

# Enhanced Neural Speech Tracking in Aging and Hearing Loss: The Role of Stochastic Resonance

Björn Herrmann

ORCID: 0000-0001-6362-3043

Rotman Research Institute,  
Baycrest Academy for Research and Education, M6A 2E1, North York, ON, Canada

Department of Psychology,  
University of Toronto, M5S 1A1, Toronto, ON, Canada

Correspondence concerning this article should be addressed to Björn Herrmann, Rotman Research Institute, Baycrest, 3560 Bathurst St, North York, ON, M6A 2E1, Canada. E-mail: [bherrmann@research.baycrest.org](mailto:bherrmann@research.baycrest.org)

**Keywords:** Electroencephalography, aging, hearing loss speech encoding, story listening, temporal response function, background noise

**Conflict of interest statement:** The author declares no competing interests.

**Acknowledgments:** I thank Priya Pandey and Saba Junaid for their help with story generation and data collection. The research was supported by the Canada Research Chair program (CRC-2019-00156) and the Natural Sciences and Engineering Research Council of Canada (Discovery Grant: RGPIN-2021-02602).

**Author contributions: Björn Herrmann:** Conceptualization, methodology, formal analysis, investigation, data curation, writing - original draft, writing - review and editing, visualization, supervision, project administration, funding acquisition.

**Word count:** Abstract (239), Significance Statement (120), Introduction (717), Discussion (1645)

**Figure/table count:** Figures (7), Tables (0)

### Abstract

Tracking the envelope of speech in the brain is important for speech comprehension. Recent research suggests that acoustic background noise can enhance neural speech tracking, enabling the auditory system to robustly encode speech even under unfavorable conditions. Aging and hearing loss are associated with internal, neural noise in the auditory system, which raises the question whether additional acoustic background noise can enhance neural speech tracking in older adults. In the current electroencephalography study, younger (~25.5 years) and older adults (~68.5 years) listened to spoken stories either in quiet (clear) or in the presence of background noise at a wide range of different signal-to-noise ratios. In younger adults, neural speech tracking was enhanced by minimal background noise, indicating the presence of stochastic resonance, that is, the response facilitation through noise. In contrast, older adults, compared to younger adults, showed enhanced neural speech tracking for clear speech and speech masked by minimal background noise, but the acoustic noise led to little enhancement in neural tracking in older people. The data demonstrate different sensitivity of the auditory cortex to speech masked by noise between younger and older adults. The results are consistent with the idea that the auditory cortex of older people exhibits more internal, neural noise that enhances neural speech tracking – through stochastic resonance – but that additional acoustic noise does not further support speech encoding. The work points to a highly non-linear auditory system that differs between younger and older adults.

35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54

55 **Significance statement**

56 Acoustic background noise can enhance neural speech tracking in younger adults to facilitate  
57 robust speech encoding in unfavorable situations. Aging and hearing loss increase neural noise,  
58 potentially making the auditory system less sensitive to acoustic noise. Here, younger and older  
59 adults listened to spoken stories in quiet or background noise while electroencephalography  
60 was recorded. Neural speech tracking was larger for older than younger adults for speech in  
61 quiet and under minimal background noise. However, noise enhanced neural tracking only for  
62 younger, but not older adults. The results support the idea that the auditory cortex of older  
63 people exhibits more neural noise that enhances neural speech tracking through stochastic  
64 resonance, but that additional acoustic noise does not further amplify speech encoding.

65

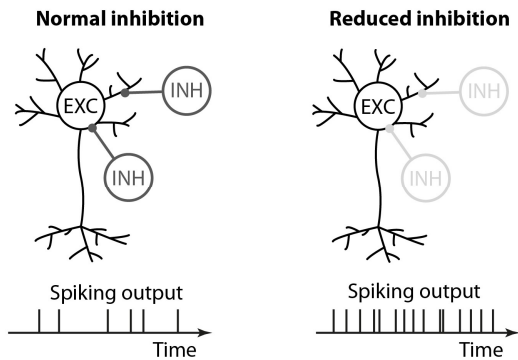
## Introduction

66

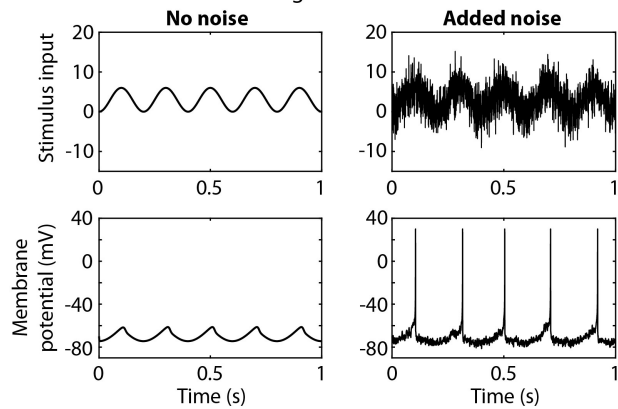
67 Many older adults live with some form of hearing loss (Feder et al., 2015; Goman and Lin, 2016)  
68 that leads to difficulties comprehending speech in the presence of background noise, such as in  
69 crowded places (Pichora-Fuller et al., 2016; Herrmann and Johnsrude, 2020). Understanding  
70 how speech in noisy situations is encoded in the brains of older people is critical for developing  
71 effective treatments for speech comprehension challenges.

72 Much research has focused on how the auditory cortex tracks the envelope of speech  
73 (Lalor and Foxe, 2010; Ding et al., 2014; Ding and Simon, 2014; Brodbeck and Simon, 2020),  
74 because accurate envelope encoding is thought to facilitate speech understanding (Rosen,  
75 1992; Shannon et al., 1995; Ding et al., 2014; Vanthornhout et al., 2018; Lesenfants et al., 2019).  
76 However, recent works suggest non-linearities in how neural speech tracking is affected by  
77 different levels of background noise (Yasmin et al., 2023; Herrmann, 2024; Panela et al., 2024).  
78 Neural speech tracking exhibits an inverted u-shaped profile, where tracking is highest for  
79 moderate signal-to-noise ratios (SNRs) that are associated with intelligible, but challenging  
80 speech, whereas neural tracking decreases for more unfavorable SNRs (poor intelligibility) and  
81 more favorable SNRs (high intelligibility; Yasmin et al., 2023; Herrmann, 2024). Attentional effort  
82 required to understand speech at moderate SNRs has been suggested to lead to the neural-  
83 tracking enhancement (Hauswald et al., 2022; Yasmin et al., 2023; Panela et al., 2024), but  
84 recent work demonstrates little impact of attention on the inverted u-shape (Herrmann, 2024).  
85 Instead, it was suggested that noise per se increases neural speech tracking at moderate SNRs,  
86 and that tracking only decreases when speech intelligibility significantly declines for unfavorable  
87 SNRs (Herrmann, 2024). Stochastic resonance – the response facilitation through noise  
88 (McDonnell and Abbott, 2009; McDonnell and Ward, 2011; Krauss et al., 2016) – was proposed  
89 as the critical mechanism that leads to the neural tracking enhancement (Herrmann, 2024;  
90 Figure 1).

**A** Schematic of microcircuit with different levels of inhibition



**B** Stochastic resonance: Single neuron simulation



**Figure 1: Schematic of increased neural noise due to a loss of neural inhibition and simulation of the impact of increased neural noise on neural outputs.**

**A:** Schematic of an auditory microcircuit with normal and reduced inhibition (INH), regulating the spiking output of an excitatory neuron (EXC). Reduced inhibition in auditory cortex is associated with aging and hearing loss (Ouellet and de Villers-Sidani, 2014; Salvi et al., 2017; Herrmann and Butler, 2021), leading to less regulated – more noisy – spiking output. **B:** Visualization of stochastic resonance (McDonnell and Abbott, 2009; McDonnell and Ward, 2011; Krauss et al., 2016) through the simulation of the membrane potential of a single neuron driven by a periodic (5 Hz) stimulus input. An integrate-and-fire neuron model was used (Izhikevich, 2003, 2004). Periodic stimulation in the absence of noise (top left) does not elicit spikes (i.e., action potentials; bottom left). Periodic stimulation in the presence of background noise (top right) leads to spiking activity at the periodicity of the stimulus input (bottom right). This response facilitation through noise is referred to as stochastic resonance (McDonnell and Abbott, 2009; McDonnell and Ward, 2011; Krauss et al., 2016).

91 Aging and hearing loss are associated with a loss of neural inhibition and an increase in  
92 neural excitation in auditory cortex, resulting from reduced inputs to the neural pathway caused  
93 by peripheral damage (Caspary et al., 2008; Ouellet and de Villers-Sidani, 2014; Zhao et al.,  
94 2016; Resnik and Polley, 2017; Salvi et al., 2017; Herrmann and Butler, 2021; McClaskey, 2024).  
95 A loss of inhibition and increased excitation can manifest as hyperresponsivity to sound  
96 (Auerbach et al., 2014; Chambers et al., 2016; Salvi et al., 2017). Consistently, the neural  
97 tracking of the speech envelope is enhanced in older compared to younger adults (Presacco et  
98 al., 2016a, b; Brodbeck et al., 2018; Decruy et al., 2019; Broderick et al., 2021; Panella et al.,  
99 2024), highlighting the impact on the encoding of relevant features of speech. Reduced  
100 inhibition and increased excitation also increase spontaneous activity – and thus neural noise –  
101 in the absence of sound (Kaltenbach and Afman, 2000; Eggermont and Roberts, 2004;  
102 Eggermont, 2015; Parthasarathy et al., 2019). Neural noise – although difficult to observe directly  
103 in humans using non-invasive recording techniques, such as electroencephalography (EEG) –  
104 could drive the age-related enhancement of neural speech tracking through stochastic  
105 resonance (for discussions of the role of stochastic resonance in hearing loss see Krauss et al.,

106 2016; Schilling et al., 2023). Some neurons may receive insufficient input to elicit a response  
107 when an individual listens to clear speech but may be pushed beyond their firing threshold by  
108 acoustically elicited neural noise (e.g., younger) or intrinsic neural noise (e.g., older) in the  
109 auditory system (Figure 1B). Critically, a neural system with increased internal, neural noise  
110 (e.g., older) may show reduced sensitivity to external, acoustic noise than a system with lower  
111 neural noise (e.g., younger), because the neural distortions introduced through acoustic noise  
112 may not necessarily add further output amplification to a system that is already amplified  
113 through stochastic resonance. However, whether the auditory system of older adults enhances  
114 neural speech tracking through acoustic noise is unknown.

115 The current study uses EEG to investigate in younger and older adults how neural speech  
116 tracking is affected by background noise ranging from very high (i.e., intelligible) to low SNRs  
117 (i.e., less intelligible). Enhanced speech tracking for speech in quiet and a reduced sensitivity of  
118 neural speech tracking to background noise for older compared to younger adults would  
119 indicate that changes in the aged auditory system, such as increased neural noise, reduces  
120 stochastic resonance driven by external, acoustic noise.

