

18 **Abstract**

19 Despite the distortion of speech signals caused by unavoidable noise in daily life, our ability to comprehend
20 speech in noisy environments is relatively stable. However, the neural mechanisms underlying reliable
21 speech-in-noise comprehension remain to be elucidated. The present study investigated the neural tracking
22 of acoustic and semantic speech information during noisy naturalistic speech comprehension. Participants
23 listened to narrative audio recordings mixed with spectrally matched stationary noise at three signal-to-ratio
24 (SNR) levels (no noise, 3 dB, -3 dB), and 60-channel electroencephalography (EEG) signals were recorded.
25 A temporal response function (TRF) method was employed to derive event-related-like responses to the
26 continuous speech stream at both the acoustic and the semantic levels. Whereas the amplitude envelope of
27 the naturalistic speech was taken as the acoustic feature, word entropy and word surprisal were extracted
28 via the natural language processing method as two semantic features. Theta-band frontocentral TRF
29 responses to the acoustic feature were observed at around 400 ms following speech fluctuation onset over
30 all three SNR levels, and the response latencies were more delayed with increasing noise. Delta-band frontal
31 TRF responses to the semantic feature of word entropy were observed at around 200 to 600 ms leading to
32 speech fluctuation onset over all three SNR levels. The response latencies became more leading with
33 increasing noise and were correlated with comprehension performance and perceived speech intelligibility.
34 While the following responses to speech acoustics were consistent with previous studies, our study revealed
35 the robustness of leading responses to speech semantics, which suggests a possible predictive mechanism
36 at the semantic level for maintaining reliable speech comprehension in noisy environments.

37

38 **Keywords:** speech-in-noise comprehension, semantic processing, neural tracking, temporal response
39 function, EEG

40 **Highlights**

41 1. Leading responses were observed in the semantic-level neural tracking, with more leading latencies as
42 noise increased.

43 2. Following responses were observed in the acoustic-level neural tracking, with more delayed latencies as
44 noise increased.

45 3. Semantic-level neural tracking is correlated with comprehension performance and perceived
46 intelligibility.

47 4. Distinct frequency bands were involved in speech semantic and acoustic processing.

48 **1 Introduction**

49 Noise is an inevitable part of daily life, from car horns on the streets to background music at parties,
50 and it presents a significant challenge to verbal communication. Reliable speech comprehension in noisy
51 environments is crucial in various situations such as education or emergency service. Despite the distortion
52 of auditory information, individuals with normal hearing can comprehend speech with ease. Understanding
53 the adaptive neural mechanisms that enable robust speech-in-noise comprehension is essential for clinical
54 intervention for hearing/language-impaired groups and for developing hearing-aid techniques.

55 Neurophysiological studies have revealed important insights into how noise affects speech processing.
56 Using the event-related techniques, cortical auditory evoked potentials (CAEP) elicited by auditory and
57 speech stimuli have been found to show delayed latencies and reduced amplitudes under adverse conditions,
58 including the early component P1-N1-P2 complex related to primary sound processing (Billings et al., 2009,
59 2011), and the later components such as the N2 component related to phonological analysis (Billings et al.,
60 2009; Martin & Stapells, 2005; Tomé et al., 2015; Whiting et al., 1998) and the P3 component related to
61 speech discrimination (Kaplan-Neeman et al., 2006; Koerner et al., 2017; Martin & Stapells, 2005; Whiting
62 et al., 1998). In recent years, studies have focused more on the neural tracking of continuous speeches. i.e.,
63 the alignment between neural activities and the quasi-rhythmic fluctuations of continuous speech (see
64 reviews, Brodbeck & Simon, 2020; Ding & Simon, 2014; Giraud & Poeppel, 2012; Lakatos et al., 2019;
65 Obleser & Kayser, 2019). Specific temporal dynamics of neural tracking can be described via system
66 identification methods such as the temporal response function (TRF; Crosse et al., 2016, 2021) by relating
67 neural signals with speech features such as acoustic envelope. Neural tracking has been found to remain
68 stable under mild and moderate noise, and it is regarded as an essential tool for segregating speech from
69 the noisy background (Ding & Simon, 2013). Nevertheless, the TRF-based studies have also reported
70 delayed latencies and/or reduced amplitudes of the neural tracking in noisy conditions (Gillis, Decruy, et
71 al., 2022; Mirkovic et al., 2019; Muncke et al., 2022; Zou et al., 2019), similar to previous event-related
72 studies. These results suggest an impaired acoustic processing efficiency in noisy environments (Gillis,
73 Decruy, et al., 2022; Kaplan-Neeman et al., 2006). In addition to auditory processing, semantic processing
74 also plays a vital role in speech-in-noise comprehension and has been paid substantial emphasis.

75 Semantic processing could be a crucial factor in robust speech comprehension against noisy
76 environments. Numerous research has shown that coherent semantic context enabling anticipating
77 upcoming stimuli contributes to an effective understanding of degraded speech (Miller et al., 1951; Obleser
78 & Kotz, 2010, 2011; Sohoglu et al., 2012; Zekveld et al., 2011). For example, Miller et al. (1951) found
79 that words in coherent sentences had higher intelligibility compared with the same words in unrelated word
80 lists during speech-in-noise comprehension. Regarding the influence of noise on semantic processing, such
81 as the N400 component (Kutas & Federmeier, 2011; Kutas & Hillyard, 1984), several studies have reported
82 robust or increased amplitude of N400 under mild degradation, which might be related to additional
83 cognitive effort (Jamison et al., 2016; Romei et al., 2011; Zendel et al., 2015), while other studies reported
84 reduced/delayed N400 for degraded speech, which might be related to damaged signal quality (Aydelott et
85 al., 2006; Connolly et al., 1992; Daltrozzo et al., 2012; Obleser & Kotz, 2011; Strauß et al., 2013). These
86 mixed results provided valuable information on the complex relationship between noise and semantic
87 processing. Moreover, it was discovered that semantic processing includes early responses before the onset
88 of the stimulus, which was considered to be associated with semantic prediction (Grisoni et al., 2017, 2021;
89 Pulvermüller & Grisoni, 2020). Nevertheless, it is still unknown how this pre-onset response is modulated
90 by noise at various signal-to-ratios (SNRs). These inconsistent results and inadequate explorations of noise
91 effect on semantic processing may be due to limitations inherent in the event-related design. This design
92 typically uses highly-controlled and short-duration speech units, such as individual words (e.g., Romei et
93 al., 2011) or disconnected sentences (e.g., Strauß et al., 2013), which only contain limited
94 semantic/contextual information.

95 The recent rise of the naturalistic speech paradigm is expected to expand our knowledge of the neural
96 mechanisms of semantic processing during speech-in-noise comprehension (Z. Li & Zhang, 2023).
97 Compared to the highly-controlled and short-duration speech units, continuous naturalistic speech stimuli
98 provide a better resemblance to our daily communications because of a longer duration, more flexible
99 content, and less deliberate semantic violations (Alday, 2019; Alexandrou et al., 2020; Hartley & Poeppel,
100 2020; Sonkusare et al., 2019; Willems et al., 2020; Wöstmann et al., 2017). Most of all, the continuous
101 naturalistic speech stimuli provide rich context-based semantic information (Alday, 2019; Alexandrou et
102 al., 2020; Hamilton & Huth, 2020; Sonkusare et al., 2019), which is indispensable for semantic prediction

103 and reliable speech comprehension in chaotic daily environments. In addition, via state-of-art
104 computational linguistic models, the semantic information of naturalistic speech can be quantified, and the
105 semantic-level neural tracking can be directly measured (Broderick et al., 2018, 2019, 2021; Gillis et al.,
106 2021; Koskinen et al., 2020; Mesik et al., 2021; Weissbart et al., 2020), presenting a powerful tool to
107 investigating how semantic processing is affected by noise at different SNRs.

108 The two frequently adopted semantic features in speech-related neuroscience research are entropy and
109 surprisal derived from information theory (Brodbeck et al., 2022; Donhauser & Baillet, 2020; Goldstein et
110 al., 2022; Heilbron et al., 2022), which respectively measures the semantic uncertainty of the upcoming
111 stimuli and the unexpectedness of the current stimulus (Pickering & Gambi, 2018; Willems et al., 2016).
112 The word surprisal was found to be associated with the superior temporal gyrus and inferior frontal sulcus,
113 etc. (Willems et al., 2016), and is linked to an N400-like neural response, i.e., negativity at around 400 ms
114 within the central-parietal electrodes (Broderick et al., 2021; Gillis et al., 2021; Heilbron et al., 2022). The
115 word entropy was associated with neural activities within the left ventral premotor cortex, left middle
116 frontal gyrus and right inferior frontal gyrus, etc. (Willems et al., 2016). Furthermore, Goldstein et al. (2022)
117 derived word entropy from deep language models (GPT-2) and correlated them with electrocorticography
118 (ECOG) signals. The results indicated that entropy was related to neural activities in the left-lateralized
119 channels at several hundred milliseconds before the word onset. This pre-onset response is consistent with
120 the semantic prediction potential (SPP) in event-related studies as a direct neural signature for semantic
121 prediction (Grisoni et al., 2021; Pulvermüller & Grisoni, 2020). A recent study by Yasmin et al. (2023)
122 discovered that the N400-like response in semantic-level neural tracking remained robust under mild and
123 moderate noise conditions and declined abruptly at the high-noise level (SNR = -3 dB). However, the noise
124 effect on the pre-onset response in semantic-level neural tracking is still unexplored.

125 The current study aimed to investigate the neural mechanisms of speech-in-noise comprehension by
126 simultaneously focusing on both the acoustic and semantic levels as well as both the pre-onset and the post-
127 onset stages. A naturalistic speech comprehension paradigm was employed, as the naturalistic speech
128 stimuli were expected to provide better ecological validity and contextual information (Alday, 2019;
129 Sonkusare et al., 2019). 60-channel EEGs were recorded while the participants listened to spoken narratives
130 at three SNRs (no noise, 3 dB, -3 dB). Following previous studies, the amplitude envelopes of the speech

131 stimuli were extracted as the acoustic feature (Di Liberto et al., 2015; O’Sullivan et al., 2015). Two typical
132 semantic features were calculated by a Chinese NLP model, i.e., word entropy and word surprisal (Gillis et
133 al., 2021; Weissbart et al., 2020; Willems et al., 2016; Koskinen et al., 2020; Mesik et al., 2021; Broderick
134 et al., 2021). The neural responses to the acoustic and semantic features were estimated using the TRF
135 method (Crosse et al., 2016), which yields the spatiotemporal dynamics of how our brain tracks these
136 features in naturalistic speeches. The pre-onset and post-onset responses in the current study were defined
137 as significant TRF responses with negative and positive time lags, respectively. Especially, we conducted
138 TRF analyses and detected significant TRF responses separately at different SNR levels to capture all
139 potential neural signatures. We hypothesize that the acoustic-level TRF could be related to delayed peak
140 latencies or reduced amplitudes under noisy conditions as in previous studies (Gillis, Decruy, et al., 2022;
141 Mirkovic et al., 2019; Muncke et al., 2022; Zou et al., 2019). As for the semantic-level TRF, we hypothesize
142 that both the pre-onset and post-onset response could show resilience against noise (Yasmin et al., 2023).
143 By exploring the pre-onset and post-onset temporal dynamics of low- and high-level processing, this study
144 hopes to gain a more complete overview of the noise effect on neural processing during naturalistic speech
145 comprehension.

