# The impact of aging, cognition, and sensory function on speech

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The aim of this study is to better understand the relation between cognition, sensory functions and motor control for speech in older age. Thereby, we strive for improved speech-based discrimination of typical aging from neurodegenerative declines.

Ethical review Approved WMO

**Status** Pending

**Health condition type** Other condition

**Study type** Observational non invasive

## **Summary**

## ID

NL-OMON57444

#### Source

**ToetsingOnline** 

#### **Brief title**

Speech changes in typical aging and neurodegenerative diseases

## **Condition**

- Other condition
- Movement disorders (incl parkinsonism)

#### Synonym

dysarthria, Pathological speech

#### **Health condition**

neurodegenerative disorders

## **Research involving**

Human

## **Sponsors and support**

**Primary sponsor:** Rijksuniversiteit Groningen

Source(s) of monetary or material Support: NWO

## Intervention

**Keyword:** aging, Mild Cognitive Impairment, Parkinson's disease, speech motor control

## **Outcome measures**

#### **Primary outcome**

The most important research parameters are acoustic changes of speech in response to feedback perturbations and kinematic measures of breathing.

## **Secondary outcome**

The secondary study endpoints are acoustic measurements of voice pitch, acoustic characteristics of articulation during unperturbed speech and subjective speech complaints.

# **Study description**

## **Background summary**

Previous research suggests that speech motor control for pitch declines as we age. However, as voice-level control differs from articulatory-level and breathing control, it is unclear how aging affects speech motor control for articulation and respiration. While such changes do usually not lead to speech problems in typical older adults, declining speech motor control may cause speech impairments in neurodegenerative diseases such as Parkinson\*s and Alzheimer\*s disease. The underlying mechanisms of those speech impairments are unclear. While people with Parkinson\*s Disease primarily show sensory but also some cognitive impairments, people with Alzheimer\*s Disease primarily show cognitive but also some sensory impairments. Yet, the relation of cognitive and sensory decline, and declining speech motor control remains to be clarified.

## **Study objective**

The aim of this study is to better understand the relation between cognition,

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sensory functions and motor control for speech in older age. Thereby, we strive for improved speech-based discrimination of typical aging from neurodegenerative declines.

## Study design

The study is designed as a cross-sectional observational study. The participants will complete several tasks during two experimental sessions. They will complete behavioural tasks targeting (1) various cognitive abilities, and (2) auditory as well as somatosensory perception. Furthermore, they will complete (3) speech tasks, while being recorded acoustically. At the same time, respiratory patterns will be recorded through electromagnetic articulography (EMA).

## Study burden and risks

Participation in this study is not related to any known risks or benefits. A slight burden could be fatigue due to the duration of the experiment (appr. 2 hours per session). To reduce this burden, tasks are varied, and breaks are included in the experimental procedure. Breaks will also be added or prolonged on the participants\* request. Moreover, the session has been split into two sessions in order to reduce the burden for participants. For the behavioural assessment of sensory functions, sanitised plastic domes will be gently pressed against participants\* tongue and lips. For several tasks, participants will wear a microphone and headphones to listen to sounds or their own voice in a regular, conversational amplitude. For the speech tasks specifically, EMA sensors will be attached to their chest, the back (i.e., spine), and (if possible) the abdomen in order to record respiratory and articulatory kinematics. Those sensors are easily removable. If participants with Parkinson\*s disease have undergone Deep Brain Stimulation (DBS) in the past, they cannot participate in the study due to the EMA recordings which rely on an electromagnetic field.

## **Contacts**

#### **Public**

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## **Scientific**

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## **Trial sites**

## **Listed location countries**

**Netherlands** 

# **Eligibility criteria**

## Age

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

## Inclusion criteria

- Native speaker of Dutch

Only for the group of people with a sensorimotor disorder:

- Diagnosed with Parkinson's disease

Only for the group of people with a cognitive disorder:

- Diagnosed with mild cognitive impairment

Only for the group of control participants:

- No prior speech or language impairments

## **Exclusion criteria**

- Lack of ability to give informed consent (i.e., incapacitation) in any case of doubt, the (in-)capacitation will be determined by a physician
- History of stroke or other neurological or psychological disorder
- Self-reported signs of depression
- Stuttering or other speech and language problems that are not related to PD or  $\mbox{MCI}$
- Severe impairment of vision
- Metal at or close to the head, chest or abdomen (e.g., piercings, pacemaker, etc.)

- History of Deep Brain Stimulation (DBS)

# 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: Pending

Start date (anticipated): 01-12-2024

Enrollment: 120

Type: Anticipated

## **Ethics review**

Approved WMO

Date: 25-04-2025

Application type: First submission

Review commission: METC Universitair Medisch Centrum Groningen (Groningen)

# **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 NL87470.042.24