## 121 **Methods and materials**

### 122 **Participants**

123 Twenty-six younger adults (median: 25.5 years; range: 18–34 years; 8 male or man, 16 female or  
124 woman, 1 transgender man, 1 non-binary) and 26 older adults (median: 68.5 years; range: 57–78  
125 years; 9 male or man, 17 female or woman) participated in the current study. Participants were  
126 native English speakers or grew up in English-speaking countries (mostly Canada) and have  
127 been speaking English since early childhood (<5 years of age). Participants reported having  
128 normal hearing abilities and no neurological disease (one person reported having ADHD, but this  
129 did not affect their participation). Participants gave written informed consent prior to the  
130 experiment and were compensated for their participation. The study was conducted in  
131 accordance with the Declaration of Helsinki, the Canadian Tri-Council Policy Statement on  
132 Ethical Conduct for Research Involving Humans (TCPS2-2014), and was approved by the  
133 Research Ethics Board of the Rotman Research Institute at Baycrest Academy for Research and  
134 Education.

### 135 **Acoustic environment and stimulus delivery**

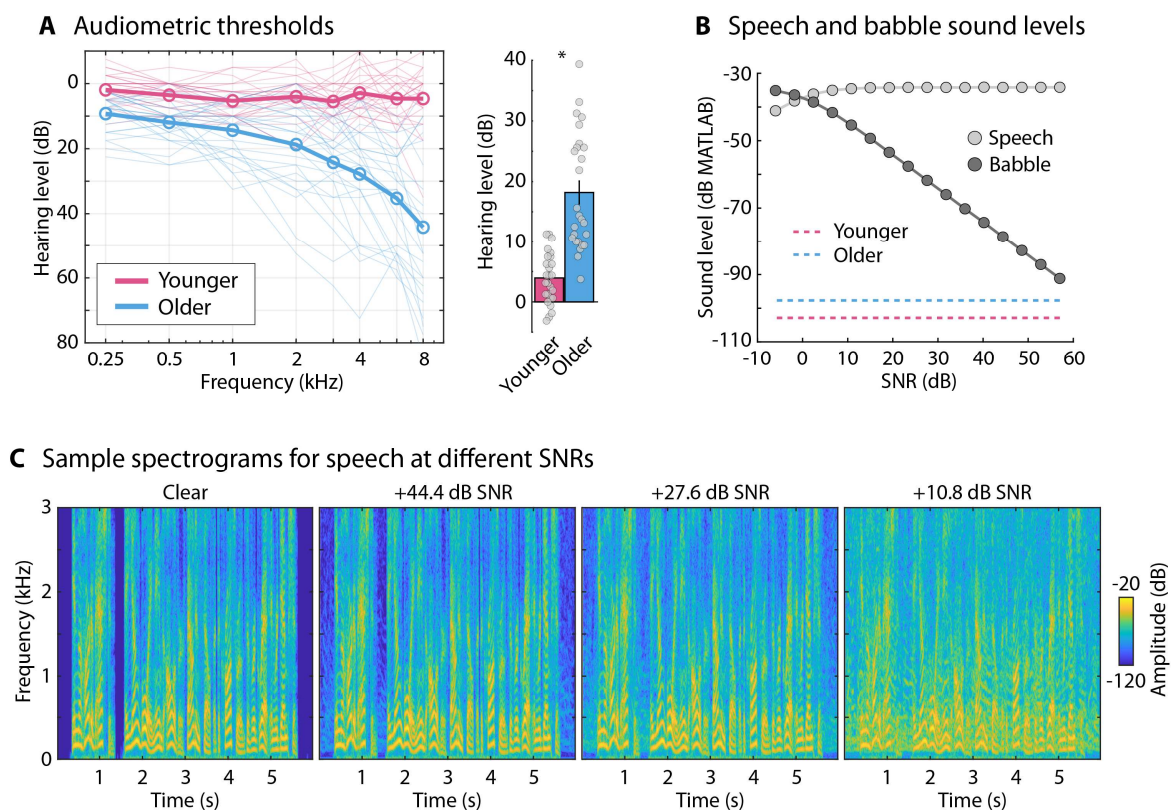
136 Data were gathered in a sound-attenuating booth to reduce external sound interference. Sounds  
137 were delivered using Sennheiser HD 25-SP II headphones connected via an RME Fireface 400



138 audio interface. The experiment was implemented using Psychtoolbox (version 3.0.14) running  
139 in MATLAB (MathWorks Inc.) on a Lenovo T480 laptop with Windows 7. Visual stimuli were  
140 projected into the booth via a mirrored display. Auditory stimuli were played at approximately 70  
141 dB SPL.

## 142 Hearing assessment

143 Pure-tone audiometry was administered for each participant at frequencies of 0.25, 0.5, 1, 2, 3,  
144 4, 6, and 8 kHz. Pure-tone average thresholds (PTA: average across 0.5, 1, 2, and 4 kHz; Stevens  
145 et al., 2013; Humes, 2019) were higher for older compared to younger adults ( $t_{50} = 6.893$ ,  $p = 8.8$   
146  $\cdot 10^{-9}$ ,  $d = 1.912$ ; Figure 2A). Elevated thresholds are consistent with the presence of mild-to-  
147 moderate hearing loss in the current sample of older adults, as would be expected (Moore,  
148 2007; Plack, 2014; Presacco et al., 2016a; Herrmann et al., 2018, 2022). A few older adults of the  
149 current sample also appeared to have ‘clinical’ hearing loss as indicated by thresholds above 20  
150 dB HL (Stevens et al., 2013; Humes, 2019), but none of them were prescribed with hearing aids.  
151 Although the main analyses focus on the originally intended comparisons of younger and older  
152 adults, in explorative analyses, data were analyzed separately for older adults with clinically  
153 normal hearing (PTA < 20 dB HL) and those with hearing impairment (PTA > 20 dB HL).



**Figure 2: Audiograms, stimulus sound levels, and sample spectrograms. A:** Left: Pure-tone

audiometric thresholds for younger and older adults. Circles and thick lines reflect the mean across participants. Thin lines are the thresholds for each participant. Right: Pure-tone average threshold (PTA; across 0.5, 1, 2, and 4 kHz). Bars reflect the mean across participants. Error bars reflect the standard error of the mean. Dots reflect the PTAs for individual participants. **B:** Sound levels of the speech and background babble for each signal-to-noise ratio (SNR) used in the current study. Sound levels are provided in dB based on MATLAB calculations. More negative values reflect softer sound intensities. Values can be interpreted relative to each other, whereas the absolute magnitude is related to hardware and software conditions, such as sound card, transducers, and MATLAB internal settings. The colored dashed lines show the mean sensation level for a babble noise stimulus for both age groups. **C:** Sample spectrograms for the first 6 seconds of one story under different speech-clarity conditions (clear, 44.4, 27.6, and 10.8 dB SNR). Note that the magnitudes in panels B and C are not comparable.

154 In order to obtain a reference threshold in MATLAB software for speech and babble  
155 presentation during the main experimental procedures, the sensation level for a 12-talker  
156 babble noise was estimated using a method-of-limits procedure (Leek, 2011; Herrmann and  
157 Johnsrude, 2018; Herrmann et al., 2022). Participants listened to a 14-s babble noise that  
158 changed continuously in intensity at a rate of 5.4 dB/s (either decreased [i.e., starting at  
159 suprathreshold levels] or increased [i.e., starting at subthreshold levels]). Participants pressed a  
160 button when they could no longer hear the tone (intensity decrease) or when they started to hear  
161 the tone (intensity increase). The sound stopped after the button press. The sound intensity at  
162 the time of the button press was noted for 6 decreasing sounds and 6 increasing sounds  
163 (decreasing and increasing sounds alternated), and these were averaged to determine the  
164 sensation level. Due to technical issues this threshold was only available for 21 younger and 25  
165 older adults. As expected, given the audiometric pure-tone average thresholds (Figure 2A),  
166 sensation levels for the babble noise were elevated for older compared to younger adults ( $t_{44} =$   
167 2.573,  $p = 0.014$ ,  $d = 0.762$ , mean difference: 5.2 dB; Figure 2B).

## 168 **Story materials**

169 Participants listened to 20 unique audio stories, each with a duration between 1.5 to 2.5  
170 minutes. These stories were crafted using OpenAI's GPT-3.5 (OpenAI et al., 2023), which also  
171 generated four comprehension questions per story, alongside four answer options (one correct,  
172 three distractors). The themes varied widely across stories, encompassing scenarios such as  
173 making an unexpected friendship on a plane, a boy finding a knitting talent, and a linguist  
174 deciphering ancient text. To ensure high quality of both the content and questions, the AI-  
175 generated materials underwent manual verification. Google's AI-based text-to-speech  
176 synthesizer was employed to produce the auditory version of the stories, using the male English-  
177 speaking voice "en-US-Neural2-J" with default settings ([https://cloud.google.com/text-to-](https://cloud.google.com/text-to-speech/docs/voices)  
178 [speech/docs/voices](https://cloud.google.com/text-to-speech/docs/voices)).



179 Participants listened to the 20 stories in 5 blocks, each comprising 4 stories. Four of the  
180 20 stories were played under clear conditions, that is, in quiet. Twelve-talker babble was added  
181 to the other 16 stories (Bilger, 1984; Bilger et al., 1984; Wilson et al., 2012). Twelve-talker babble  
182 simulates a crowded restaurant, but does not permit identifying individual words in the masker  
183 (Mattys et al., 2012). The babble masker was added at SNRs ranging from +57 to -6 dB in 16  
184 steps of 4.2 dB SNR (Figure 2B, C). Speech in background babble above +15 dB SNR is highly  
185 intelligible (Holder et al., 2018; Rowland et al., 2018; Spyridakou et al., 2020; Irsik et al., 2022),  
186 and listeners had no trouble understanding speech at the highest SNRs used in the current  
187 study. All stimuli (clear speech; the mixed speech and babble signals) were normalized to the  
188 same root-mean-square amplitude and presented at about 70 dB SPL. Figure 2B shows that for  
189 high SNRs, the root-mean-square amplitude of the speech signal in the sound mixture remained  
190 relatively constant, because the babble at high SNRs had little impact on the root-mean-square  
191 amplitude of the sound mixture. Assignment of speech-clarity levels (clear speech and SNRs) to  
192 specific stories was randomized for each participant.

193 After each story, participants rated two statements regarding their speech  
194 comprehension using a 9-point scale (1 = strongly disagree, 9 = strongly agree): ‘I understood the  
195 gist of the story’ and ‘I was able to comprehend the speech well’. They were instructed to rate  
196 the statements independently from other stories they had heard. Ratings were linearly  
197 normalized to a 0 to 1 scale for statistical purposes, making them comparable to proportion-  
198 correct measures (Herrmann, 2024; Mathiesen et al., 2024; Panella et al., 2024). Such gist  
199 ratings have previously been shown to strongly correlate with word-report speech intelligibility  
200 measures (Davis and Johnsruide, 2003; Ritz et al., 2022). The ratings of the two statements were  
201 averaged to obtain one comprehension rating per story and participant. After rating the two  
202 statements, participants answered four multiple-choice questions about the content of the  
203 story. The comprehension questions offered four answer choices (chance level of 25%). The  
204 proportion of correct answers was calculated.