146 **2 Methods**

147 **2.1 Participants**

148 Twenty college students (10 females, ages ranging from 19 to 28 years old) participated in the study.
149 The sample size was determined to be sufficient by reference to previous TRF-based studies on the human
150 speech processing (e.g., Broderick et al., 2018; Di Liberto et al., 2015). One male participant was excluded
151 due to technical problems during data recording. The data of the remaining nineteen participants (age: mean
152 \pm SD = 21.79 ± 1.99) were included in the subsequent analyses. All participants were native Chinese
153 speakers, right-handed, with normal hearing and normal or corrected-to-normal vision by self-report. The
154 study was conducted in accordance with the Declaration of Helsinki and was approved by the local Ethics
155 Committee of Tsinghua University. Written informed consent was obtained from all participants.

156

157 **2.2 Materials**

158 Thirty narrative audio recordings from our previous studies (Z. Li et al., 2021, 2022) were used as
159 stimuli. These audio recordings were recorded from six native Chinese speakers with professional training
160 in broadcasting. The participants were unfamiliar with the content of these narrative audio recordings,
161 which were about speakers' personal experiences on daily-life topics adapted from the National Mandarin
162 Proficiency Test. Each narrative audio recording lasted for around 90 s and was recorded by a regular
163 microphone at a sampling rate of 44,100 Hz in a sound-attenuated room.

164 These speech stimuli were further processed into three versions at three different SNR levels: no-noise
165 (NN), low-noise (SNR = 3 dB), and high-noise (SNR = -3 dB), where speech intensity percentage was
166 100%, 60%, and 40%, respectively. This procedure was achieved by adding spectrally matched stationary
167 noise, which was generated based on a 50th-order linear predictive coding (LPC) model estimated from the
168 original speech recording (Broderick et al., 2018). The SNR levels were selected following previous studies
169 (Ding & Simon, 2013), and were produced by varying the noise intensity while keeping the intensity of
170 original speech (measured by its root mean square) constant (Ding & Simon, 2013).

171 For each narrative audio recording, two four-choice questions were prepared by the experimenters to
172 assess one's speech comprehension performance. These questions and the corresponding choices were
173 targeted at detailed narrative contents that would demand significant attentional efforts. For instance, one

174 question following a narrative audio recording about one's major was, "What is the speaker's most likely
175 major as a graduate student? (说话人的研究生专业最可能是什么?)" and the four choices were 1) Social
176 science, 2) International politics, 3) Pedagogy and 4) Psychology (1. 社会科学, 2. 国际政治, 3. 教育学 and
177 4. 心理学).

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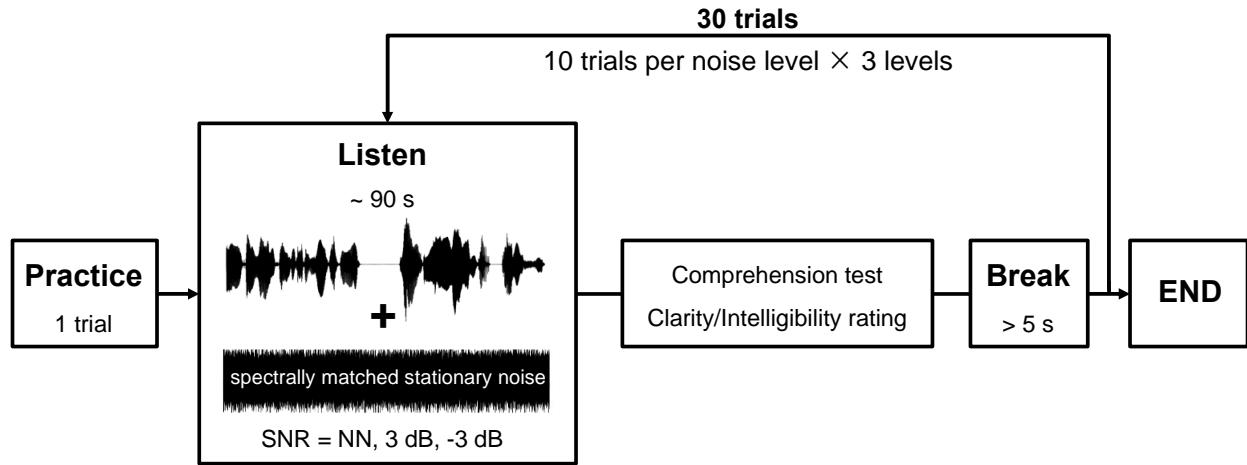
179 **2.3 Procedure**

180 Before the start of the experiment, the participants had one practice trial to get familiar with the
181 procedure, with an additional narrative audio recording at the no-noise level not used in the formal
182 experiment. The formal experiment consisted of 30 trials, with 10 trials per SNR level. In each trial, the
183 participants listened to narrative audio recordings at one of the three SNR levels. The participants were
184 required to maintain visual fixation on a fixation cross displayed on the computer screen in front of them
185 and to minimize eye blinks and all other motor activities during listening. The order of the narrative audio
186 recordings and their assigned SNR levels was randomized for each participant.

187 After each trial, the participants were instructed to answer two four-choice questions about the content
188 of the narrative audio recording using the computer keyboard. The averaged accuracies across all trials
189 (separately for each SNR level) were used to reflect the participants' comprehension performance. After
190 completing these questions, the participants were instructed to rate the perceived clarity and intelligibility
191 of the narrative audio recording on a 100-point rating scale and rested for at least 5 s before moving on to
192 the next trial. No feedback was given to the participants about their comprehension performance during the
193 experiment.

194 The experimental procedure was programmed in MATLAB using the Psychophysics Toolbox 3.0
195 (Brainard, 1997). The speech stimuli were delivered to listeners seated in a sound-attenuated room via an
196 air-tube earphone (Etymotic ER2, Etymotic Research, Elk Grove Village, IL, USA) to avoid environmental
197 noise and equipment electromagnetic interference. The volume of the audio stimuli was adjusted

198 individually for each participant to a comfortable level, and it was kept consistent across trials. The
199 experimental procedure is illustrated in Figure 1.



200 **Figure 1.** Experimental procedure. The participants listened to 30 naturalistic narrative audio recordings
201 which each lasted around 90 s. These audio recordings were mixed with three levels of spectrally matched
202 stationary noise: no-noise (NN), low-noise (SNR = 3 dB), and high-noise (SNR = -3 dB). The 60-channel
203 EEG signals were recorded during listening. After listening to each narrative audio recording, the
204 participants were required to complete a comprehension test and report the clarity and intelligibility rating.
205 In the comprehension test, two four-choice questions per audio recording based on the narrative content
206 were used.

207 2.4 EEG recording and preprocessing

EEG signals were recorded from 60 channels with a NeuroScan amplifier (SynAmp II, NeuroScan, Compumedics, USA) at a sampling rate of 1000 Hz. Electrodes were positioned according to the international 10–20 system, including FP1/2, FPZ, AF3/4, F7/8, F5/6, F3/4, F1/2, FZ, FT7/8, FC5/6, FC3/4, FC1/2, FCZ, T7/8, C5/6, C3/4, C1/2, CZ, TP7/8, CP5/6, CP3/4, CP1/2, CPZ, P7/8, P5/6, P3/4, P1/2, PZ, PO7/8, PO5/6, PO3/4, POZ, Oz, O1/2, referenced to an electrode between CZ and CPZ with a forehead ground at FZ. Electrode impedances were kept below 10 kOhm for all electrodes throughout the experiment.

214 The recorded EEG data were first notch filtered to remove the 50 Hz powerline noise. Independent
215 Component Analysis (ICA) was performed to remove artifacts such as eye blinks and eye movements based
216 on visual inspection. Around 4–12 independent components (ICs; mean = 6.6) were removed per participant.
217 The remaining ICs were then back-projected onto the scalp EEG channels to reconstruct the artifact-free
218 EEG signals. The EEG signals were then re-referenced to the average of all scalp channels and
219 downsampled to 128 Hz. Afterward, EEG signals were filtered into the delta (1–4 Hz) and theta (4–8 Hz)

220 bands, which have been previously reported to be important for speech neural tracking (Ding et al., 2014;
221 Etard & Reichenbach, 2019; Keitel et al., 2017; Koskinen et al., 2020; J. Li et al., 2023; Peelle et al., 2013).
222 For comprehensiveness, we also included the alpha (8–12 Hz) and beta (12–30 Hz) bands into analyses.
223 Causal FIR (Finite Impulse Response) filters were employed to ensure that the filtered EEG signals were
224 determined only by the current and previous data samples, which was important for the present study
225 focusing on the fine-grained time course, particularly considering the pre-onset neural responses (de
226 Cheveigné & Nelken, 2019).

227 These preprocessed EEG signals were segmented into 30 trials, from 5 to 90 s (duration = 85 s),
228 relative to the speech onsets of each trial to avoid possible onset and offset effects. All preprocessing was
229 conducted offline using MATLAB and the Fieldtrip toolbox (Oostenveld et al., 2011).

230

231 **2.5 Feature extraction**

232 Three types of features were extracted to represent the acoustic (amplitude envelope) and semantic
233 (word entropy, word surprisal) information for each narrative audio recording. An example of these speech
234 features is illustrated in Figure 2A.

235 *Acoustic feature.* The amplitude envelope for each narrative audio recording was calculated as the
236 absolute values after a Hilbert transform and then downsampled to the sampling rate of 128 Hz to match
237 that of the EEG signals.

238 *Semantic features.* Before feature extraction, the narrative audio recordings were converted to text by
239 Iflyrec software (Iflytek Co., Ltd, Hefei, Anhui) and then segmented into words based on the THU Lexical
240 Analyzer for Chinese (THULAC) toolbox (Sun et al., 2016).

