in normal hearing adults and unilaterally implanted adult CI patients.

Study design

This is a cross-sectional study that intends to gather and analyze audiometric

data obtained with a new apparatus for measuring speech in noise and spatial hearing capabilities. All participants will be tested with the new audiometric test set up once.

Study burden and risks

Participation in this study will take some time, but it does not carry any risks.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years)
Elderly (65 years and older)

Inclusion criteria

Normal hearing subjects:

1. Age ≥ 18 and ≤ 70 years.
2. Normal hearing as defined by;
 - a. Pure tone audiometry threshold ≤ 15 dB HL at octave frequencies from 500-4000 Hz.
 - b. Speech intelligibility threshold $\geq 95\%$ at 50 dB SPL
3. Dutch language proficiency.
4. Willingness and ability to participate in all scheduled procedures outlined in the protocol.
5. Written informed consent; Cochlear implantees;
 1. Age ≥ 18 and ≤ 70 years.
 2. Postlingual onset of hearing loss, defined as: the patient attended mainstream education.
 3. At least 1 year of hearing experience with the cochlear implant.
 4. Dutch language proficiency.
 5. Willingness and ability to participate in all scheduled procedures outlined in the protocol.
 6. Written informed consent

Exclusion criteria

1. Disability which could interfere with the completion of the tests.

Study design

Design

Study type:	Observational non invasive
Intervention model:	Other
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active
Primary purpose:	Diagnostic

Recruitment

NL	
Recruitment status:	Recruitment stopped
Start date (anticipated):	12-04-2012
Enrollment:	40
Type:	Actual

Ethics review

Approved WMO

Date: 11-04-2012

Application type: First submission

Review commission: METC Universitair Medisch Centrum Utrecht (Utrecht)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
CCMO	NL39472.041.12

Study results

Date completed: 26-03-2013

Actual enrolment: 37

Summary results

Trial is ongoing in other countries