## 205 **Electroencephalography (EEG) acquisition and preprocessing**

206 A BioSemi system (BioSemi, Netherlands) was used to record electroencephalographic data  
207 from 16 Ag/Ag-Cl electrodes (10-20 system) and two additional electrodes, one positioned on  
208 the left and one on the right mastoid. Data were recorded at a 1024 Hz sampling rate and with a  
209 208 Hz online low-pass filter. Reference electrodes were part of the BioSemi CMS-DRL (common  
210 mode sense-driven right leg) system for optimal referencing and noise reduction.

211 Offline processing was performed in MATLAB. A 60-Hz elliptic notch filter was used to  
212 reduced power-line noise. EEG signals were re-referenced to the average of the left and right  
213 mastoid electrodes, which enhances auditory responses at fronto-central electrodes (Ruhnau et

214 al., 2012; Herrmann, 2024). EEG data were high-pass filtered at 0.7 Hz (length: 2449 samples,  
215 Hann window) and low-pass filtered at 22 Hz (length: 211 samples, Kaiser window, window  
216 parameter 4). The data were time-locked to the onset of each story, downsampled to 512 Hz,  
217 and subjected to an Independent Component Analysis (ICA) to remove blink and eye movement  
218 artifacts (Bell and Sejnowski, 1995; Makeig et al., 1995; Oostenveld et al., 2011). Signal  
219 segments showing fluctuations greater than 80  $\mu\text{V}$  within a 0.2-second window in any EEG  
220 channel were set to 0  $\mu\text{V}$  to remove artifacts not removed by the ICA (cf. Dmochowski et al.,  
221 2012; Dmochowski et al., 2014; Cohen and Parra, 2016; Irsik et al., 2022; Yasmin et al., 2023;  
222 Panela et al., 2024). Finally, EEG data were further low-pass filtered at 10 Hz (251 points, Kaiser  
223 window) because neural signals in the low-frequency range are most sensitive to the speech  
224 envelope (Luo and Poeppel, 2007; Di Liberto et al., 2015; Zuk et al., 2021; Karunathilake et al.,  
225 2023; Synigal et al., 2023; Yasmin et al., 2023).

#### 226 **Calculation of amplitude-onset envelopes**

227 Each story's audio signal (devoid of background noise) was processed through a basic auditory  
228 model, which included 30 cochlear-like auditory filters and cochlear compression by a factor of  
229 0.6 (McDermott and Simoncelli, 2011). The resulting 30 envelopes were averaged and smoothed  
230 with a 40-Hz low-pass filter (Butterworth, 4<sup>th</sup> order). Such a computationally simple peripheral  
231 model has been shown to be sufficient, as compared to complex, more realistic models, for  
232 envelope-tracking approaches (Biesmans et al., 2017). The amplitude-onset envelope was  
233 computed since it elicits strong neural speech tracking (Hertrich et al., 2012) and was used in  
234 the previous studies in younger adults that showed noise-related enhancements in neural  
235 speech tracking (Yasmin et al., 2023; Herrmann, 2024; Panela et al., 2024). The amplitude-onset  
236 envelope was obtained by calculating the first derivative of the averaged amplitude envelope  
237 and subsequently setting negative values to zero (Hertrich et al., 2012; Fiedler et al., 2017;  
238 Daube et al., 2019; Fiedler et al., 2019; Yasmin et al., 2023; Panela et al., 2024). It was then  
239 downsampled to match the EEG data's temporal resolution and transformed to z-scores  
240 (subtraction by the mean and division by the standard deviation).

#### 241 **EEG analysis: Temporal response function and prediction accuracy**

242 The relationship between EEG signals and auditory stimuli was assessed using a linear temporal  
243 response function (TRF) model (Crosse et al., 2016; Crosse et al., 2021). Ridge regression with a  
244 regularization parameter of  $\lambda = 10$  was applied based on previous work (Fiedler et al., 2017;  
245 Fiedler et al., 2019; Yasmin et al., 2023; Panela et al., 2024). Pre-selection of  $\lambda$  based on previous  
246 work avoids extremely low and high  $\lambda$  on some cross-validation iterations and avoids

247 substantially longer computational time. Pre-selection of  $\lambda$  also avoids issues if limited data per  
248 condition are available, as in the current study (Crosse et al., 2021).

249 For each story, 50 random 25-second segments of the EEG data were extracted and  
250 paired with corresponding segments of the amplitude-onset envelope. A leave-one-out cross-  
251 validation approach was employed, with one segment reserved for testing and the other non-  
252 overlapping segments used to train the TRF model for lags ranging from 0 to 0.4 s. The model's  
253 performance was evaluated by correlating the predicted EEG signals with the actual EEG in the  
254 test segment, and this procedure was repeated across all 50 segments to derive the mean  
255 prediction accuracy. Overlapping segments were used to increase the amount of data for  
256 training given the short duration of the stories (Herrmann, 2024). Critically, speech-clarity levels  
257 were randomized across stories and analyses were the same for all conditions. Hence, no  
258 impact of overlapping training data on the results is expected (consistent with noise-related  
259 enhancements observed previously when longer stories and non-overlapping data were used;  
260 Yasmin et al., 2023).

261 To investigate the neural-tracking response amplitude, TRFs for each training dataset  
262 were calculated for lags ranging from -0.15 to 0.5 s. Baseline correction was performed by  
263 subtracting the mean signal from -0.15 to 0 seconds from the TRF data at each time point.  
264 Analysis concentrated on the fronto-central electrodes (F3, Fz, F4, C3, Cz, C4), which are known  
265 to reflect auditory cortical activity (Näätänen and Picton, 1987; Picton et al., 2003; Herrmann et  
266 al., 2018; Irsik et al., 2021). Key metrics were the P1-N1 and P2-N1 amplitude differences. To this  
267 end, the P1, N1, and P2 latencies were estimated for each SNR from the averaged time courses  
268 across participants (separately for each group). P1, N1, and P2 amplitudes were calculated for  
269 each participant and condition as the mean amplitude in the 0.02 s time window centered on the  
270 peak latency. The P1-minus-N1 and P2-minus-N1 amplitude differences were calculated. The  
271 amplitude of individual TRF components (P1, N1, P2) was not analyzed because the TRF time  
272 courses for the clear condition had an overall positive shift (see also Herrmann, 2024; Panela et  
273 al., 2024) that could bias analyses more favorably towards response differences which may,  
274 however, be harder to interpret. The P1-N1 amplitude is of particular interest in the current  
275 study, because this early auditory response has previously shown the response amplification  
276 through background noise (Yasmin et al., 2023; Herrmann, 2024; Panela et al., 2024).

## 277 **Statistical analyses**

278 Behavioral data (comprehension accuracy, comprehension ratings), TRFs, and EEG prediction  
279 accuracy for the four clear stories were averaged. For the stories in babble, a sliding average  
280 across SNR levels was calculated for behavioral data, TRFs, and EEG prediction accuracy, such  
281 that data for three neighboring SNR levels were averaged to reduce noise in the data.

282 For the statistical analyses of behavioral data (comprehension accuracy, comprehension  
283 ratings), P1-N1 amplitude, P2-N1 amplitude, and EEG prediction accuracy, the clear condition  
284 was compared to each SNR level (resulting from the sliding average) using a paired samples t-  
285 test. False discovery rate (FDR) was used to account for multiple comparisons (Benjamini and  
286 Hochberg, 1995; Genovese et al., 2002). Age groups were compared at each SNR level  
287 individually using an independent-samples t-test and FDR thresholding. To investigate the  
288 overall effect of background babble and interaction with group, a repeated-measures analysis of  
289 variance (rmANOVA) was calculated with the within-participants factor Speech Clarity (clear,  
290 babble [averaged across SNR levels]) and Group (younger, older). In addition, analyses also  
291 explored neural responses for the older adult group split into those with clinically normal hearing  
292 (N = 15; PTA < 20 dB HL) and those with hearing impairment (N = 11; PTA > 20 dB HL).

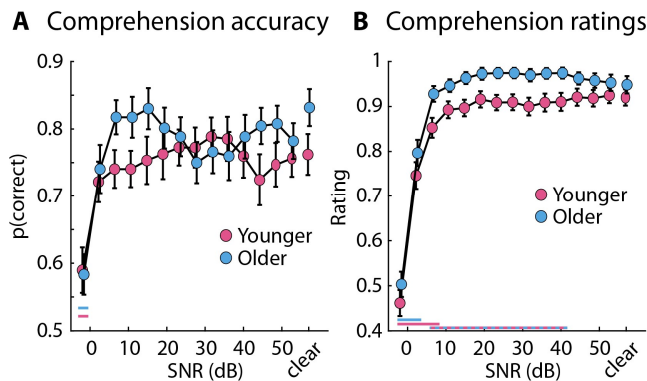
293 All statistical analyses were carried out using MATLAB (MathWorks) and JASP software  
294 (JASP, 2024; version 0.19.1). Note that for post hoc tests of an rmANOVA, JASP uses the  
295 rmANOVA degrees of freedom. The reported degrees of freedom may thus be higher than for  
296 direct contrasts had they been calculated independently from the rmANOVA.