241 Two semantic features, word entropy and word surprisal, were extracted. Word entropy measures the
242 uncertainty of predicting the upcoming word based on the context so far and was calculated as equation (1):

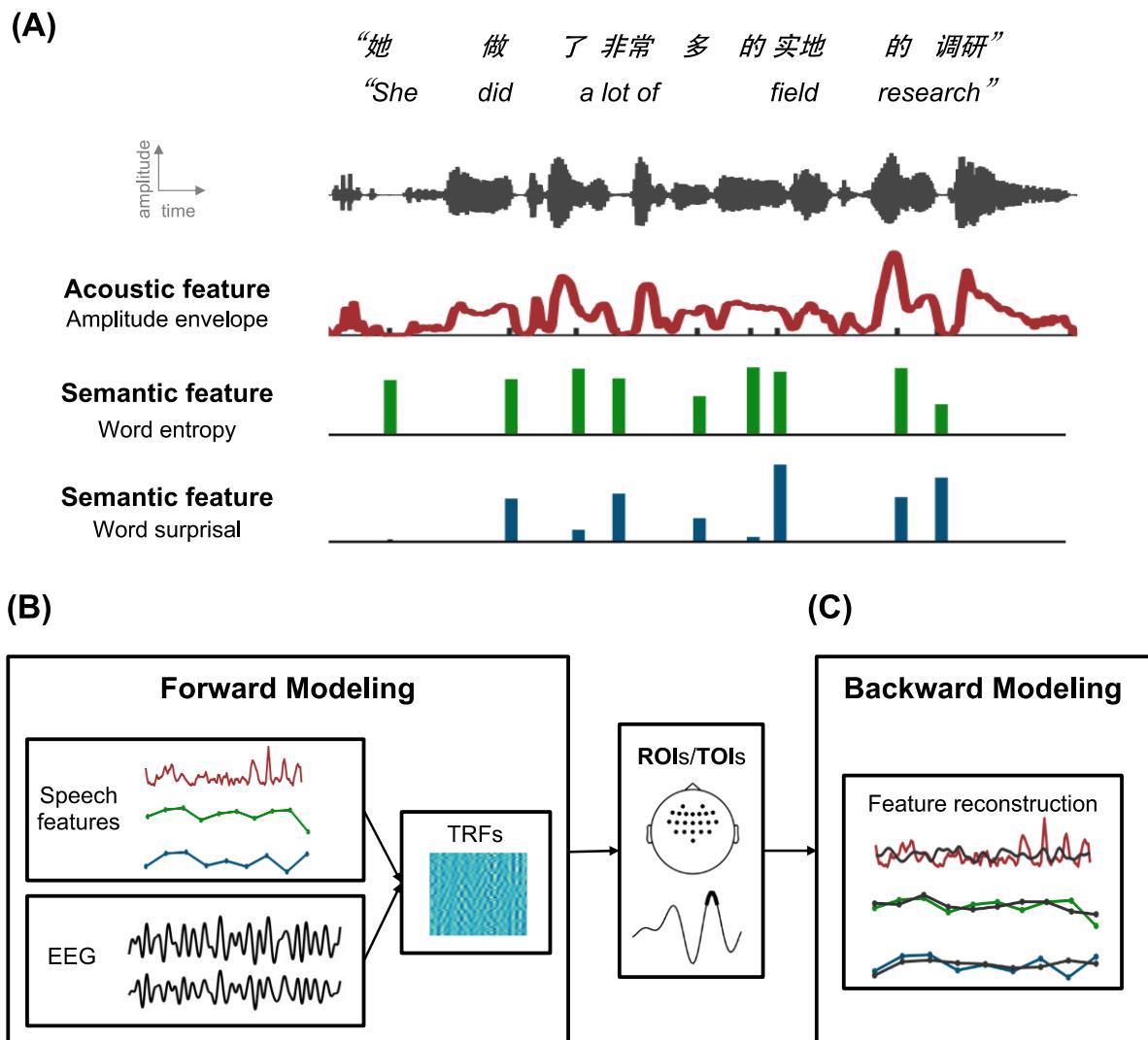
$$243 \text{Entropy}(t) = - \sum_{w_{t+1}} P(w_{t+1}|w_1, \dots, w_t) \log P(w_{t+1}|w_1, \dots, w_t) \quad (1)$$

244 Word surprisal measures how surprising the current word is given the previously encountered words and
245 was calculated as equation (2):

$$246 \text{Surprisal}(t) = - \log P(w_t|w_1, \dots, w_{t-1}) \quad (2)$$

247 Where w_1, \dots, w_{t-1} are the existing word sequence and $P(w_t|w_1, \dots, w_{t-1})$ is the conditional probability of
248 next word (Willems et al., 2016). These NLP calculations were conducted by ADAM, a widely accepted
249 Long-Short Term Memory (LSTM) Neural Network model (Kingma & Ba, 2015). The model was trained
250 on the corpora corpus of the People's Daily with 534,246 Chinese words. See Supplementary Table S1 for
251 more information about the model and Supplementary Table S2 for more information about the descriptive
252 statistics of semantic features.

253 After extracting the semantic features of each word, the word onset timings were estimated via Iflyrec
254 software. Impulses at the word onset time were manipulated with corresponding semantic feature values to
255 generate one-dimensional “semantic vectors” (e.g., Broderick et al., 2018; Gillis et al., 2021). The sampling
256 rate of the semantic vectors was 128 Hz to match the EEG signals.



257 **Figure 2.** Speech feature extraction and Temporal Response Function analyses. (A) Three types of speech
258 features were extracted, including one acoustic feature (amplitude envelope) and two semantic features

259 (word entropy and word surprisal). The two semantic features were derived from a computational linguistic
260 model and one-dimensional vectors were generated with impulses manipulated with semantic feature values
261 at the corresponding word onsets time. (B) Forward modeling. TRFs were extracted by regressing each of
262 the three types of speech features against the EEG signals separately. The significance of these TRFs was
263 estimated by comparing them with the corresponding control TRFs, which were modeled based on EEG
264 signals and shuffled speech features. The resulting spatiotemporal ranges were identified as regions of
265 interest (ROIs) and time lags of interest (TOIs). (C) Backward modeling. These three types of speech
266 features were separately reconstructed through backward TRF models, and the reconstruction accuracy
267 (Pearson's r) depicted the strength of neural tracking. Control backward models were constructed with EEG
268 signals and shuffled features.

269 **2.6 Modeling of the stimulus-response relationship**

270 The Temporal Response Function (TRF) modeling method based on ridge regression was adopted to
271 explore the relationship between the neural activities and the three types of stimulus features (Crosse et al.,
272 2016, 2021). Forward modeling was first used to illustrate the specific spatiotemporal response patterns
273 and identify key electrodes and time lags in TRF responses of the corresponding speech feature, and then
274 backward modeling was adopted to verify the possible contribution of these identified neural correlates
275 (e.g., Broderick et al., 2019; Etard & Reichenbach, 2019). The overall procedure of the modeling analyses
276 is shown in Figure 2B and 2C.

277 *Forward modeling.* With a forward modeling approach, we described neural response patterns to
278 different speech features by linear spatiotemporal filters called TRFs, which measure how neural signals
279 from different regions are modulated by stimulus features at different time lags (Crosse et al., 2016). The
280 estimated TRF together with the corresponding speech feature was used to predict the EEG responses from
281 each electrode. The prediction accuracy measured as the Pearson's correlation between the actual and
282 predicted EEG signals represents the performance of the forward model. The TRF, w , is measured by
283 equation (3):

$$284 \quad w = (S^T S + \lambda I)^{-1} S^T r \quad (3)$$

285 Where S is the lagged time series of the stimulus features, r is the neural signals, and I is the identity
286 matrix. The time lags for forward modeling were chosen to cover a relatively broad time range, from -1000
287 to 1000 ms (Goldstein et al., 2022; J. Li et al., 2021, 2023). The λ is the regularization parameter used to
288 prevent overfitting and ranged between 0.1 and 1000 in steps of the powers of 10 empirically (Gillis, Van

289 Canneyt, et al., 2022). The cross-validation procedure was implemented in a leave-one-trial-out procedure
290 within each participant: each time, the model was trained based on 9 trials and tested on the left-out trial,
291 which was repeated for each of the 10 trials at three SNR levels separately. The λ value that produced the
292 highest prediction accuracy averaged across trials after cross-validation was selected as the regularization
293 parameter for all trials at a certain SNR level per participant. TRF amplitudes were further transformed into
294 z -scores before statistical analyses (Ding et al., 2014; Gillis et al., 2021; J. Li et al., 2021).

295 The statistical significance of the estimated TRFs was estimated by constructing control TRF models
296 (Weissbart et al., 2020). We built control models by constructing TRF models using shuffled stimulus
297 features and the EEG recordings in the same way as for the computation of the actual TRFs. The shuffled
298 amplitude envelope was constructed by randomly shuffling the feature value within a trial while keeping
299 the timing of the quiet fragments. The shuffled word entropy and word surprisal were constructed by
300 randomly shuffling the feature value within a trial while keeping the timing of impulses. Therefore the
301 speech features that described acoustic and linguistic word onsets were not altered and thus left no impact
302 on TRFs' significance (Weissbart et al., 2020). The shuffling was repeated 1,000 times and resulted in
303 1,000 control TRFs for a corresponding actual TRF.

304 A nonparametric cluster-based permutation test was applied to account for multiple comparisons
305 (Maris & Oostenveld, 2007). For each electrode-time bin in the actual and control TRFs, a one-sample t -
306 test was used to examine whether the TRF amplitudes significantly differed from 0. Then neighboring
307 electrode-time bins with an uncorrected p -value less than 0.01 were combined into clusters. The minimum
308 number of neighboring significant channels that was required for inclusion in a cluster was 2. For each
309 cluster, the sum of the t -statistics was obtained. A null distribution was created from the 1,000 control test
310 statistics, i.e., the maximum cluster-level t -statistics. The corrected p -value for each cluster was calculated
311 as the proportion of control test statistics greater than the actual cluster-level t -statistics. Clusters with p -
312 values below 0.05 were selected for further analyses. We implemented the same statistical analyses
313 procedure for each of the 18 TRFs (3 stimulus features \times 2 frequency bands \times 3 SNR levels). The EEG
314 electrodes and time lags from significant clusters were regarded as ROIs/TOIs. Then peaks were estimated
315 within these ROIs/TOIs, and the peak amplitudes and peak latencies were compared across different SNRs.

316 *Backward modeling.* With a backward modeling approach, we simultaneously took neural signals from
317 several electrodes to reconstruct stimulus features with a decoder. The reconstruction accuracy was
318 measured as the Pearson’s correlation between the actual and reconstructed stimulus features. The decoder,
319 g , is calculated by equation (4), where R is the lagged time series of the EEG data. The reconstructed feature,
320 $\hat{s}(t)$, is calculated by equation (5) where n is the EEG electrodes, and τ is the time lags (Broderick et al.,
321 2019).

$$g = (R^T R + \lambda I)^{-1} R^T s \quad (4)$$

$$\hat{s}(t) = \sum_n \sum_{\tau} r_n(t + \tau) g_n(\tau) \quad (5)$$

Only the exact ROIs/TOIs from significant clusters found in the forward modeling were included in the backward modeling. The EEG signals and stimulus features were downsampled to 64 Hz for better computational efficiency. The same leave-one-trial-out cross-validation procedure as in the forward modeling approach was conducted to obtain the optimal regularization parameter and calculate the reconstruction accuracy. We also estimated the control decoders using the same shuffling methods in forward modeling. The reconstruction accuracies from the 1,000 control decoders were averaged and compared with the actual decoder via a one-tailed paired t -test, and the p values of clusters were corrected via the false discovery rate (FDR) method (Benjamini & Hochberg, 1995).

332 In sum, the analyses of neural tracking followed two main steps. (1) We calculated the actual forward
333 model and control forward models and identified ROIs/TOIs according to clusters with significant
334 differences between them. (2) We estimated the reconstruction accuracy based on these ROIs/TOIs. This
335 procedure resulted in (1) the specific spatiotemporal TRF response pattern and (2) the strength of neural
336 tracking (reconstruction accuracy) for analyses.

