Bimodal Zoom

No registrations found.

Ethical review	Positive opinion
Status	Other
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON21299

Source NTR

Health condition

Cochlear implant, Hearing aid, Bimodal fitting, Directional microphone

Sponsors and support

Primary sponsor: Maastricht University Medical Center (MUMC+) **Source(s) of monetary or material Support:** Maastricht University Medical Center (MUMC+) Advanced Bionics inc.

Intervention

Outcome measures

Primary outcome

The primary outcome is the effect of microphone configuration on speech recognition performance in noise. Two directional microphone systems will be addressed, a monaural and a binaural system, and compared to the standard omnidirectional microphone setting. The microphone systems will be tested in asymmetric configuration (same setting across ears) and/or asymmetric configuration (different setting across ears).

Secondary outcome

Aside from the primary outcome also listening effort, defined as the effort it takes to listen to speech in noise, will be tested. Furthermore the degree of bimodal benefit will be addressed as well as the effect of noise type, stationary versus fluctuating. Results between the two patient groups, bimodal versus bilateral, will be compared.

Study description

Background summary

Rationale:

Cochlear implantation (CI) has become standard practice to restore hearing in severely hearing-impaired patients by providing multi-channel electrical stimulation to the auditory nerve. Although CI-patients can achieve high levels of speech recognition in guiet, speech recognition in noise still remains one of the most challenging tasks. One way to improve performance in noise is to benefit from binaural hearing (hearing with two ears). For individuals with residual contralateral hearing, a CI in one ear can be combined with an acoustic hearing aid (HA) in the other ear. This is known as bimodal hearing. For individuals without residual hearing a second CI can be an opportunity in the rare case. This is known as bilateral hearing. A second way to improve speech recognition in noise is to improve the quality of the signal before it is offered to the ear. To that end directional microphone systems are designed as they focus on the speech signal in front and reduce the noise from other directions. Nowadays,

directional microphone algorithms are available for HA's as well as for CI's. Both approaches (binaural hearing and directional systems) are considered complementary, however they are not yet evaluated conjointly.

Objective:

Evaluate the performance of directional microphone systems in binaural (bimodal and bilateral) cochlear implant users

Study population:

One group consists of users of a cochlear implant (CI) in one ear and a conventional hearing aid (HA) in the other ear. The other group consist users of a cochlear implant (CI) in both ears.

Study design:

A cross-over repeated measures design is carried out to single-blind evaluate the performance of directional microphone systems. During two test sessions bimodal subjects are provided with the latest speech processor for the CI ear and a state-of- the-art hearing aid in the other ear. The bilateral group will be provided with the latest CI speech processor in both ears during a single test session. Both CI and HA devices allow different microphone configurations: standard omnidirectional processing and directional multimicrophone processing in each ear separately (monaural) or combined cross ears (binaural). For each directional setting, speech perception in noise is assessed using two different masking materials (stationary noise versus fluctuating talker).

Primary study parameter:

The primary outcome is the effect of microphone configuration on speech recognition performance in noise.

Secondary study parameters:

Secondary outcomes in this study are bimodal benefit, the

effect of masker type and listening effort. Results between

the two patient groups, bimodal versus bilateral, will be

compared.

Nature and extent of the burden and risks associated with

participation, benefit and group relatedness:

There are no known health risks associated with

participation in this study. CE-marked hearing devices (CI and HA) are used within the scope of standard care.

Participation however takes time, effort and attention from

subjects. As a result of the study subjects can be advised

towards the use of a directional microphone setting to

improve their speech recognition performance in noise.

Amendments 23-aug-2016

Study objective

For cochlear implanted (CI) patients speech recognition in noise still remains one of the most challenging tasks. One way to improve performance in noise is to benefit from binaural hearing (hearing with two ears). For individuals with residual contralateral hearing, a CI in one ear can be combined with an acoustic hearing aid (HA) in the other ear, known as bimodal hearing. For individuals without residual hearing a second CI can be an opportunity in the rare case. This is known as bilateral hearing.

A second way to improve speech recognition in noise is to

improve the quality of the signal before it is offered to the

ear. To that end directional microphone systems are

designed as they focus on the speech signal in front and

reduce the noise from other directions. Nowadays,

directional microphone algorithms are available for HA's as

well as for CI's. Both approaches (binaural hearing and

directional systems) are considered complementary, however they are not yet evaluated conjointly.

Therefore the hypothesis assessed in this study is that

directional microphone systems can improve speech

recognition performance in binaural (bimodal or bilateral)

cochlear implant users.

Study design

NA

Intervention

During two test session bimodal subjects are provided with the latest speech processor for the CI ear and a state-of- the-art hearing aid in the other ear. The bilateral group will be provided with the latest CI speech processor in both ears during a single test session. Both CI and HA devices allow different microphone configurations: standard omnidirectional processing and directional multi-microphone processing in each ear separately (monaural) or combined cross ears (binaural).

Contacts

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Eligibility criteria

Inclusion criteria

Bimodal Group:

- 1. capacitated adult (>18 years of age)
- 2. patient of CI-team South-East Netherlands
- 3. user of a unilateral cochlear implant (CI) of the brand Advanced Bionics (AB)
- 4. first fit CI >= 6 months ago
- 5. wearing CI speech processor (almost) always (i.e. circa 10 hours a day)

6. wearing a contralateral hearing aid >50% of the time (i.e. circa 5 hours a day)

7. able to perform the speech-in-noise test (i.e. speech recognition in quiet <50%)

8. willing and able to visit hospital and participate in testing

9. agreed to participate in this study (by informed consent)

Bilateral group:

- 1. capacitated adult (>18 years of age)
- 2. patient of CI-team azM, RadboudUMC or UMCU

3. Former subject in the study NL24660.018.08/NTR1722 who completed the full fuollow-up period of four years since first implantation

4. user of a bilateral cochlear implants (CI's) of the brand Advanced Bionics (AB)

- 5. first fit of second Cl >= 6 months ago
- 6. wearing CI speech processor in both ears (almost) always (i.e. circa 10 hours a day)
- 7. able to perform the speech-in- noise test (i.e. speech recognition in quiet >50%)
- 8. willing and able to visit hospital and participate in testing

9. agreed to participate in this study (by informed consent) And additionally in case of patient Radboud UMC or

UMCU:

1. Agreed to let research team inform own CI-team of participation in current study (by informed consent)

2. Agreed to let research team retrieve basic audiological information from own CI-team (by informed consent)

Exclusion criteria

Bimodal group:

- 1. non fluent in Dutch
- 2.<18 years of age or incapacitated
- 3. bilateral cochlear implant user (CI+CI)

Bilateral group:

- 1. non fluent in Dutch
- 2. <18 years of age or incapacitated

Study design

Design

Study type:	Interventional
Intervention model:	Crossover
Allocation:	Non controlled trial
Masking:	Single blinded (masking used)
Control:	N/A , unknown

Recruitment

NL	
Recruitment status:	Other
Start date (anticipated):	01-01-2015
Enrollment:	24
Туре:	Unknown

IPD sharing statement

Plan to share IPD: Undecided

Ethics review

Positive opinion	
Date:	13-11-2014
Application type:	First submission

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

RegisterIDNTR-newNL4696NTR-oldNTR4901OtherMETC azZM/UM : 141130

Study results

Summary results N/A