297

## Results

### 298 **Older adults show reduced noise-related enhancements of neural speech tracking**

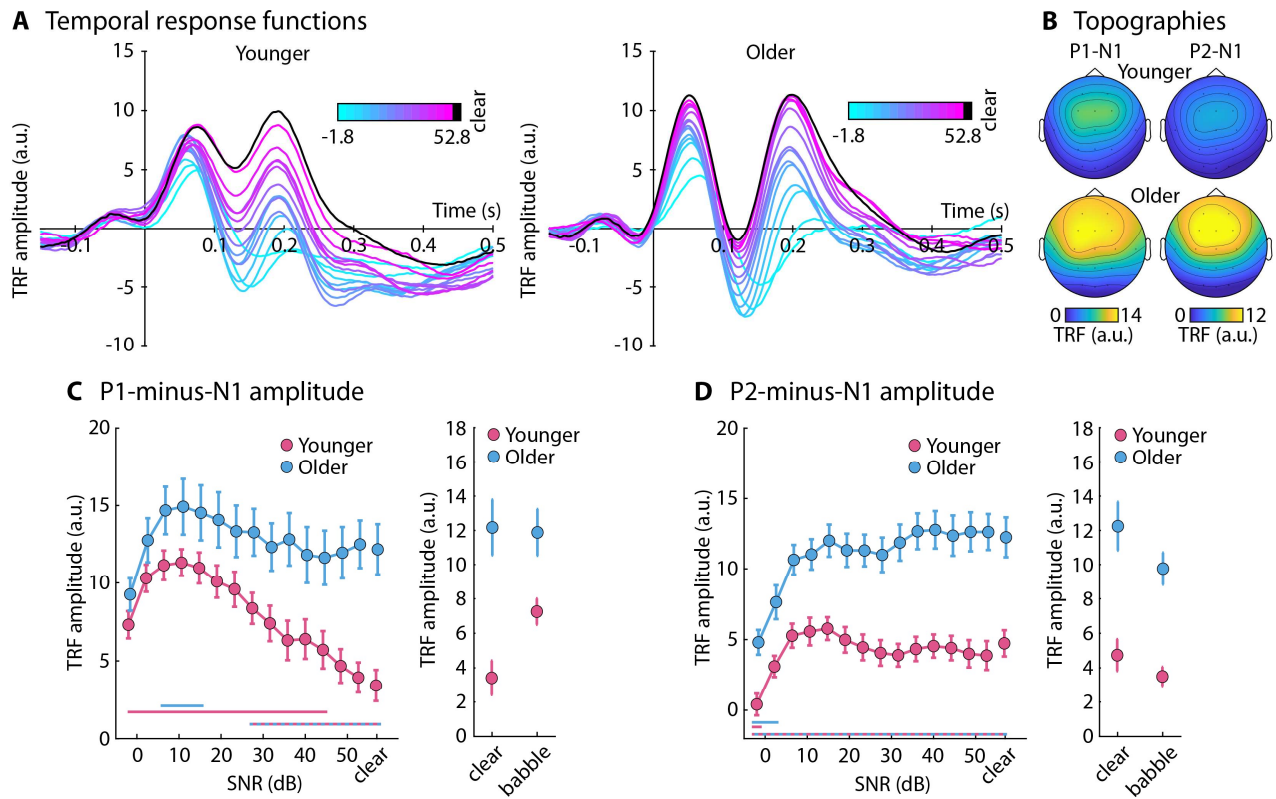
299 For both younger and older adults, story comprehension accuracy decreased for the most  
300 difficult SNRs relative to clear speech, but there were no differences between age groups (FDR-  
301 thresholded; Figure 3A). Ratings of speech comprehension/gist understanding decreased for  
302 SNRs below +10 dB compared to clear speech, for both age groups. Older adults also rated  
303 speech comprehension/gist understanding higher than younger adults for SNRs between +6.6  
304 and +40.2 dB, which may be related to the known higher subjective ratings of hearing abilities  
305 relative to objective hearing abilities in older compared to younger adults (Helfer et al., 2017;  
306 Helfer and Jesse, 2021).



**Figure 3: Behavioral results. A:** Story comprehension accuracy. **B:** Ratings of speech comprehension and gist understanding. The colored, horizontal lines close to the x-axis reflect a significant difference between clear speech and the different SNRs (FDR-thresholded). The two-colored (dashed) line reflects a significant difference between age groups (FDR-thresholded).

307

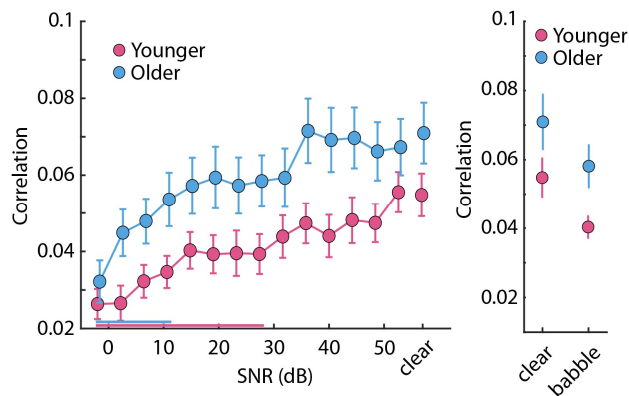
308 Figures 4A and 4B show the temporal responses functions and topographical  
309 distributions. Figures 4C displays P1-N1 amplitudes as they relate to speech-clarity conditions.  
310 For younger adults, P1-N1 amplitudes increased with decreasing SNR relative to clear speech,  
311 up to about +10 dB SNR, whereas amplitudes decreased for yet lower SNRs (Figure 4C, left). For  
312 older adults, the increase in P1-N1 amplitudes associated with background babble was only  
313 significant around +10 dB SNR, with amplitudes decreasing for lower SNRs (Figure 4C, left). In  
314 fact, P1-N1 amplitudes were greater for older compared to younger adults only for clear speech  
315 and for speech at high SNRs (i.e., >28 dB), because the auditory cortex of older adults showed a  
316 reduced sensitivity to background babble (Figure 3C, left). This reduced noise-sensitivity is also  
317 evidenced by the rmANOVA for the P1-N1 amplitude (clear speech vs speech in babble  
318 [collapsed across SNRs]). The Speech Clarity  $\times$  Group interaction ( $F_{1,50} = 15.714$ ,  $p = 2.3 \cdot 10^{-4}$ ,  $\omega^2$   
319 = 0.025) showed that the P1-N1 amplitude for younger adults was greater for speech in babble  
320 than for clear speech ( $t_{25} = 6.893$ ,  $p_{Holm} = 2.1 \cdot 10^{-5}$ ,  $d = 0.611$ ), whereas this was not the case for  
321 older adults ( $t_{25} = 0.392$ ,  $p_{Holm} = 0.697$ ,  $d = 0.046$ ; Figure 3C, right; effect of Speech Clarity:  $F_{1,50} =$   
322 11.628,  $p = 0.001$ ,  $\omega^2 = 0.018$ ; effect of Group:  $F_{1,50} = 16.207$ ,  $p = 1.9 \cdot 10^{-4}$ ,  $\omega^2 = 0.13$ ).



**Figure 4: Results for metrics derived from temporal response functions.** **A:** Temporal response functions for different speech-clarity conditions and for younger and older adults. **B:** Topographies for P1-N1 and P2-N1 TRF amplitudes. **C:** Left: P1-N1 amplitudes for different speech-clarity conditions and both age groups. The colored, horizontal lines close to the x-axis reflect a significant difference between clear speech and the SNR conditions (FDR-thresholded). The two-colored (dashed) line reflects a significant difference between age groups (FDR-thresholded). Right: P1-N1 TRF amplitude for clear speech and the mean across SNR conditions (babble). **D:** Same as in panel C for the P2-N1 TRF amplitudes.

323 Figures 4D displays the relation between speech-clarity conditions and P2-N1  
 324 amplitudes. For both younger and older adults, the P2-N1 amplitudes were smaller for SNRs  
 325 around 0 dB and below compared to clear speech, but older adults showed overall larger P2-N1  
 326 amplitudes for all speech-clarity conditions. This is also shown by the rmANOVA for the P2-N1  
 327 amplitude, revealing smaller amplitudes for speech in babble than clear speech ( $F_{1,50} = 13.780$ ,  $p$   
 328  $= 5.2 \cdot 10^{-4}$ ,  $\omega^2 = 0.030$ ) and larger amplitudes for older compared to younger adults ( $F_{1,50} =$   
 329  $26.698$ ,  $p = 4.2 \cdot 10^{-6}$ ,  $\omega^2 = 0.201$ ). The interaction was not significant ( $F_{1,50} = 1.540$ ,  $p = 0.220$ ,  $\omega^2$   
 330  $= 0.001$ ; Figure 3D).



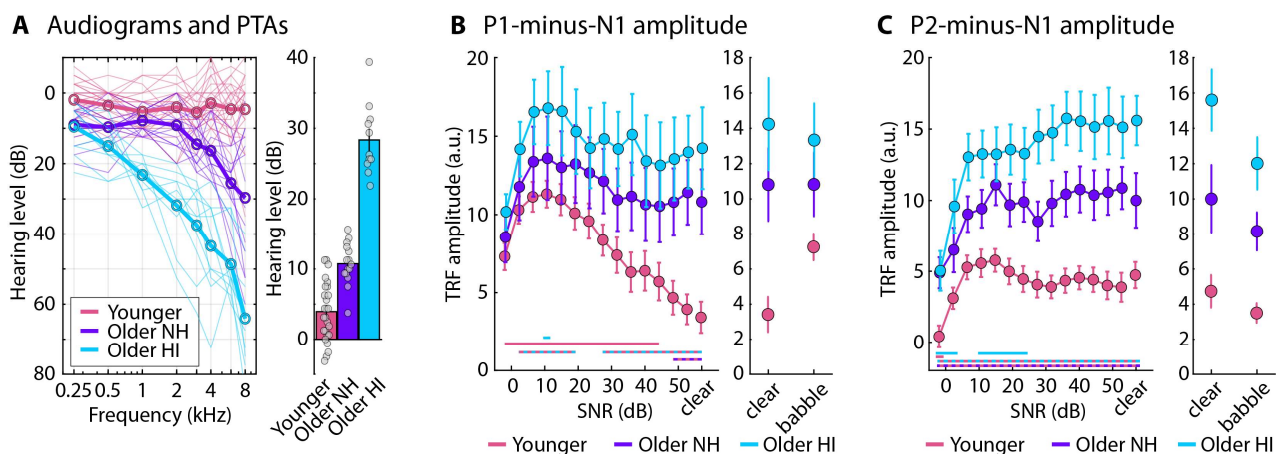


**Figure 5: EEG prediction accuracy.** **Left:** EEG prediction accuracy for each speech-clarity condition and age group. The colored, horizontal lines close to the x-axis reflect a significant difference between clear speech and the SNR conditions (FDR-thresholded). There was no significant difference between age groups for individual speech-clarity conditions (FDR-thresholded). **Right:** EEG prediction accuracy for clear speech and the mean across SNR conditions (babble).

331 Figure 5 shows the relation between speech-clarity conditions and EEG prediction  
 332 accuracy. Prediction accuracy decreased with decreasing SNR relative to clear speech (Figure 4,  
 333 left). This was also reflected in the rMANOVA, revealing smaller EEG prediction accuracies for  
 334 speech in babble than clear speech ( $F_{1,50} = 16.881$ ,  $p = 1.5 \cdot 10^{-4}$ ,  $\omega^2 = 0.046$ ) and younger  
 335 compared to older adults ( $F_{1,50} = 4.832$ ,  $p = 0.033$ ,  $\omega^2 = 0.036$ ). The interaction was not significant  
 336 ( $F_{1,50} = 0.064$ ,  $p = 0.801$ ,  $\omega^2 < 0.001$ ; Figure 4, right).

### 337 Comparing older adults with clinically ‘normal’ hearing to those with hearing impairment

338 Audiograms for younger adults, older adults with clinically ‘normal’ hearing, and older adults  
 339 with hearing impairment are shown in Figure 6A. Despite the group separation, the older adult  
 340 group with clinically ‘normal’ hearing still had greater pure-tone average thresholds compared to  
 341 younger adults ( $t_{39} = 5.536$ ,  $p = 2.3 \cdot 10^{-6}$ ,  $d = 1.795$ ), revealing subclinical hearing impairments  
 342 that are common among older individuals (Dubno et al., 2013; Plack, 2014; Helfer and Jesse,  
 343 2021).