Given that the neural signatures of speech processing could exhibit different spatiotemporal patterns at various SNR levels (e.g., Bidelman & Howell, 2016; Billings et al., 2009; Strauß et al., 2013), we conducted separate statistical tests for identifying significant clusters in the TRF responses at different SNR levels, in order to capture all potentially significant results without missing anything.

341 We classified these significant clusters into two types based on their spatiotemporal dynamics: those
342 with largely overlapped spatiotemporal patterns across all SNR levels, which could represent a reliable
343 response across all SNR levels, and those with unique patterns at a certain SNR level, which might signify

344 distinct processing mechanisms under certain circumstances. This was achieved by visual inspection and
345 calculating a similarity index, which derived from the average of the temporal and spatial similarity. See
346 Supplementary Figure S2 for more information. The consistent clusters were compared to explore how the
347 commonly shared neural signature adapted to noise, while the unique clusters received less attention in
348 subsequent analyses. Linear-mixed effect (LME) models and Spearman's correlation was conducted to
349 examine the relationship between neural tracking and behavioral performance.

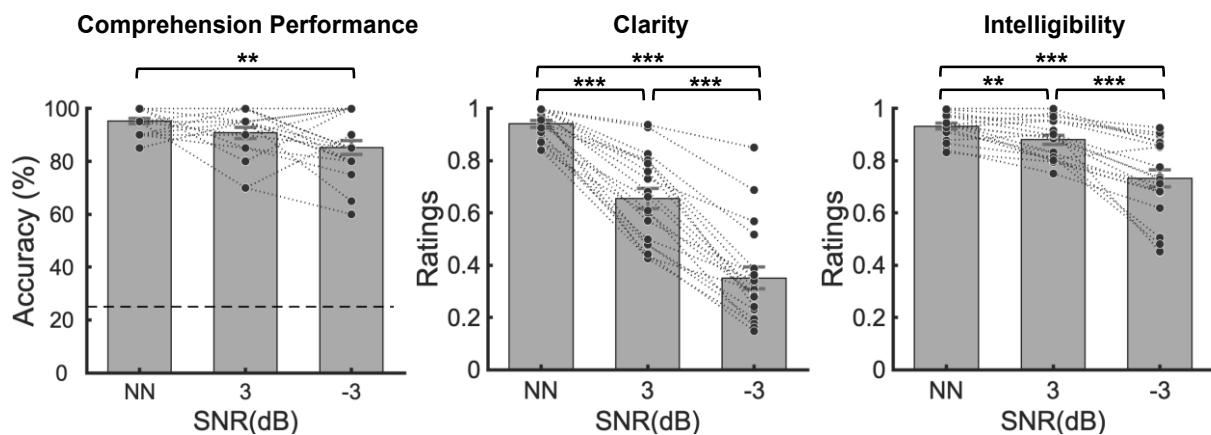
350 Before modeling, the three types of stimulus features across all trials and EEG signals across all
351 channels were z-scored as recommended to ensure consistent scaling (Crosse et al., 2016, 2021). Modeling
352 and analyses for different stimulus features were conducted independently. Considering we focused on the
353 neural tracking of underlying hierarchical information in speech rather than physical stimulus, we adopted
354 the same stimulus features of no-noise speech for the other two SNR levels (Ding & Simon, 2013; Fuglsang
355 et al., 2017). The forward and backward modeling was conducted in MATLAB using the Multivariate
356 Temporal Response Function (mTRF) toolbox (Crosse et al., 2016). The cluster-based permutation test was
357 conducted in the FieldTrip toolbox (Oostenveld et al., 2011). Other statistical analyses were conducted via
358 MATLAB functions and IBM SPSS Statistics software (IBM corp., 2019).

359 **3 Results**

360 **3.1 Behavioral performance**

361 The speech comprehension performance was measured as the averaged response accuracy of the four-
362 choice questions and was found to be significantly different among the three SNR levels (rmANOVA, $F(2,$
363 $36) = 6.74, p = .003$). The speech comprehension performance was $95.26 \pm 1.05\%$, $90.79 \pm 2.10\%$, and
364 $85.26 \pm 2.63\%$ (mean \pm SE) at the no-noise level, low-noise level, and high-noise level, respectively. The
365 comprehension performance at the high-noise level was significantly lower than that at the no-noise level
366 (post-hoc t -test, $p = .006$, Bonferroni corrected). In addition, it should be noted that the comprehension
367 performance was still well above chance level even at the high-noise level (one-tailed t -test, $t(18) = 22.88$,
368 $p < .001$).

369 The subjective ratings of clarity and intelligibility showed a similar pattern with significant differences
370 among the SNR levels (rmANOVA, $F(2, 36) = 148.32$ and $35.31, ps < .001$). The normalized clarity rating
371 scores were 0.94 ± 0.01 , 0.66 ± 0.04 , and 0.35 ± 0.04 (mean \pm SE), and the normalized intelligibility rating
372 scores were 0.93 ± 0.01 , 0.88 ± 0.02 , and 0.73 ± 0.03 (mean \pm SE) at the no-noise, low-noise, and high-
373 noise level, respectively. Post hoc t -tests revealed significant pairwise differences for all possible
374 comparisons ($ps < .01$, Bonferroni corrected). The behavioral performance is illustrated in Figure 3. These
375 results suggested that the effect of noise on speech comprehension and perception was effectively
376 manipulated.

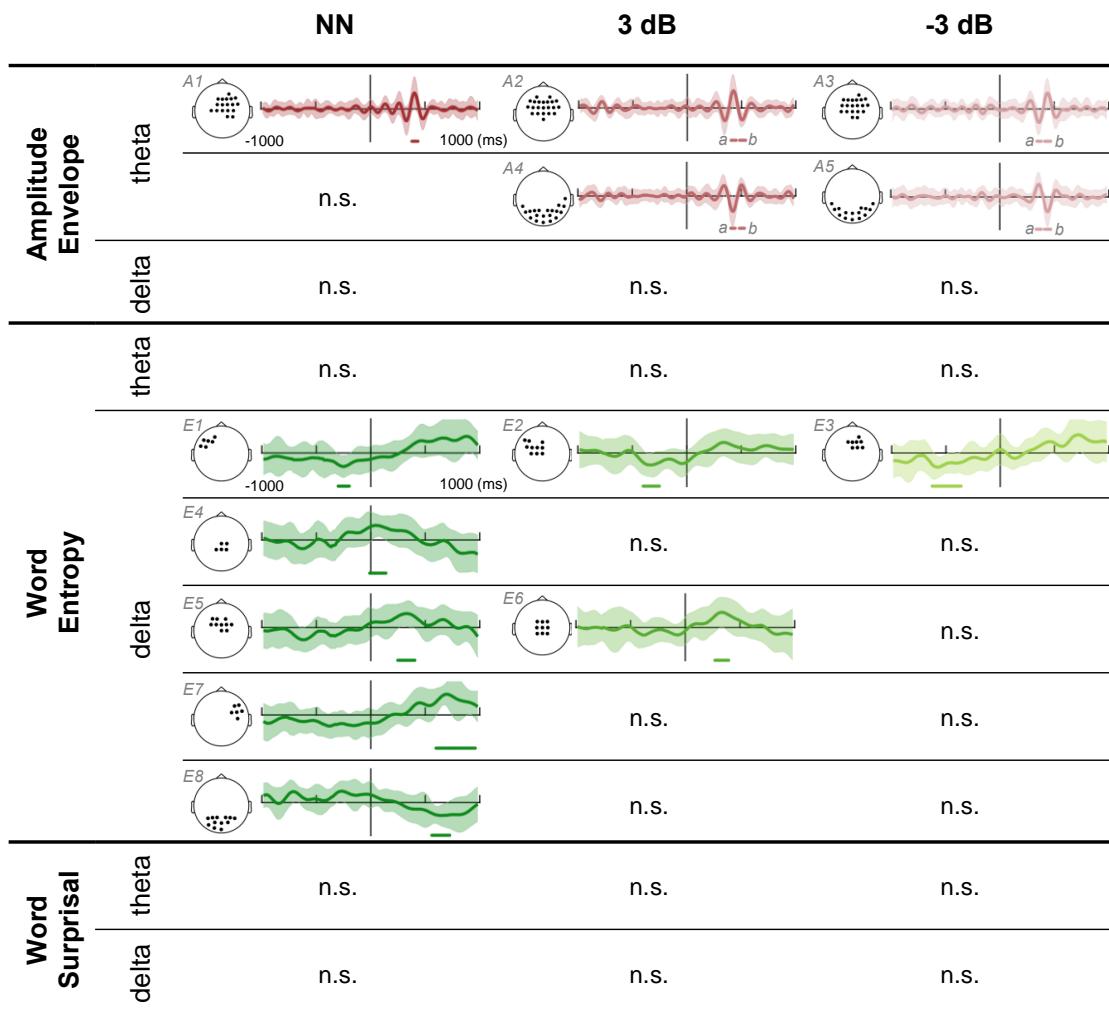


377 **Figure 3.** Behavioral results. Black dots indicate data points from each participant. Error bars denote the
378 standard error. *: $p < .05$, **: $p < .01$, ***: $p < .001$.

380 **3.2 Summary of all significant clusters in the acoustic- and semantic-level TRF responses**

381 We summarized all significant clusters in the acoustic- and semantic-level TRF responses in Figure 4.
382 Significant clusters were only found in the delta/theta bands but not alpha/beta bands (see Supplementary
383 Figure S1 for more information). The specific time lags of TOIs of clusters were listed in Supplementary
384 Table S3-4. According to visual inspection and the similarity index (shown in Supplementary Figure S2),
385 these significant clusters were classified into responses that exhibited relative consistency across different
386 SNR levels, as well as distinctive response at a certain SNR level.

387 Clusters with largely overlapped spatiotemporal patterns across all SNR levels were found in theta-
388 band acoustic-level TRFs (i.e., A1, A2, A3) and delta-band entropy-based semantic-level TRFs (i.e., E1,
389 E2, E3). Detailed analyses of them are demonstrated in the section 3.3 and 3.4, respectively. Several clusters
390 with similar spatiotemporal patterns shared by certain SNR levels, such as the acoustic-level TRF response
391 within the occipital electrodes (i.e., A4 and A5) and the post-onset entropy-based semantic-level TRF
392 responses within the central electrodes (i.e., E5 and E6). Analyses of them are demonstrated in
393 Supplementary Figure S3 and S4. No further analysis was done for the other unique clusters at the no-noise
394 level. No significant TRF responses to word surprisal were found.