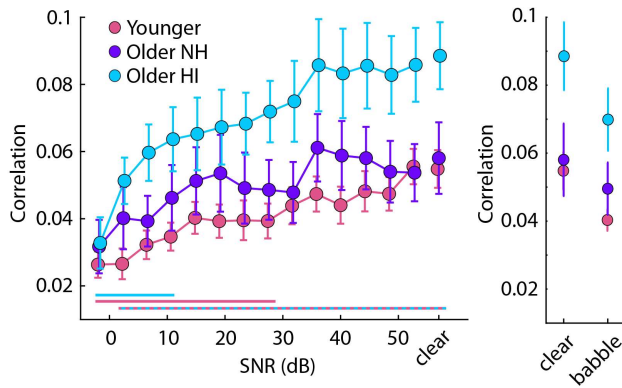


**Figure 6: Audiograms and neural-tracking amplitudes for groups split by presence vs absence of hearing impairment.** **A:** Audiograms (left) and pure-tone average thresholds for younger adults, older

adults with ‘normal’ hearing (NH), and older adults with clinical hearing impairment (HI). **B:** Left: P1-N1 TRF amplitudes for different speech-clarity conditions and groups. The colored, horizontal lines close to the x-axis reflect a significant difference between clear speech and the SNR conditions (FDR-thresholded). The two-colored (dashed) lines reflect a significant difference between groups (FDR-thresholded). Right: P1-N1 TRF amplitude for clear speech and the mean across SNR conditions (babble). **C:** Same as in panel B for the P2-N1 TRF amplitudes.

344 For neither of the two groups of older adults did the P1-N1 amplitudes show much  
345 sensitivity to background babble relative to clear speech, with the exception around +10 dB SNR  
346 for older adults with hearing impairment (Figure 6B, left). Critically, for clear speech and speech  
347 at high SNRs, the P1-N1 amplitude was larger in both older adult groups compared to younger  
348 adults, although the difference was significant for more SNRs for older adults with hearing  
349 impairment (FDR-thresholded; Figure 6B). This is also reflected in the rmANOVA for the P1-N1  
350 amplitude, showing a larger amplitude for both groups of older adults compared to younger  
351 adults (normal hearing:  $t_{49} = 2.844$ ;  $p_{\text{Holm}} = 0.013$ ;  $d = 0.877$ ; hearing impairment:  $t_{49} = 3.95$ ;  $p_{\text{Holm}} =$   
352  $7.5 \cdot 10^{-4}$ ;  $d = 1.352$ ), whereas no difference between the two older adult groups was found ( $t_{49} =$   
353  $1.256$ ;  $p_{\text{Holm}} = 0.215$ ;  $d = 0.474$ ; main effect of Group:  $F_{2,49} = 9.134$ ,  $p = 4.3 \cdot 10^{-4}$ ,  $\omega^2 = 0.098$ ).  
354 Group interacted with Speech Clarity ( $F_{2,49} = 7.963$ ,  $p = 0.001$ ,  $\omega^2 = 0.024$ ; Figure 5B, right):  
355 Younger adults showed a larger P1-N1 amplitude when speech was masked by babble  
356 compared to clear speech ( $t_{49} = 5.083$ ;  $p_{\text{Holm}} = 8.7 \cdot 10^{-5}$ ;  $d = 0.614$ ), whereas this was not  
357 significant in older adults with normal hearing ( $t_{49} = 0.009$ ;  $p_{\text{Holm}} = 1$ ;  $d = 0.001$ ) nor in older adults  
358 with hearing impairments ( $t_{49} = 0.776$ ;  $p_{\text{Holm}} = 1$ ;  $d = 0.144$ ).

359 A decrease in P2-N1 amplitudes with SNR, particularly for very low SNRs, relative to clear  
360 speech was observed for younger adults and older adults with hearing impairment (FDR-  
361 thresholded; Figure 6C left; this was significant for older adults without hearing impairment for  
362 uncorrected p-values). Moreover, P2-N1 amplitudes were larger for both older adult groups  
363 compared to younger adults for all SNRs (FDR-thresholded; Figure 6C left). The rmANOVA  
364 further corroborated this, revealing larger P2-N1 amplitudes for older adults with hearing  
365 impairment relative to those with normal hearing ( $t_{49} = 2.62$ ;  $p_{\text{Holm}} = 0.012$ ;  $d = 0.967$ ) and younger  
366 adults ( $t_{49} = 5.921$ ;  $p_{\text{Holm}} = 9.3 \cdot 10^{-7}$ ;  $d = 1.98$ ), and larger amplitudes for older adults with normal  
367 hearing relative to younger adults ( $t_{49} = 3.36$ ;  $p_{\text{Holm}} = 0.003$ ;  $d = 1.013$ ; main effect of Group:  $F_{2,49} =$   
368  $18.640$ ,  $p = 9.5 \cdot 10^{-7}$ ,  $\omega^2 = 0.190$ ). P2-N1 amplitudes were lower for speech in babble than clear  
369 speech ( $F_{1,49} = 17.527$ ,  $p = 1.2 \cdot 10^{-4}$ ,  $\omega^2 = 0.043$ ), but there was no interaction ( $F_{2,49} = 1.643$ ,  $p =$   
370  $0.204$ ,  $\omega^2 = 0.003$ ; Figure 5C, right).



**Figure 7: EEG prediction accuracy for groups split by the presence vs absence of hearing impairment. Left:** EEG prediction accuracy for each speech-clarity condition and age group. The colored, horizontal lines close to the x-axis reflect a significant difference between clear speech and the SNR conditions (FDR-thresholded). The two-colored (dashed) line reflects a significant difference between groups (FDR-thresholded). **Right:** EEG prediction accuracy for clear speech and the mean across SNR conditions (babble). NH – normal hearing; HI – hearing impairment

371 For EEG prediction accuracy, an SNR-related decrease relative to clear speech was  
372 observed for younger adults and older adults with hearing impairment (FDR-thresholded; Figure  
373 7 left). Prediction accuracy was greater for older adults with hearing impairment relative to  
374 younger adults for most speech-clarity conditions (FDR-thresholded; Figure 7 left). The  
375 rmANOVA for EEG prediction accuracy showed smaller accuracies for speech in babble than  
376 clear speech ( $F_{1,49} = 15.474$ ,  $p = 2.6 \cdot 10^{-4}$ ,  $\omega^2 = 0.046$ ). Prediction accuracy was greater for older  
377 adults with hearing impairment than younger adults ( $t_{49} = 3.307$ ;  $p_{\text{Holm}} = 0.005$ ;  $d = 1.085$ ) and  
378 older adults with ‘normal’ hearing ( $t_{49} = 2.408$ ;  $p_{\text{Holm}} = 0.040$ ;  $d = 0.872$ ), whereas there was no  
379 difference between the two latter groups ( $t_{49} = 0.721$ ;  $p_{\text{Holm}} = 0.474$ ;  $d = 0.213$ ; effect of Group:  
380  $F_{2,49} = 5.546$ ,  $p = 0.007$ ,  $\omega^2 = 0.057$ ; Figure 7 right).

## 381 Discussion

382 The current study investigated the extent to which auditory cortex of older adults shows noise-  
383 related enhancements in neural speech tracking. Younger and older adults listened to spoken  
384 stories either in quiet (clear) or in the presence of background noise. In younger adults, neural  
385 speech tracking, as evidenced by the P1-N1 amplitude of the temporal response functions, was  
386 enhanced when speech was presented in minimal background noise. Although neural tracking  
387 of speech in quiet and speech masked by minimal background noise was enhanced for older  
388 adults compared to younger adults, older adults showed little evidence of noise-related  
389 enhancements. The data demonstrate different sensitivity of the auditory cortex between  
390 younger and older people to speech masked by acoustic background noise. The data are  
391 consistent with the idea that internal, neural noise enhances neural speech tracking in older  
392 adults – possibly through stochastic resonance – that reduces the impact of additional external,  
393 acoustic noise.

394 **Noise-related enhancement of neural speech tracking**

395 The current study shows that, for younger adults, minimal background noise increases the  
396 neural tracking of the amplitude-onset envelope of speech compared to speech presented in  
397 quiet (P1-N1 amplitude; Figure 4). This is remarkable, given that the background babble overlaps  
398 spectrally with the speech, but a noise-related enhancement has been shown recently in a few  
399 other works for speech and more simple sounds (Alain et al., 2009; Ward et al., 2010; Parbery-  
400 Clark et al., 2011; Alain et al., 2012; Alain et al., 2014; Shukla and Bidelman, 2021; Herrmann,  
401 2024; Panella et al., 2024).

402 Minor background noise appears to be sufficient to enhance neural tracking, because the  
403 enhancement is present for very high SNRs (>40 dB; Figure 4) for which speech is as intelligible  
404 as speech in quiet (Holder et al., 2018; Rowland et al., 2018; Spyridakou et al., 2020; Irsik et al.,  
405 2022; Figures 3 and 4). Neural speech tracking decreases only for low SNRs for which speech  
406 comprehension declines (Figures 3 and 4). Critically, a recent study shows that the noise-related  
407 enhancement in speech tracking is present even when participants attend to a demanding visual  
408 task rather than to the speech (Herrmann, 2024), making it unlikely that attentional effort  
409 explains the enhancement, especially at high SNRs (Rowland et al., 2018). Moreover, the  
410 enhancement was present only for early sensory responses (P1-N1), but not for later responses  
411 (P2-N1) and EEG prediction accuracy (which integrates responses over time), consistent with a  
412 sensory-driven nature of the enhancement.

413 A few neural speech tracking studies used noise that matched the spectrum of speech as  
414 a background masker but did not find a noise-related enhancement (Ding and Simon, 2013; Zou  
415 et al., 2019; Synigal et al., 2023). However, these studies used low SNRs for which speech is less  
416 intelligible (<10 dB) and neural tracking reduced (Figure 4). Moreover, babble noise appears to  
417 enhance the neural tracking more than speech-matched noise (Herrmann, 2024), potentially  
418 because the babble facilitates neural activity in the same speech-relevant auditory regions that  
419 are recruited by the speech. This points to some specificity of the spectral noise properties in  
420 facilitating the amplification (cf. Krauss and Tziridis, 2021)

421 Stochastic resonance – that is the response facilitation of a non-linear system through  
422 noise (Stocks, 2000; Ward et al., 2002; Moss et al., 2004; Stein et al., 2005; McDonnell and  
423 Abbott, 2009; McDonnell and Ward, 2011; Krauss et al., 2016; Schilling et al., 2023; Figure 1) –  
424 has been suggested to underlie the enhancement of neural speech tracking in the presence of  
425 background noise (Herrmann, 2024). Observing neural speech tracking with scalp EEG requires  
426 the synchronized activity of more than 10,000 neurons (Niedermeyer and da Silva, 2005; da  
427 Silva, 2010). Some neurons may not have been driven enough to elicit a response for speech in  
428 quiet, whereas the presence of noise – through stochastic resonance – may have pushed them  
429 beyond their firing threshold (Figure 1). Stochastic resonance may help individuals to hear  
430 robustly even when background noise is present.

431 **Age-related enhancement in neural speech tracking**

432 Neural speech tracking was enhanced in older compared to younger adults for all metrics (P1-  
433 N1, P2-N1, EEG prediction accuracy), especially for speech in quiet (clear) and speech masked  
434 by minimal background noise. An age-related enhancement in neural speech tracking is  
435 consistent with previous work (Presacco et al., 2016a, b; Brodbeck et al., 2018; Broderick et al.,  
436 2021; Panela et al., 2024) and with work showing larger neural responses to tones and noises for  
437 older adults (Sörös et al., 2009; Alain et al., 2014; Bidelman et al., 2014; Stothart and Kazanina,  
438 2016; Irsik et al., 2021; Herrmann et al., 2023). Such hyperactivity is thought to result from a loss  
439 of inhibition and an increase in excitation in the auditory pathway due to reduced peripheral  
440 inputs associated with aging and hearing loss (Caspary et al., 2008; Caspary et al., 2013; Ouellet  
441 and de Villers-Sidani, 2014; Zhao et al., 2016; Resnik and Polley, 2017; Salvi et al., 2017;  
442 Herrmann and Butler, 2021; McClaskey, 2024).

443 The age-related enhancement of the P1-N1 amplitude appears mostly to be due to aging  
444 (Figure 6B), whereas the P2-N1 amplitude and the EEG prediction accuracy seem to be also or  
445 exclusively driven by hearing loss (Figures 6C and 7). Previous studies have shown mixed results  
446 regarding the effects of aging versus hearing loss on neural speech tracking. Some studies have  
447 found that neural speech tracking is greater for older adults with hearing loss compared to those  
448 without (Decruy et al., 2020; Fuglsang et al., 2020; Gillis et al., 2022; Schmitt et al., 2022),  
449 whereas other studies point to age-related enhancements per se (Presacco et al., 2019).  
450 Counterintuitively, the current data suggest that earlier, sensory responses (P1-N1) are less  
451 affected by hearing loss than later responses (P2-N1). However, distinguishing between the  
452 impacts of hearing loss versus aging per se may not be possible (Humes et al., 2012). Even minor  
453 hearing loss or peripheral damage that is less-well detectable with pure-tone audiometry, such  
454 as damage to synapses connecting to auditory nerve fibers (Kujawa and Liberman, 2009;  
455 Bharadwaj et al., 2014; Liberman and Kujawa, 2017), can lead to hyperactivity in the auditory  
456 system (Qiu et al., 2000; Munguia et al., 2013; Resnik and Polley, 2017; Salvi et al., 2017). The  
457 current enhancements for older adults with clinically ‘normal’ hearing compared to younger  
458 adults may thus still be related to differences in hearing abilities (i.e., audiometric thresholds  
459 were elevated even in normal-hearing older adults; Figure 6A). Speculatively, minor hearing loss  
460 is sufficient to enhance early sensory responses, which then do not amplify further with  
461 worsening hearing abilities.

462 **Age-related differences in sensitivity of neural speech tracking to background noise**

463 The main purpose of the current study was to investigate whether auditory cortex of older adults  
464 shows enhancements in speech tracking due to acoustic background noise. However, there was  
465 little evidence that neural speech tracking in older adults with or without hearing loss is



466 enhanced by acoustic background noise (Figures 4C and 6B). There was only a minor increase in  
467 the P1-N1 amplitude around +10 dB SNR, but speech comprehension for this and lower SNRs is  
468 more difficult for listeners (Irsik et al., 2022; Herrmann, 2023) and the increase could thus be  
469 due to attentional effort (Pichora-Fuller et al., 2016; Hauswald et al., 2022; Yasmin et al., 2023).  
470 Neural speech tracking decreased for both younger and older adults for lower SNRs, for which  
471 speech comprehension decreased as well; this is consistent with previous work (Ding and  
472 Simon, 2013; Zou et al., 2019; Yasmin et al., 2023).

473 A loss of neural inhibition and an increase in neural excitation due to aging and hearing  
474 loss can lead to a gain or amplification of a neuron's output, because under such circumstances  
475 a smaller input to the neuron is already sufficient for it to fire compared to a less excitable and  
476 inhibited neuron. This gain – or central gain, because the amplification increases along the  
477 auditory pathway – is often referred to as the cause for hyperresponsivity to sound and speech  
478 (Auerbach et al., 2014; Zhao et al., 2016; Salvi et al., 2017; Herrmann and Butler, 2021;  
479 McClaskey, 2024). The reduced sensitivity to acoustic noise for older adults observed in the  
480 current study could potentially be the result of a maxed-out central gain, such that any noisy  
481 neural activity driven by the acoustic noise that would normally facilitate enhancements due to  
482 stochastic resonance may not be as effective under conditions of maxed-out central gain.  
483 However, this may not be a sufficient explanation, since there was a minor effect of background  
484 noise around +10 dB SNR (Figures 4C and 6B) and hearing loss appeared to increase the P1-N1  
485 amplitude slightly (but non-significantly; Figure 6B), suggesting that central gain may not be  
486 entirely at its maximum.

487 Alternatively or in addition, spontaneous activity – that is, neural noise – in the auditory  
488 system is known to increase with age and hearing loss (Kaltenbach and Afman, 2000; Eggermont  
489 and Roberts, 2004; Munguia et al., 2013; Eggermont, 2015; Parthasarathy et al., 2019). Internal,  
490 neural noise in older adults may have a comparable effect on the neural tracking of speech in  
491 quiet as the external, acoustic noise has in younger adults. Indeed, increased neural noise in  
492 hearing loss has been suggested to lead to stochastic resonance phenomena where the neural  
493 noise can increase sensitivity to sound (Krauss et al., 2016; Krauss and Tziridis, 2021; Schilling et  
494 al., 2023). Increased internal, neural noise in older adults could reduce the sensitivity to  
495 external, acoustic noise, because the noisy neural activity elicited by acoustic noise may not  
496 necessarily add further output amplification to the aged auditory system if it is already amplified  
497 through stochastic resonance (resulting from neural noise). Although, central gain and  
498 increased neural noise are not independent phenomena and distinguishing one from the other  
499 with non-invasive recording techniques may be challenging (Schilling et al., 2023), the current  
500 results are the first to point at the possibility that internal neural noise in the auditory cortex of  
501 older people could underlie enhanced neural speech tracking.



502

### Acknowledgments

503 I thank Priya Pandey and Saba Junaid for their help with story generation and data collection. The  
504 research was supported by the Canada Research Chair program (CRC-2019-00156) and the  
505 Natural Sciences and Engineering Research Council of Canada (Discovery Grant: RGPIN-2021-  
506 02602).

507

### Author contributions

508 **Björn Herrmann:** Conceptualization, methodology, formal analysis, investigation, data  
509 curation, writing - original draft, writing - review and editing, visualization, supervision, project  
510 administration, funding acquisition.

511

### Statements and Declarations

512 The author has no conflicts or competing interests.

513

### Data availability

514 Data are available at <https://osf.io/> (upon publication). One younger and one older participant  
515 declined to share their data publicly (we employ separate consents for study participation and  
516 data sharing in line with Canadian Tri-Council Policies for Ethical Conduct for Research Involving  
517 Humans – TCPS 2 from 2022). Their data are thus not made available.

518

### References

519 Alain C, McDonald K, Van Roon P (2012) Effects of age and background noise on processing a  
520 mistuned harmonic in an otherwise periodic complex sound. *Hearing Research* 283:126-  
521 135.  
522 Alain C, Roye A, Salloum C (2014) Effects of age-related hearing loss and background noise on  
523 neuromagnetic activity from auditory cortex. *Frontiers in Systems Neuroscience* 8:Article 8.  
524 Alain C, Quan J, McDonald K, Van Roon P (2009) Noise-induced increase in human auditory  
525 evoked neuromagnetic fields. *European Journal of Neuroscience* 30:132-142.  
526 Auerbach BD, Rodrigues PV, Salvi RJ (2014) Central gain control in tinnitus and hyperacusis.  
527 *Frontiers in Neurology* 5:Article 206.  
528 Bell AJ, Sejnowski TJ (1995) An information maximization approach to blind separation and blind  
529 deconvolution. *Neural Computation* 7:1129-1159.

- 530 Benjamini Y, Hochberg Y (1995) Controlling the false discovery rate: a practical and powerful  
531 approach to multiple testing. *Journal of the Royal Statistical Society Series B* 57:289-300.
- 532 Bharadwaj HM, Verhulst S, Shaheen L, Liberman MC, Shinn-Cunningham BG (2014) Cochlear  
533 neuropathy and the coding of supra-threshold sound. *Frontiers in Systems Neuroscience*  
534 8:Article 26.
- 535 Bidelman GM, Villafuerte JW, Moreno S, Alain C (2014) Age-related changes in the  
536 subcortical encoding and categorical perception of speech. *Neurobiology of Aging*  
537 35:2526-2540.
- 538 Biesmans W, Das N, Francart T, Bertrand A (2017) Auditory-Inspired Speech Envelope Extraction  
539 Methods for Improved EEG-Based Auditory Attention Detection in a Cocktail Party Scenario.  
540 *IEEE Transactions on Neural Systems and Rehabilitation Engineering* 25:402-412.
- 541 Bilger RC (1984) *Manual for the clinical use of the revised SPIN Test*. Champaign, IL, USA: The  
542 University of Illinois.
- 543 Bilger RC, Nuetzel JM, Rabinowitz WM, Rzeczkowski C (1984) Standardization of a Test of  
544 Speech Perception in Noise. *Journal of Speech, Language, and Hearing Research* 27:32-48.
- 545 Brodbeck C, Simon JZ (2020) Continuous speech processing. *Current Opinion in Physiology*  
546 18:25-31.
- 547 Brodbeck C, Presacco A, Anderson S, Simon JZ (2018) Over-representation of speech in older  
548 adults originates from early response in higher order auditory cortex. *Acta Acust United*  
549 *Acust* 104:774-777.
- 550 Broderick MP, Di Liberto GM, Anderson AJ, Rofes A, Lalor EC (2021) Dissociable  
551 electrophysiological measures of natural language processing reveal differences in speech  
552 comprehension strategy in healthy ageing. *Scientific Reports* 11:4963.
- 553 Caspary DM, Hughes LF, Ling LL (2013) Age-related GABAA receptor changes in rat auditory  
554 cortex. *Neurobiology of Aging* 34:1486-1496.
- 555 Caspary DM, Ling L, Turner JG, Hughes LF (2008) Inhibitory neurotransmission, plasticity and  
556 aging in the mammalian central auditory system. *The Journal of Experimental Biology*  
557 211:1781-1791.
- 558 Chambers AR, Resnik J, Yuan Y, Whitton JP, Edge AS, Liberman MC, Polley DB (2016) Central  
559 Gain Restores Auditory Processing following Near-Complete Cochlear Denervation. *Neuron*  
560 89:867-879.
- 561 Cohen SS, Parra LC (2016) Memorable Audiovisual Narratives Synchronize Sensory and  
562 Supramodal Neural Responses. *eNeuro* 3:e0203.
- 563 Crosse MJ, Di Liberto GM, Bednar A, Lalor EC (2016) The Multivariate Temporal Response  
564 Function (mTRF) Toolbox: A MATLAB Toolbox for Relating Neural Signals to Continuous  
565 Stimuli. *Frontiers in human neuroscience* 10:604.