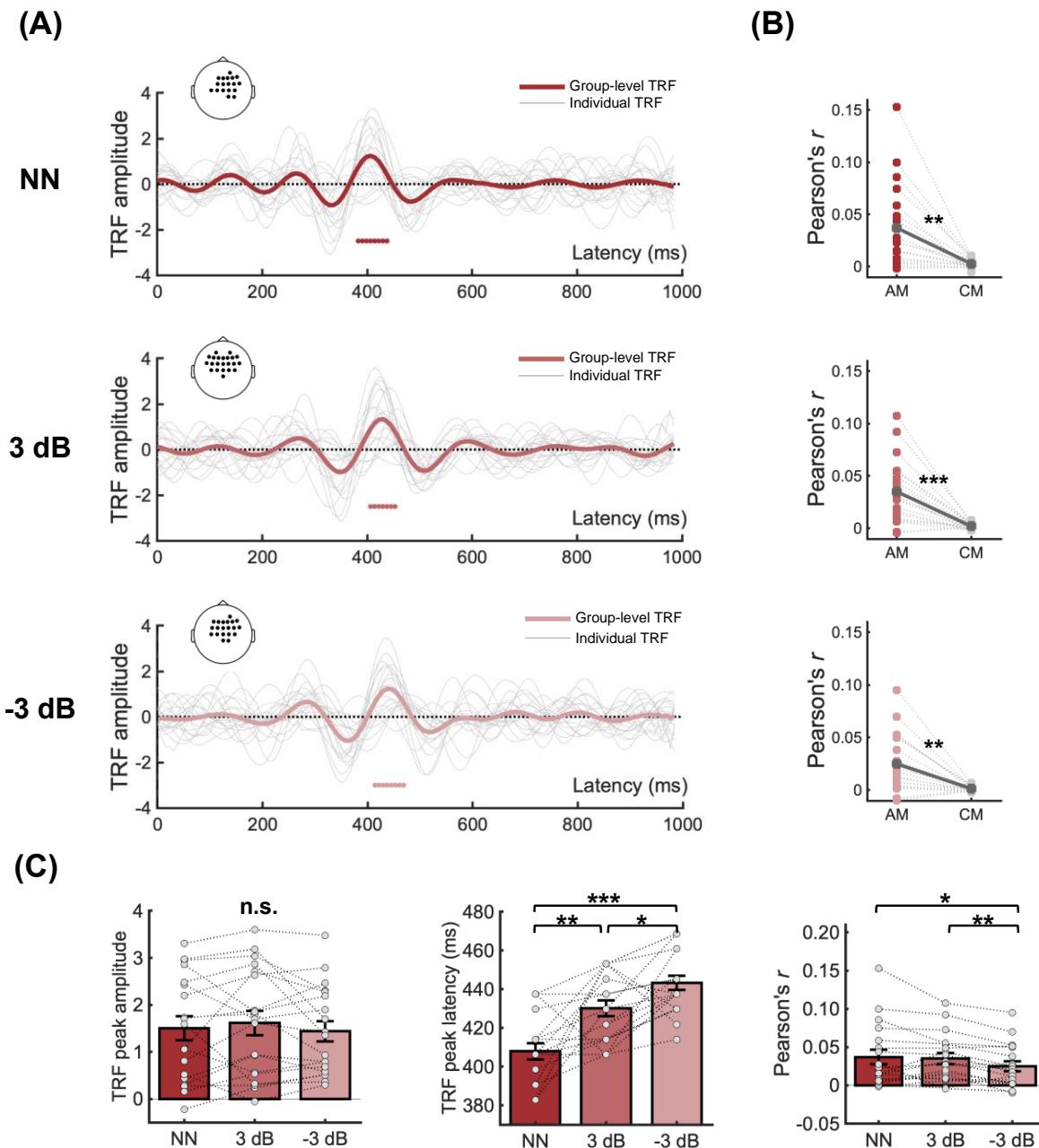


395 **Figure 4.** Significant clusters in TRF responses to (A) amplitude envelope, (B) word entropy, and (C) word
396 surprisal at different SNR levels and frequency bands. Significant clusters are numbered A1–A5 and E1–E8.
397 Clusters with similar spatiotemporal patterns are organized in the same row. ROIs of clusters are shown as
398 black dots in the corresponding topography. The colored curves are the mean of TRFs averaged among
399 ROIs across participants. The shaded areas denote the standard error of TRFs. The colored horizontal line
400 below the TRF curve indicates the TOIs of the cluster. The *a* and *b* refer to clusters with the same ROIs but
401 different TOIs. n.s.: no significant cluster was found.

402 **3.3 Acoustic-level TRF responses with delayed latencies as noise increases**

403 Significant acoustic-level TRF responses in the theta band were found at all SNR levels and showed a
404 similar positivity within central electrodes at around 300~500 ms (i.e., A1, A2a, A3b), as demonstrated in
405 Figure 5A. At the no-noise level, the TRF showed positivity in the central electrodes with a latency of
406 around 400 ms (cluster-level $p < .01$). At the low-noise and high-noise levels, the TRF showed similar
407 positivity in the central electrodes with a latency of around 430 (cluster-level $p < .01$) and 440 ms (cluster-
408 level $p < .01$). We estimated the peak amplitude and peak latency for the positive peak at each SNR level.
409 The peak latencies were significantly different among the three SNR levels (rmANOVA, $F(2, 36) = 21.42$,
410 $p < .001$), and post-hoc *t*-tests revealed significantly delayed peak latencies as noise increased ($ps < .05$,
411 Bonferroni corrected), as shown in Figure 5C. No significant differences were found in the peak amplitudes
412 (rmANOVA, $F(2, 36) = 0.64$, $p = .535$).

413 The reconstruction accuracies from the corresponding ROIs/TOIs were significantly higher in the
414 actual decoders than in the control decoders ($p < .01$, FDR corrected, Figure 5B). Comparing the
415 reconstruction accuracies revealed significant differences among the three SNR levels (rmANOVA, $F(2,$
416 $36) = 6.19, p = .010$ with Greenhouse-Geisser correction), and post-hoc t -tests revealed significantly
417 weakened neural tracking at the high-noise level compared with the no-noise level and the low-noise level
418 ($p < .05$, Bonferroni corrected, Figure 5C).



419 **Figure 5.** Acoustic-level TRF responses in the theta band at different SNR levels. (A) The bold curves in
420 different shades of red are the mean of TRFs averaged among ROIs across participants at the three SNR
421 levels. The grey curves are TRFs averaged among the ROIs of each participant. The colored horizontal line

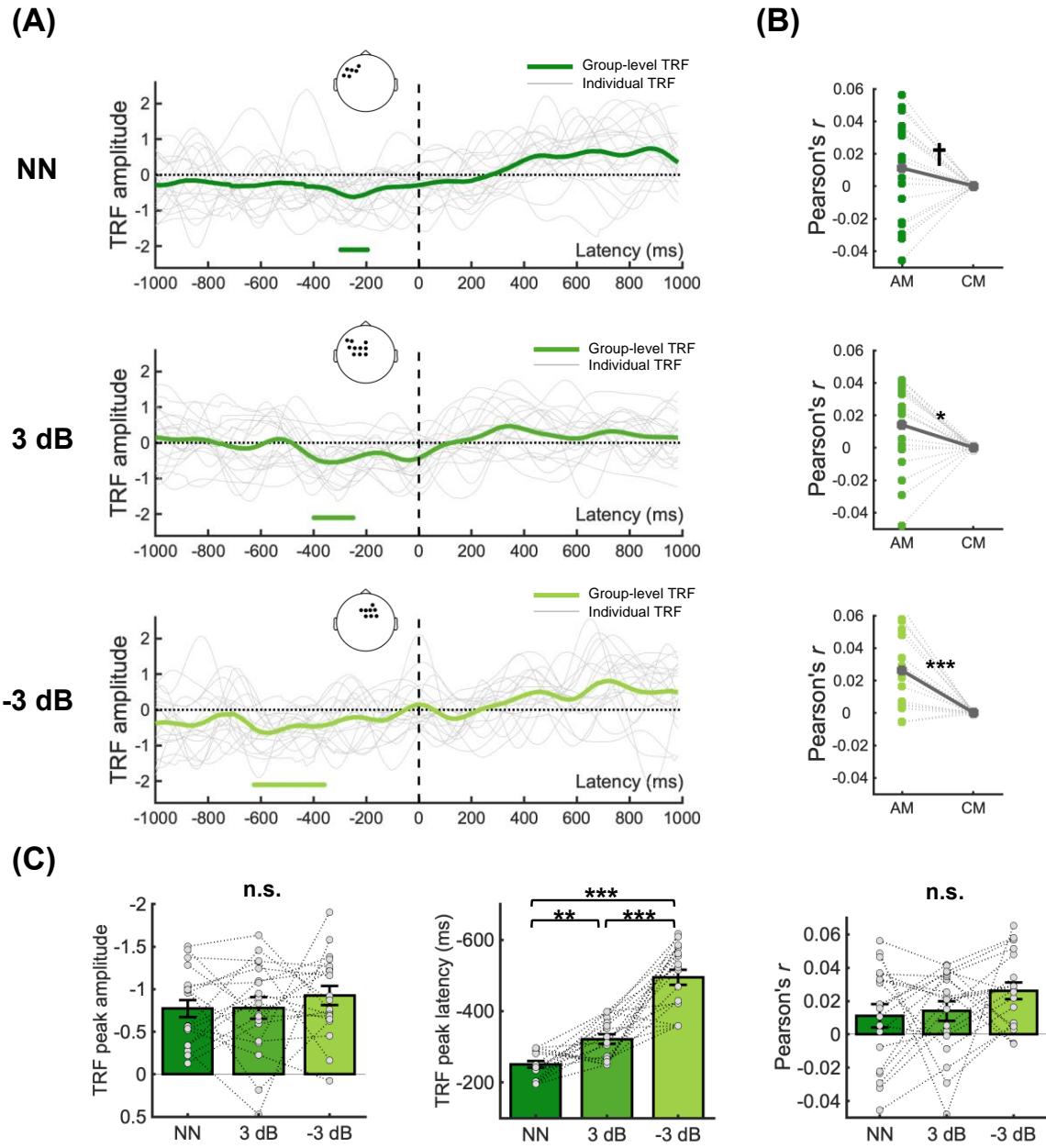
422 at the bottom of each plot indicates TOIs over which TRFs differed significantly from the control models.
423 Dots in the corresponding topographies depicted the ROIs. (B) Reconstruction accuracy calculated from
424 the ROIs/TOIs in (A). AM means actual models. CM means control models. (C) Noise effect on the peak
425 amplitude, peak latency, and reconstruction accuracy. Grey dots indicate data points from each participant.
426 Error bars denote the standard error. n.s.: not significant, *: $p < .05$, **: $p < .01$, ***: $p < .001$.

427 **3.4 Semantic-level TRF responses with earlier latencies as noise increases**

428 Significant semantic-level TRF responses to word entropy in the delta band were found at all SNR
429 levels. They showed similar negativity at around 200~600 ms leading to speech fluctuation onset, as
430 demonstrated in Figure 6A. The time lags of pre-onset processing to word entropy showed a gradual
431 advanced trend as noise increased. The time lag was approximately from around -300 ms to -180 ms at the
432 no-noise level (cluster-level $p < .05$) and was from around -400 ms to -250 ms at the low-noise level
433 (cluster-level $p < .05$), from around -630 ms to -360 ms at the high-noise level (cluster-level $p < .01$). We
434 estimated the peak amplitude and peak latency for the pre-onset negative peak at each SNR level. The peak
435 latencies were significantly different among the three SNR levels (rmANOVA, $F(2, 36) = 58.08, p < .001$),
436 and post-hoc t -tests revealed that as noise increased the peak latencies were gradually earlier ($p < .05$,
437 Bonferroni corrected), as shown in Figure 6C. No significant differences were found in the peak amplitudes
438 (rmANOVA, $F(2, 36) = 0.63, p = .538$).