- 566 Crosse MJ, Zuk NJ, Di Liberto GM, Nidiffer AR, Molholm S, Lalor EC (2021) Linear Modeling of  
567 Neurophysiological Responses to Speech and Other Continuous Stimuli: Methodological  
568 Considerations for Applied Research. *Frontiers in Neuroscience* 15.
- 569 da Silva FL (2010) EEG: Origin and Measurement. In: *EEG - fMRI: Physiological Basis, Technique,*  
570 *and Applications* (Mulert C, Lemieux L, eds), pp 19-38. Berlin, Heidelberg: Springer Berlin  
571 Heidelberg.
- 572 Daube C, Ince RAA, Gross J (2019) Simple Acoustic Features Can Explain Phoneme-Based  
573 Predictions of Cortical Responses to Speech. *Current Biology* 29:1924-1937.
- 574 Davis MH, Johnsrude IS (2003) Hierarchical Processing in Spoken Language Comprehension.  
575 *The Journal of Neuroscience* 23:3423-3431.
- 576 Decruy L, Vanthornhout J, Francart T (2019) Evidence for enhanced neural tracking of the speech  
577 envelope underlying age-related speech-in-noise difficulties. *Journal of Neurophysiology*  
578 122:601-615.
- 579 Decruy L, Vanthornhout J, Francart T (2020) Hearing impairment is associated with enhanced  
580 neural tracking of the speech envelope. *Hearing Research* 393:107961.
- 581 Di Liberto Giovanni M, O'Sullivan James A, Lalor Edmund C (2015) Low-Frequency Cortical  
582 Entrainment to Speech Reflects Phoneme-Level Processing. *Current Biology* 25:2457-2465.
- 583 Ding N, Simon JZ (2013) Adaptive temporal encoding leads to a background-insensitive cortical  
584 representation of speech. *The Journal of Neuroscience* 33:5728-5735.
- 585 Ding N, Simon JZ (2014) Cortical entrainment to continuous speech: functional roles and  
586 interpretations. *Frontiers in Human Neuroscience* 8.
- 587 Ding N, Chatterjee M, Simon JZ (2014) Robust cortical entrainment to the speech envelope relies  
588 on the spectro-temporal fine structure. *NeuroImage* 88:41-46.
- 589 Dmochowski JP, Sajda P, Dias J, Parra LC (2012) Correlated components of ongoing EEG point to  
590 emotionally laden attention – a possible marker of engagement? *Frontiers in Human*  
591 *Neuroscience* 6:Article 112.
- 592 Dmochowski JP, Bezdek MA, Abelson BP, Johnson JS, Schumacher EH, Parra LC (2014)  
593 Audience preferences are predicted by temporal reliability of neural processing. *Nature*  
594 *Communications* 29:4567.
- 595 Dubno JR, Eckert MA, Lee F-S, Matthews LJ, Schmiedt RA (2013) Classifying Human Audiometric  
596 Phenotypes of Age-Related Hearing Loss from Animal Models. *Journal of the Association for*  
597 *Research in Otolaryngology* 14:687-701.
- 598 Eggermont JJ (2015) Animal models of spontaneous activity in the healthy and impaired auditory  
599 system. *Frontiers in Neural Circuits* 9:19.
- 600 Eggermont JJ, Roberts LE (2004) The neuroscience of tinnitus. *Trends in Neurosciences* 27:676-  
601 682.

- 602 Feder K, Michaud D, Ramage-Morin P, McNamee J, Beauregard Y (2015) Prevalence of hearing  
603 loss among Canadians aged 20 to 79: Audiometric results from the 2012/2013 Canadian  
604 Health Measures Survey. *Health Reports* 26:18-25.
- 605 Fiedler L, Wöstmann M, Herbst SK, Obleser J (2019) Late cortical tracking of ignored speech  
606 facilitates neural selectivity in acoustically challenging conditions. *Neuroimage* 186:33-42.
- 607 Fiedler L, Wöstmann M, Graversen C, Brandmeyer A, Lunner T, Obleser J (2017) Single-channel  
608 in-ear-EEG detects the focus of auditory attention to concurrent tone streams and mixed  
609 speech. *Journal of Neural Engineering* 14:036020.
- 610 Fuglsang SA, Märcher-Rørsted J, Dau T, Hjortkjær J (2020) Effects of sensorineural hearing loss  
611 on cortical synchronization to competing speech during selective attention. *The Journal of*  
612 *Neuroscience*:1936-1919.
- 613 Genovese CR, Lazar NA, Nichols T (2002) Thresholding of statistical maps in functional  
614 neuroimaging using the false discovery rate. *NeuroImage* 15:870–878.
- 615 Gillis M, Decruy L, Vanthornhout J, Francart T (2022) Hearing loss is associated with delayed  
616 neural responses to continuous speech. *European Journal of Neuroscience* 55:1671-1690.
- 617 Goman AM, Lin FR (2016) Prevalence of Hearing Loss by Severity in the United States. *American*  
618 *Journal of Public Health* 106:1820-1822.
- 619 Hauswald A, Keitel A, Chen Y-P, Rösch S, Weisz N (2022) Degradation levels of continuous  
620 speech affect neural speech tracking and alpha power differently. *European Journal of*  
621 *Neuroscience* 55:3288-3302.
- 622 Helfer KS, Jesse A (2021) Hearing and speech processing in midlife. *Hearing Research*  
623 402:108097.
- 624 Helfer KS, Merchant GR, Wasiuk PA (2017) Age-Related Changes in Objective and Subjective  
625 Speech Perception in Complex Listening Environments. *Journal of Speech, Language, and*  
626 *Hearing Research* 60:3009-3018.
- 627 Herrmann B (2023) The perception of artificial-intelligence (AI) based synthesized speech in  
628 younger and older adults. *International Journal of Speech Technology* 26:395-415.
- 629 Herrmann B (2024) Minimal background noise enhances neural speech tracking: Evidence of  
630 stochastic resonance. *BioRxiv*.
- 631 Herrmann B, Johnsrude IS (2018) Attentional State Modulates the Effect of an Irrelevant Stimulus  
632 Dimension on Perception. *Journal of Experimental Psychology: Human Perception and*  
633 *Performance* 44:89-105.
- 634 Herrmann B, Johnsrude IS (2020) A Model of Listening Engagement (MoLE). *Hearing Research*  
635 397:108016.
- 636 Herrmann B, Butler BE (2021) Hearing Loss and Brain Plasticity: The Hyperactivity Phenomenon.  
637 *Brain Structure & Function* 226:2019-2039.

- 638 Herrmann B, Maess B, Johnsrude IS (2018) Aging Affects Adaptation to Sound-Level Statistics in  
639 Human Auditory Cortex. *The Journal of Neuroscience* 38:1989-1999.
- 640 Herrmann B, Maess B, Johnsrude IS (2022) A Neural Signature of Regularity in Sound is Reduced  
641 in Older Adults. *Neurobiology of Aging* 109:1-10.
- 642 Herrmann B, Maess B, Johnsrude IS (2023) Sustained responses and neural synchronization to  
643 amplitude and frequency modulation in sound change with age. *Hearing Research*  
644 428:108677.
- 645 Hertrich I, Dietrich S, Trouvain J, Moos A, Ackermann H (2012) Magnetic brain activity  
646 phase-locked to the envelope, the syllable onsets, and the fundamental frequency of a  
647 perceived speech signal. *Psychophysiology* 49:322-334.
- 648 Holder JT, Levin LM, Gifford RH (2018) Speech Recognition in Noise for Adults With Normal  
649 Hearing: Age-Normative Performance for AzBio, BKB-SIN, and QuickSIN. *Otology &*  
650 *Neurotology* 39:e972-e978.
- 651 Humes LE (2019) Examining the Validity of the World Health Organization's Long-Standing  
652 Hearing Impairment Grading System for Unaided Communication in Age-Related Hearing  
653 Loss. *American Journal of Audiology* 28:810-818.
- 654 Humes LE, Dubno JR, Gordon-Salant S, Lister JJ, Cacace AT, Cruickshanks KJ, Gates GA, Wilson  
655 RH, Wingfield A (2012) Central Presbycusis: A Review and Evaluation of the Evidence.  
656 *Journal of the American Academy of Audiology* 23:635-666.
- 657 Irsik VC, Johnsrude IS, Herrmann B (2022) Neural activity during story listening is synchronized  
658 across individuals despite acoustic masking. *Journal of Cognitive Neuroscience* 34:933-  
659 950.
- 660 Irsik VC, Almanaseer A, Johnsrude IS, Herrmann B (2021) Cortical Responses to the Amplitude  
661 Envelopes of Sounds Change with Age. *The Journal of Neuroscience* 41:5045-5055.
- 662 Izhikevich EM (2003) Simple model of spiking neurons. *IEEE Transactions on Neural Networks*  
663 14:1569-1572.
- 664 Izhikevich EM (2004) Which model to use for cortical spiking neurons? *IEEE Transactions on*  
665 *Neural Networks* 15:1063-1070.
- 666 JASP (2024) JASP [Computer software]. In: <https://jasp-stats.org/>.
- 667 Kaltenbach JA, Afman CE (2000) Hyperactivity in the dorsal cochlear nucleus after intense sound  
668 exposure and its resemblance to tone-evoked activity: a physiological model for tinnitus.  
669 *Hearing Research* 140:165-172.
- 670 Karunathilake IMD, Dunlap JL, Perera J, Presacco A, Decruy L, Anderson S, Kuchinsky SE, Simon  
671 JZ (2023) Effects of aging on cortical representations of continuous speech. *Journal of*  
672 *Neurophysiology* 129:1359-1377.
- 673 Krauss P, Tziridis K (2021) Simulated transient hearing loss improves auditory sensitivity.  
674 *Scientific Reports* 11:14791.