439 The pre-onset TRF responses to word entropy exhibited different spatiotemporal patterns at three SNR
440 levels. At the no-noise level, the ROIs included frontal-parietal electrodes and exhibited obvious left
441 lateralization. At the low-noise level, the ROIs showed similar left-lateralized topological distribution but
442 included more electrodes, while at the high-noise level, no obvious lateralization was observed in the
443 frontal-parietal ROIs.

444 The reconstruction accuracies from corresponding ROIs/TOIs were significantly higher in the actual
 445 decoders than in the control decoders at both the low-noise level ($p < .05$, FDR corrected) and the high-
 446 noise level ($p < .001$, FDR corrected), but only marginally significant at the no-noise level ($p = .073$, FDR
 447 corrected), as shown in Figure 6B. Comparing the reconstruction accuracies among different SNR levels
 448 revealed no significant differences (rmANOVA, $F(2, 36) = 1.64$, $p = .208$, Figure 6C).



449 **Figure 6.** Semantic-level TRF responses to word entropy in the delta band at different SNR levels. (A) The
 450 bold curves in different shades of green are the mean of TRFs averaged among ROIs across participants at
 451 the three SNR levels. The grey curves are TRFs averaged among the ROIs of each participant. The colored
 452 horizontal line at the bottom of each plot indicates TOIs over which TRFs differed significantly from the
 453 control models. Dots in the corresponding topographies depicted the ROIs. (B) Reconstruction accuracy

454 calculated from the ROIs/TOIs in (A). AM means actual models. CM means control models. (C) Noise
455 effect on the peak amplitude, peak latency, and reconstruction accuracy. Grey dots indicate data points from
456 each participant. Error bars denote the standard error. n.s.: not significant, †: $p < .1$, *: $p < .05$, **: $p < .01$,
457 ***: $p < .001$.

458 **3.5 Correlation between TRF responses and behavioral performance**

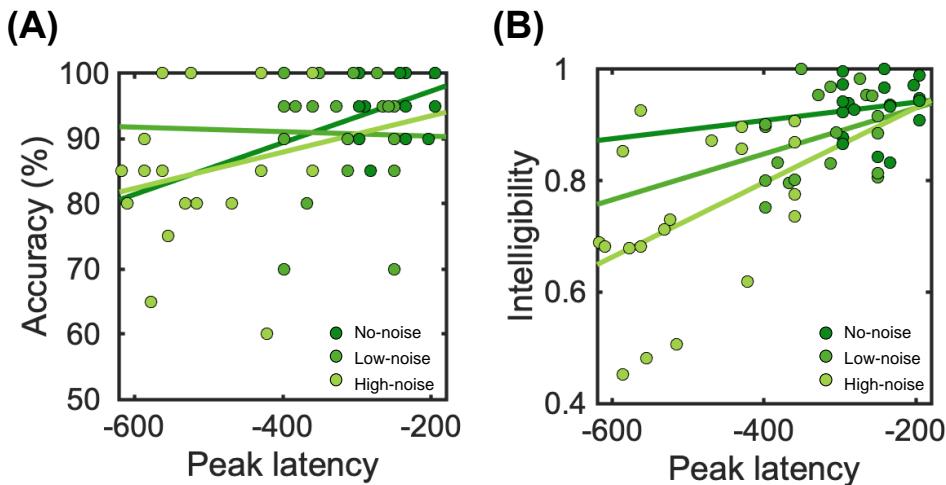
459 As the peak latencies of both the post-onset positive peak of acoustic-level TRFs and the pre-onset
460 negative peak of semantic-level TRFs showed significant differences among SNR levels, we then created
461 linear mixed effect models to explore whether these peak latencies were sensitive predictors of the
462 behavioral performance with the following general formula:

463
$$\text{Behavioral performance} \sim \text{SNR} (+\text{Peak latency}) (+\text{SNR:Peak latency}) + \text{Random} \quad (6)$$

464 where “Behavioral performance” refers to the comprehension performance, clarity, or intelligibility ratings.
465 “SNR” takes the values of speech intensity percentage, i.e., 100%, 60%, and 40%. “Peak latency” refers to
466 the peak latencies of either the acoustic-level post-onset TRFs or the semantic-level pre-onset TRFs,
467 depending on the specific model being investigated. “SNR:Peak latency” refers to the interaction between
468 them. A random intercept per participant was included in the model. “Peak latency” and “SNR:Peak latency”
469 were added between brackets to the general formula because these factors were included only if they led to
470 a lower Akaike Information Criterion (AIC) which indicated a better fitting (Verschueren et al., 2022).
471 Results showed that overall, the behavioral performance was correlated with SNR levels, which echoed the
472 behavioral results in the section 3.1. More importantly, the earlier peak latencies of semantic-level pre-
473 onset TRF response were correlated with the decreasing comprehension performance (LME, $\beta = 2.61 \times 10^{-4}$,
474 $t(52.64) = 1.853, p = .069$) and the decreasing perceived intelligibility (LME, $\beta = 9.26 \times 10^{-4}$, $t(40.48) = 3.497, p = .001$). And the correlation with intelligibility was more prominent as noise increased (LME
475 interaction, $\beta = -7.69 \times 10^{-4}$, $t(45.80) = -1.987, p = .053$), as illustrated in Table 1 and Figure 7. The
476 relationships between the reconstruction accuracies and the behavioral performance were also examined
477 through Spearman’s correlation and summarized in Supplementary Table S5 and Table S6.

479 **Table 1.** LME models of the behavioral performance as a function of SNR and peak latency. Each row
 480 indicates a different model. The SNR was given as a percentage (100%, 60%, 40%).

		SNR			Peak latency			SNR:Peak latency		
		β	t	p	β	t	p	β	t	p
Comprehension performance	TRF _{acoustic}	1.46×10 ⁻²	3.835	.001				No lower AIC		No lower AIC
	TRF _{semantic}	6.23×10 ⁻²	1.056	.297	2.61×10 ⁻⁴	1.853	.069			No lower AIC
Clarity	TRF _{acoustic}	8.87×10 ⁻¹	16.481	< .001				No lower AIC		No lower AIC
	TRF _{semantic}	8.87×10 ⁻¹	16.481	< .001				No lower AIC		No lower AIC
Intelligibility	TRF _{acoustic}	1.74×10 ⁻¹	6.050	< .001				No lower AIC		No lower AIC
	TRF _{semantic}	-1.34×10 ⁻¹	-1.147	.257	9.26×10 ⁻⁴	3.497	.001	-7.69×10 ⁻⁴	-1.987	.053



481 **Figure 7.** Correlation between the peak latencies of the semantic-level pre-onset TRFs and (A) the
 482 comprehension performance and (B) the perceived intelligibility. Colored dots indicate data points from
 483 each participant at different SNR levels.

484 **4 Discussion**

485 The current study investigated the neural tracking of hierarchical features of naturalistic speech in
486 noisy situations using a TRF-based technique. Significant post-onset acoustic-level TRF responses were
487 found within the central electrodes at around 400 ms, and the peak latencies were delayed as noise increased.
488 Significant pre-onset semantic-level TRF responses were found within the frontal electrodes at around -
489 600~200 ms. The peak latencies showed a gradually advanced trend as noise increased, and increased
490 advancement was correlated with decreasing comprehension performance and intelligibility. These findings
491 indicated that noise differently modulates acoustic and semantic processing and suggested that robust and
492 adaptive semantic pre-activation could play a vital role in reliable speech comprehension in noisy
493 environments.

494 The delayed peak latency in the acoustic-level TRF responses as noise increased was in line with
495 several previous studies, suggesting an impaired efficiency in challenging conditions with background
496 noise (Gillis, Decruy, et al., 2022; Mirkovic et al., 2019; Muncke et al., 2022; Yasmin et al., 2023; Zou et
497 al., 2019). As the frontally and centrally distributed channels (corresponding to the primary auditory cortex,
498 superior temporal gyrus, premotor cortex, etc.) have been frequently reported to be related to the processing
499 of speech acoustics (e.g., Bidelman & Howell, 2016; Broderick et al., 2019; Hickok & Poeppel, 2007; Zou
500 et al., 2019), the present TRF results would imply similar recruitment of these brain regions for acoustic-
501 level processing for naturalistic speech under various SNR levels. However, the post-onset 400-ms latency
502 was later compared to the commonly reported latency of < 300 ms in previous studies (Gillis, Decruy, et
503 al., 2022; Yasmin et al., 2023). This discrepancy could be due to the causal filter used for EEG signal
504 preprocessing in the present study, possibly resulting in a delayed TRF compared to previous studies using
505 noncausal filters, similar as reported by Etard and Reichenbach (2019). Alternatively, it could be possible
506 that the latency modulation started earlier but only reached significance later for the present dataset, as the
507 observed TRF responses exhibited an oscillatory pattern starting much earlier than 400 ms (Figure 5C).
508 While an impaired processing efficiency has been associated with both amplitude and latency modulation
509 by noise in previous studies (Muncke et al., 2022; Zion Golumbic et al., 2013; Zou et al., 2019), the present
510 study together with a series of other studies reporting latency-only results would suggest latency as a more

511 sensitive candidate for noisy speech processing (Ding & Simon, 2013; Kaplan-Neeman et al., 2006;
512 Whiting et al., 1998).

513 At the semantic level, our findings on the pre-onset response to word entropy were consistent with
514 recent studies, in which the neural responses to entropy have been reported to involve neural activities
515 within the left hemisphere at up to 800 ms before onset (Goldstein et al., 2022; Weissbart et al., 2020;
516 Willems et al., 2016). This pre-onset prediction mechanism for the upcoming stimuli was regarded as a
517 fundamental computational principle in the human language processing (Goldstein et al., 2022). Our results
518 echo these findings and highlight the potential of entropy as a promising index for exploring forward-
519 looking prediction mechanisms.

520 More importantly, our results extend the present understanding of the predictive mechanism with the
521 manipulation of the SNR levels and suggest a distinct mechanism for speech-in-noise comprehension at the
522 semantic level. The significant pre-onset response to word entropy appeared at all SNR levels, which would
523 imply that such a forward-looking prediction was robust against noise. In addition, we found that the peak
524 latencies of the pre-onset responses became earlier with increasing noise, and that increased forward shift
525 trend at each SNR level was correlated with poorer perceived intelligibility as well as decreasing
526 comprehension performance. One possible explanation for this phenomenon is that our brain could adjust
527 the timing of predictive processing in response to adverse environments. As semantic prediction can
528 facilitate speech comprehension (Mattys et al., 2012; Miller et al., 1951; Obleser & Kotz, 2010; Pickering
529 & Gambi, 2018; Zekveld et al., 2011), the brain relies on it more heavily as noise increases to counteract
530 distorted auditory input. Nevertheless, noise can increase the processing load and decrease the processing
531 efficiency (Gillis, Decruy, et al., 2022; Kaplan-Neeman et al., 2006; Kong et al., 2014; Mirkovic et al.,
532 2019). To compensate for the interference, the neural system initiates the pre-onset response earlier and
533 extends it for a longer duration, giving our brain more time for the preparation of the upcoming speech
534 information. The more degraded the speech, the greater the need for this kind of “early-bird” compensation.
535 Another possible explanation is that in noisy environments our brain relies more on longer-range prediction
536 based on higher-level context information to enhance speech comprehension. According to a recent study
537 based on GPT-2 and functional Magnetic Resonance Imaging (fMRI) (Caucheteux et al., 2023), the
538 forward-looking prediction involved hierarchical representations and multiple time scales, with the

539 maximum of forecast distance reaching 8 words (corresponding to approximately 3.15s). Future studies
540 could employ local and context-unified entropy (e.g., Brodbeck et al., 2022) and longer time windows to
541 further elucidate the noise effect on the forward-looking prediction. Overall, our findings suggest that the
542 brain has a robust and adaptive prediction mechanism for reliable speech comprehension in noisy
543 environments. As such a pre-onset signature was not observed at the acoustic level, our results suggest the
544 predictive mechanism might be mainly focused at the semantic level (Goldstein et al., 2022; Grisoni et al.,
545 2021), where the speech information is expected to be more abstract and more robust against noise (Yasmin
546 et al., 2023).

547 Interestingly, the spatial patterns of TRF responses to word entropy showed left-lateralization at the
548 no-and low-noise levels and recruited bilateral hemispheres at the high-noise level. The left-lateralization
549 has been reported in studies on speech-in-noise comprehension (Z. Li et al., 2021) and was found to be
550 sensitive to linguistic content (Peelle et al., 2013), word entropy (Willems et al., 2016), and semantic
551 expectancy (Golestani et al., 2013; Obleser & Kotz, 2010). Our results would support the left-lateralized
552 brain regions for predictive speech processing at the semantic level. Meanwhile, research has reported that
553 regions within the right hemisphere, such as the right inferior frontal gyrus, are sensitive to semantic
554 features such as entropy (Willems et al., 2016) and that the involvement of the right hemisphere increased
555 under degraded conditions (Bidelman & Howell, 2016), which was hypothesized as the possible
556 recruitment of additional regions for compensation (Shtyrov et al., 1998, 1999). Accordingly, our results
557 suggest that the involvement of bilateral hemispheric in adverse environments might reflect a semantic-
558 related compensation mechanism.

559 Our results suggested the specificity of the frequency band for processing different levels of speech
560 information. Specifically, acoustic-level TRF response was primarily associated with the theta band
561 whereas semantic-level TRF neural response was dominated by the delta band (Figure 4). This could be
562 explained as that theta- and delta-band neural tracking have different functional roles: the former is related
563 to acoustic processing while the latter is related to semantic/syntactic processing (Dai et al., 2022; Ding et
564 al., 2014; Etard & Reichenbach, 2019; Kösem & van Wassenhove, 2017; J. Li et al., 2023). Alternatively,
565 this distinction could be related to the intrinsic temporal properties of the speech features (Lalor, 2018),
566 that is, a faster acoustic-level fluctuation at the theta rhythm and a slower semantic-level fluctuation at the

567 word rate similar to the delta rhythm. Further studies could employ careful experimental manipulation to
568 clarify whether this frequency-specific neural tracking is the result of intrinsic neural oscillations or
569 stimulus-evoked responses (see a review, Obleser & Kayser, 2019). For instance, researchers could
570 manipulate the speech rate (Oganian et al., 2023) and examine whether the frequency characteristics of
571 neural tracking at the semantic level change in response to the varying word rates.

572 The present study has some limitations to be noted. First, there were several significant TRF responses
573 to word entropy not included in the above analyses and discussions, which were primarily observed at the
574 no-noise level. As the focus of the present study was speech-in-noise comprehension, these responses were
575 not further discussed. Nevertheless, they also reflected speech information processing that would deserve
576 investigations in future studies. Second, while the present study only adopted word entropy and word
577 surprisal as two semantic-level features (Gillis et al., 2021; Goldstein et al., 2022; Heilbron et al., 2022;
578 Weissbart et al., 2020; Willems et al., 2016), the rapid development in NLP methods especially the large
579 language models (LLMs) present us with a broader range of options such as semantic embedding (Heilbron
580 et al., 2022). Future studies could employ additional indexes to fully demonstrate the adaptation mechanism
581 of speech-in-noise comprehension. Furthermore, beyond feature extraction, the LLMs also could serve as
582 brain-aligned agents which could be compared with humans and help unveil shared (or unique) mechanisms
583 in the human brain (Caucheteux et al., 2023; Goldstein et al., 2022; Mahowald et al., 2023; Schrimpf et al.,
584 2020). In sum, future studies could employ the promising NLP-based approach to further extend our
585 understanding of language processing. Third, despite the advantage of the high temporal resolution of EEG
586 in exploring temporal dynamics, the relatively poor spatial resolution limits the ability to investigate brain
587 regions involved in predictive mechanisms. A more fine-grained analysis of the spatiotemporal dynamics
588 of semantic prediction would require techniques such as fMRI, ECoG, or multimodal approaches.

589 In summary, the current study investigated how noise affected acoustic and semantic processing during
590 naturalistic speech comprehension. With increasing noise, acoustic processing became increasingly delayed
591 whereas semantic processing became increasingly advanced. Our results suggest that, while the efficiency
592 of brain processing of speech information is indeed impaired by noise, the brain could compensate for the
593 associated effects through active prediction at the semantic level. Overall, the present findings are expected

594 to contribute to the growing research on the neural mechanisms of naturalistic speech comprehension in
595 noisy environments.

596 **Funding**

597 This work was supported by the National Natural Science Foundation of China (NSFC) and the
598 German Research Foundation (DFG) in project Crossmodal Learning (NSFC 62061136001/DFG TRR-
599 169/C1, B1), and the National Natural Science Foundation of China (61977041).

600

601 **Author contributions**

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603 Writing – original draft, Writing – review & editing. **Jiawei Li**: Conceptualization, Methodology,
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608 Writing – review & editing, Funding acquisition, Supervision.

609

610 **Declaration of competing interest**

611 All the co-authors declare that they have no conflict of interest.

612

613 **Data availability statement**

614 Data in the current study are openly available in the Open Science Framework at <https://osf.io/de9p6/>,
615 reference number DOI 10.17605/OSF.IO/DE9P6.

616

617 **Acknowledgment**

618 The authors would like to thank Prof. Xiaoqin Wang for providing the shielded room for the
619 experiment. The authors would like to thank Prof. Zhiyuan Liu and members of his lab for computing the
620 NLP models. The authors would like to thank Lingyi Tao and Xuelin Wang for their help with data
621 collection.

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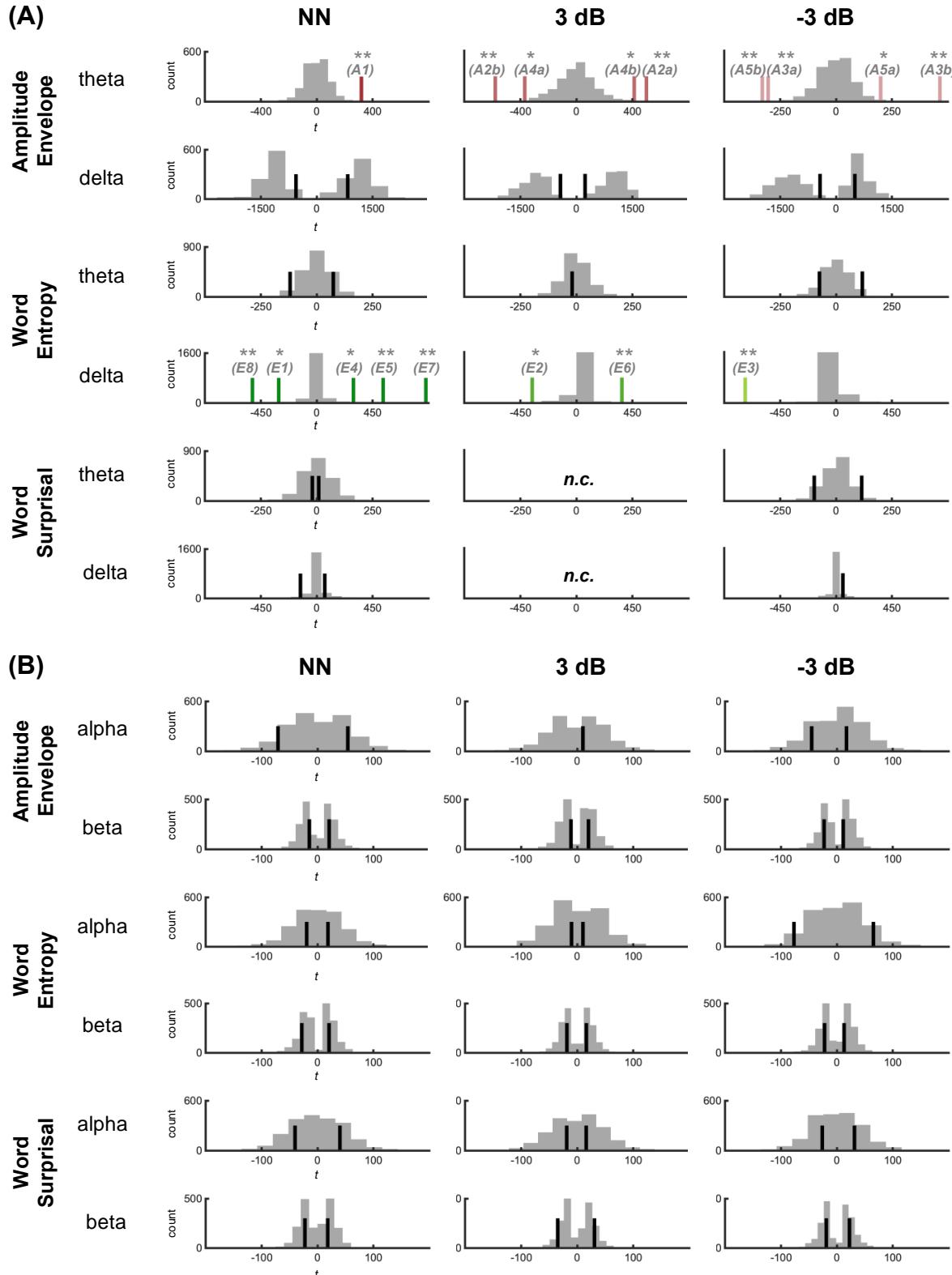
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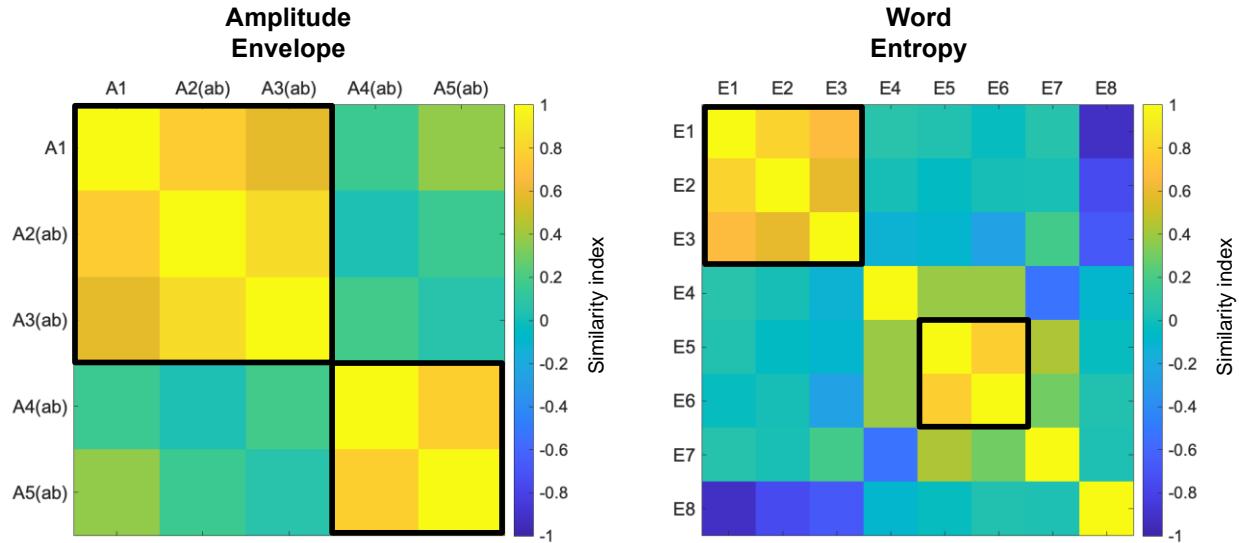
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897 **5 Supplementary materials**