- 675 Krauss P, Tziridis K, Metzner C, Schilling A, Hoppe U, Schulze H (2016) Stochastic Resonance  
676 Controlled Upregulation of Internal Noise after Hearing Loss as a Putative Cause of Tinnitus-  
677 Related Neuronal Hyperactivity. *Frontiers in Neuroscience* 10:Article 597.
- 678 Kujawa SG, Liberman MC (2009) Adding Insult to Injury: Cochlear Nerve Degeneration after  
679 “Temporary” Noise-Induced Hearing Loss. *The Journal of Neuroscience* 29:14077-14085.
- 680 Lalor EC, Foxe JJ (2010) Neural responses to uninterrupted natural speech can be extracted with  
681 precise temporal resolution. *European Journal of Neuroscience* 31:189-193.
- 682 Leek MR (2011) Adaptive procedures in psychophysical research. *Perception & Psychophysics*  
683 63:1279-1292.
- 684 Lesenfants D, Vanthornhout J, Verschueren E, Decruy L, Francart T (2019) Predicting individual  
685 speech intelligibility from the cortical tracking of acoustic- and phonetic-level speech  
686 representations. *Hearing Research* 380:1-9.
- 687 Liberman MC, Kujawa SG (2017) Cochlear synaptopathy in acquired sensorineural hearing loss:  
688 Manifestations and mechanisms. *Hearing Research* 349:138-147.
- 689 Luo H, Poeppel D (2007) Phase Patterns of Neuronal Responses Reliably Discriminate Speech in  
690 Human Auditory Cortex. *Neuron* 54:1001-1010.
- 691 Makeig S, Bell AJ, Jung T-P, Sejnowski TJ (1995) Independent component analysis of  
692 electroencephalographic data. In: *Advances in Neural Information Processing Systems*  
693 (Touretzky D, Mozer M, Hasselmo M, eds), pp 145-151. Cambridge, MA, USA: MIT Press.
- 694 Mathiesen SL, Van Hedger SC, Irsik VC, Bain MM, Johnsrude IS, Herrmann B (2024) Exploring age  
695 differences in absorption and enjoyment during story listening. *Psychology International*  
696 6:667-684.
- 697 Mattys SL, Davis MH, Bradlow AR, Scott SK (2012) Speech recognition in adverse conditions: A  
698 review. *Language and Cognitive Processes* 27:953-978.
- 699 McClaskey CM (2024) Neural hyperactivity and altered envelope encoding in the central auditory  
700 system: Changes with advanced age and hearing loss. *Hearing Research* 442:108945.
- 701 McDermott Josh H, Simoncelli Eero P (2011) Sound Texture Perception via Statistics of the  
702 Auditory Periphery: Evidence from Sound Synthesis. *Neuron* 71:926-940.
- 703 McDonnell MD, Abbott D (2009) What Is Stochastic Resonance? Definitions, Misconceptions,  
704 Debates, and Its Relevance to Biology. *PLOS Computational Biology* 5:e1000348.
- 705 McDonnell MD, Ward LM (2011) The benefits of noise in neural systems: bridging theory and  
706 experiment. *Nature Reviews Neuroscience* 12:415-425.
- 707 Moore BCJ (2007) *Cochlear Hearing Loss: Physiological, Psychological and Technical Issues*.  
708 West Sussex, England: John Wiley & Sons, Ltd.
- 709 Moss F, Ward LM, Sannita WG (2004) Stochastic resonance and sensory information processing:  
710 a tutorial and review of application. *Clinical Neurophysiology* 115:267-281.



- 711 Munguia R, Pienkowski M, Eggermont JJ (2013) Spontaneous firing rate changes in cat primary  
712 auditory cortex following long-term exposure to non-traumatic noise: Tinnitus without  
713 hearing loss? *Neuroscience Letters* 546:46-50.
- 714 Näätänen R, Picton TW (1987) The N1 wave of the human electric and magnetic response to  
715 sound: a review and an analysis of the component structure. *Psychophysiology* 24:375-425.
- 716 Niedermeyer E, da Silva FHL (2005) *Electroencephalography: Basic Principles, Clinical  
717 Applications, and Related Fields*: Lippincott Williams & Wilkins.
- 718 Oostenveld R, Fries P, Maris E, Schoffelen JM (2011) FieldTrip: Open source software for  
719 advanced analysis of MEG, EEG, and invasive electrophysiological data. *Computational  
720 Intelligence and Neuroscience* 2011:Article ID 156869.
- 721 Ouellet L, de Villers-Sidani E (2014) Trajectory of the main GABAergic interneuron populations  
722 from early development to old age in the rat primary auditory cortex. *Frontiers in  
723 Neuroanatomy* 8:Article 40.
- 724 Panela RA, Copelli F, Herrmann B (2024) Reliability and generalizability of neural speech tracking  
725 in younger and older adults. *Neurobiology of Aging* 134:165-180.
- 726 Parbery-Clark A, Marmel F, Bair J, Kraus N (2011) What subcortical–cortical relationships tell us  
727 about processing speech in noise. *European Journal of Neuroscience* 33:549-557.
- 728 Parthasarathy A, Herrmann B, Bartlett EL (2019) Aging alters envelope representations of  
729 speech-like sounds in the inferior colliculus. *Neurobiology of Aging* 73:30-40.
- 730 Pichora-Fuller MK, Kramer SE, Eckert MA, Edwards B, Hornsby BWY, Humes LE, Lemke U,  
731 Lunner T, Matthen M, Mackersie CL, Naylor G, Phillips NA, Richter M, Rudner M, Sommers  
732 MS, Tremblay KL, Wingfield A (2016) *Hearing Impairment and Cognitive Energy: The  
733 Framework for Understanding Effortful Listening (FUEL)*. *Ear & Hearing* 37 Suppl 1:5S-27S.
- 734 Picton TW, John SM, Dimitrijevic A, Purcell DW (2003) Human auditory steady-state responses.  
735 *International Journal of Audiology* 42:177-219.
- 736 Plack CJ (2014) *The sense of hearing*. New York, USA: Psychology Press.
- 737 Presacco A, Simon JZ, Anderson S (2016a) Evidence of degraded representation of speech in  
738 noise, in the aging midbrain and cortex. *Journal of Neurophysiology* 116:2346-2355.
- 739 Presacco A, Simon JZ, Anderson S (2016b) Effect of informational content of noise on speech  
740 representation in the aging midbrain and cortex. *Journal of Neurophysiology* 116:2356-2367.
- 741 Presacco A, Simon JZ, Anderson S (2019) Speech-in-noise representation in the aging midbrain  
742 and cortex: Effects of hearing loss. *PLoS ONE* 14:e0213899.
- 743 Qiu C, Salvi R, Ding D, Burkard R (2000) Inner hair cell loss leads to enhanced response  
744 amplitudes in auditory cortex of unanesthetized chinchillas: evidence for increased system  
745 gain. *Hearing research* 139:153-171.
- 746 Resnik J, Polley DB (2017) Fast-spiking GABA circuit dynamics in the auditory cortex predict  
747 recovery of sensory processing following peripheral nerve damage. *eLife* 6:e21452.

- 748 Ritz H, Wild CJ, Johnsrude IS (2022) Parametric cognitive load reveals hidden costs in the neural  
749 processing of perfectly intelligible degraded speech. *The Journal of Neuroscience* 42:4619-  
750 4628.
- 751 Rosen S (1992) Temporal Information in Speech: Acoustic, Auditory and Linguistic Aspects.  
752 *Philosophical Transactions: Biological Sciences* 336:367-373.
- 753 Rowland SC, Hartley DEH, Wiggins IM (2018) Listening in Naturalistic Scenes: What Can  
754 Functional Near-Infrared Spectroscopy and Intersubject Correlation Analysis Tell Us About  
755 the Underlying Brain Activity? *Trends in Hearing* 22:2331216518804116.
- 756 Ruhnau P, Herrmann B, Schröger E (2012) Finding the right control: The mismatch negativity  
757 under investigation. *Clinical Neurophysiology* 123:507-512.
- 758 Salvi R, Sun W, Ding D, Chen G-D, Lobarinas E, Wang J, Radziwon K, Auerbach BD (2017) Inner  
759 Hair Cell Loss Disrupts Hearing and Cochlear Function Leading to Sensory Deprivation and  
760 Enhanced Central Auditory Gain. *Frontiers in Neuroscience* 10:Article 621.
- 761 Schilling A, Sedley W, Gerum R, Metzner C, Tziridis K, Maier A, Schulze H, Zeng F-G, Friston KJ,  
762 Krauss P (2023) Predictive coding and stochastic resonance as fundamental principles of  
763 auditory phantom perception. *Brain* 146:4809-4825.
- 764 Schmitt R, Meyer M, Giroud N (2022) Better speech-in-noise comprehension is associated with  
765 enhanced neural speech tracking in older adults with hearing impairment. *Cortex* 151:133-  
766 146.
- 767 Shannon RV, Zeng F-G, Kamath V, Wygonski J, Ekelid M (1995) Speech Recognition with Primarily  
768 Temporal Cues. *Science* 270:303-304.
- 769 Shukla B, Bidelman GM (2021) Enhanced brainstem phase-locking in low-level noise reveals  
770 stochastic resonance in the frequency-following response (FFR). *Brain Research*  
771 1771:147643.
- 772 Sörös P, Treisman IK, Manemann E, Lütkenhöner B (2009) Auditory temporal processing in  
773 healthy aging: a magnetoencephalographic study. *BMC Neuroscience* 10:34.
- 774 Spyridakou C, Rosen S, Dritsakis G, Bamiou D-E (2020) Adult normative data for the speech in  
775 babble (SiB) test. *International Journal of Audiology* 59:33-38.
- 776 Stein RB, Gossen ER, Jones KE (2005) Neuronal variability: noise or part of the signal? *Nature*  
777 *Reviews Neuroscience* 6:389-397.
- 778 Stevens G, Flaxman S, Brunskill E, Mascarenhas M, Mathers CD, Finucane M, on behalf of the  
779 Global Burden of Disease Hearing Loss Expert G (2013) Global and regional hearing  
780 impairment prevalence: an analysis of 42 studies in 29 countries. *European Journal of*  
781 *Public Health* 23:146-152.
- 782 Stocks NG (2000) Suprathreshold Stochastic Resonance in Multilevel Threshold Systems.  
783 *Physical Review Letters* 84:2310-2313.

- 784 Stothart G, Kazanina N (2016) Auditory perception in the aging brain: the role of inhibition and  
785 facilitation in early processing. *Neurobiology of Aging* 47:23-24.
- 786 Synigal SR, Anderson AJ, Lalor EC (2023) Electrophysiological indices of hierarchical speech  
787 processing differentially reflect the comprehension of speech in noise. *BioRxiv*.
- 788 Vanthornhout J, Decruy L, Wouters J, Simon JZ, Francart T (2018) Speech Intelligibility Predicted  
789 from Neural Entrainment of the Speech Envelope. *Journal of the Association for Research in*  
790 *Otolaryngology* 19:181-191.
- 791 Ward LM, Neiman A, Moss F (2002) Stochastic resonance in psychophysics and in animal  
792 behavior. *Biological Cybernetics* 87:91-101.
- 793 Ward LM, MacLean SE, Kirschner A (2010) Stochastic Resonance Modulates Neural  
794 Synchronization within and between Cortical Sources. *PLoS ONE* 5:e14371.
- 795 Wilson RH, McArdle RA, Watts KL, Smith SL (2012) The Revised Speech Perception in Noise Test  
796 (R-SPIN) in a Multiple Signal-to-Noise Ratio Paradigm. *Journal of the American Academy of*  
797 *Audiology* 23:590-605.
- 798 Yasmin S, Irsik VC, Johnsrude IS, Herrmann B (2023) The effects of speech masking on neural  
799 tracking of acoustic and semantic features of natural speech. *Neuropsychologia*  
800 186:108584.
- 801 Zhao Y, Song Q, Li X, Li C (2016) Neural Hyperactivity of the Central Auditory System in Response  
802 to Peripheral Damage. *Neural Plasticity* 2016:2162105.
- 803 Zou J, Feng J, Xu T, Jin P, Luo C, Zhang J, Pan X, Chen F, Zheng J, Ding N (2019) Auditory and  
804 language contributions to neural encoding of speech features in noisy environments.  
805 *NeuroImage* 192:66-75.
- 806 Zuk NJ, Murphy JW, Reilly RB, Lalor EC (2021) Envelope reconstruction of speech and music  
807 highlights stronger tracking of speech at low frequencies. *PLOS Computational Biology*  
808 17:e1009358.
- 809