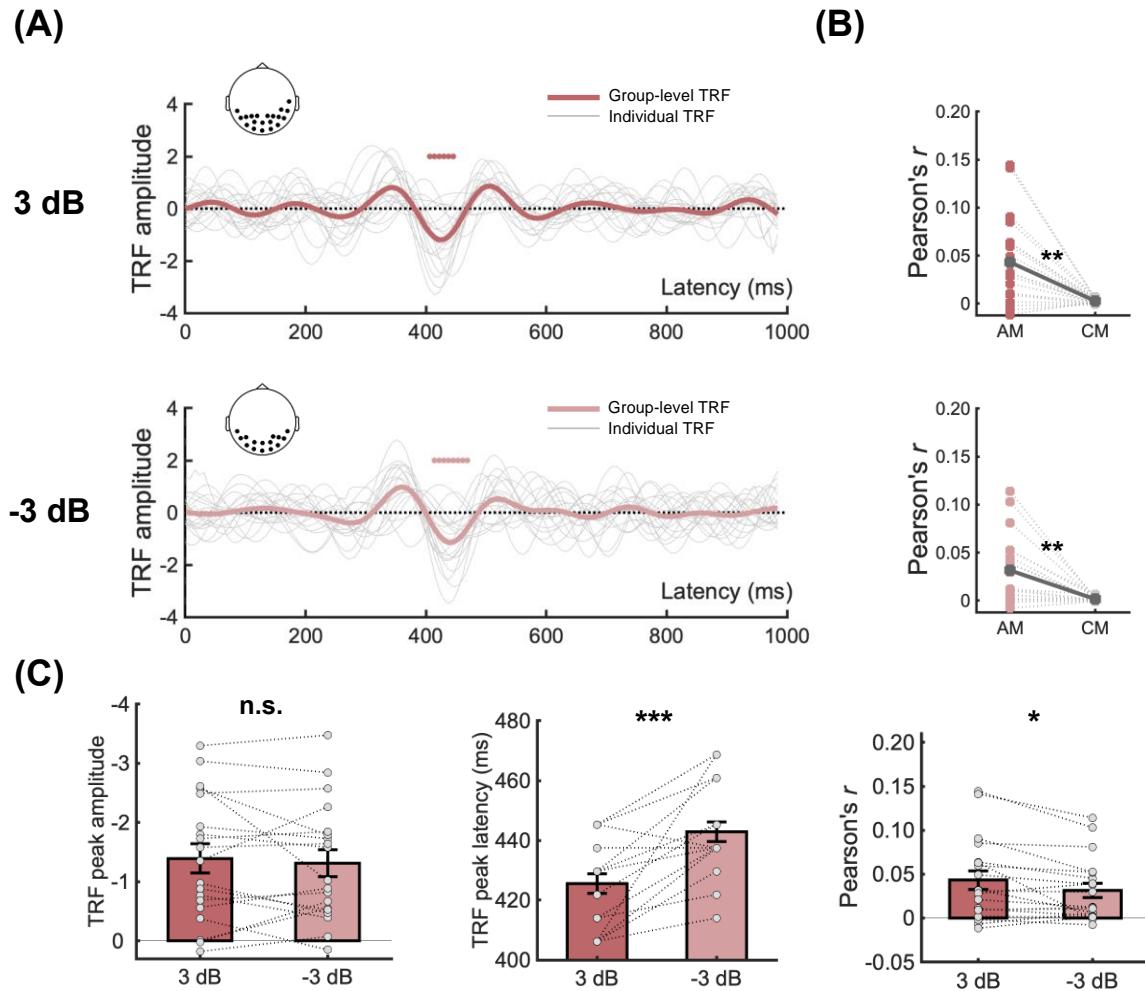
898 **Figure S1.** Clusters in TRF responses for different speech features at different SNR levels in the (A)
899 delta/theta and (B) alpha/beta bands. The grey histograms show the distribution of the cluster-level test
900 statistics from 1,000 permutations. The colored lines indicate significant clusters, and the black lines
901 indicate nonsignificant clusters. n.c.: no cluster is formed. *: $p < .05$, **: $p < .01$, ***: $p < .001$.

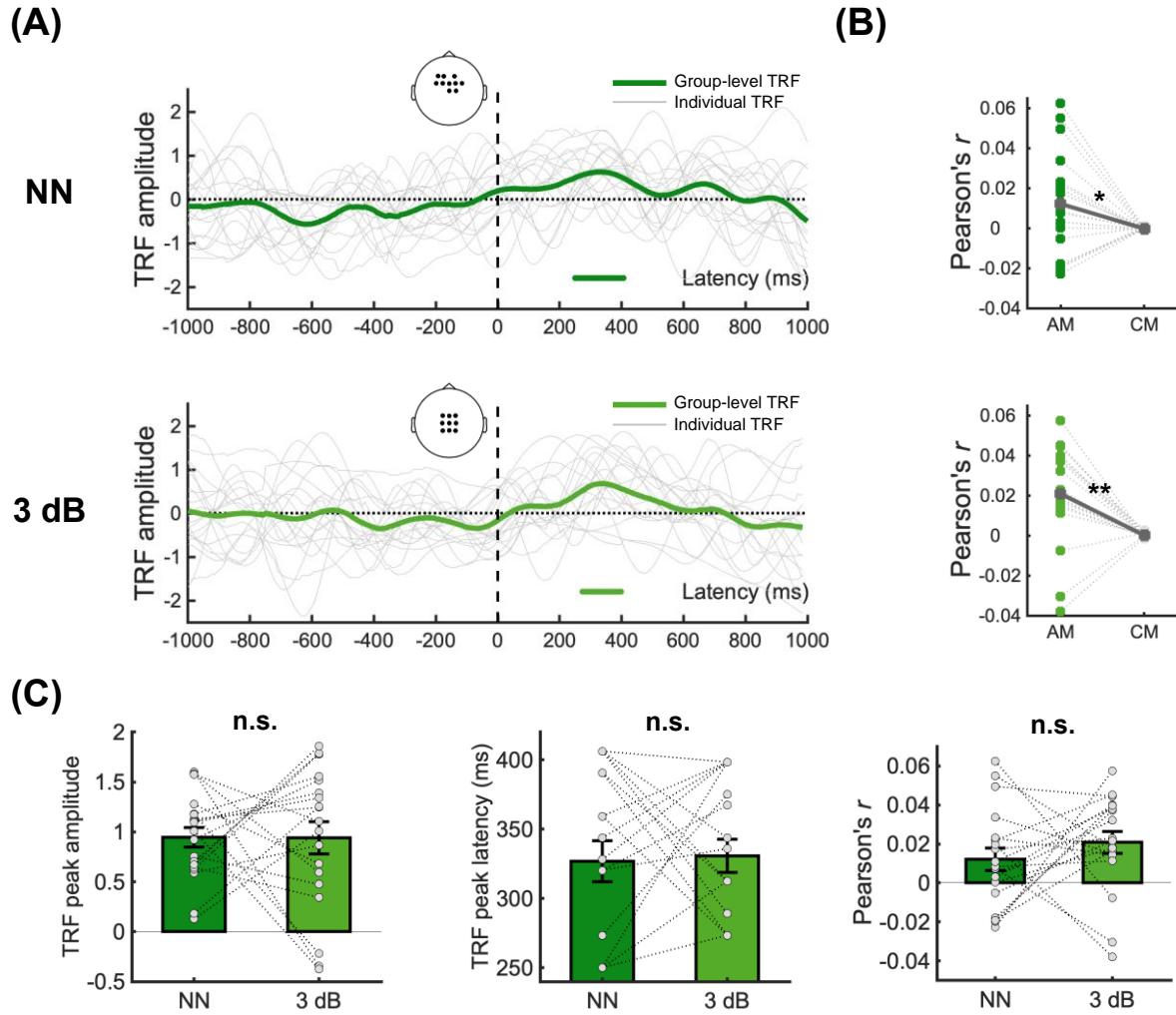


902 **Figure S2.** Similarity index of acoustic-level and semantic-level clusters. The similarity index was
903 calculated as the following formula:

904
$$\text{Similarity index} = \frac{1}{2} (S_{TRFs} + S_{topographies})$$

905 Where S_{TRFs} refers to the temporal similarity, which derived from the Pearson's correlation between the
906 averaged TRFs within the ROIs, and $S_{topographies}$ refers to the spatial similarity, which derived from the
907 cosine similarity between topographies of the peak of clusters. Black rectangles indicate the visually
908 identified clusters with similar spatiotemporal patterns.





921 **Figure S4.** Cluster E5 and E6 of the semantic-level TRF responses to word entropy in the delta band. (A)
922 The bold curves in different shades of green are the mean of TRFs averaged among ROIs across participants
923 at the no- and low-noise levels. The grey curves are TRFs averaged among the ROIs of each participant.
924 The colored horizontal line at the bottom of each plot indicates TOIs over which TRFs differed significantly
925 from the control models. Dots in the corresponding topographies depicted the ROIs. (B) Reconstruction
926 accuracy calculated from the ROIs/TOIs (A), AM means actual models. CM means control models. (C)
927 Noise effect on the peak amplitude, peak latency, and reconstruction accuracy. No significant differences
928 were found in the peak amplitudes (paired-samples t -test, $t(18) = 0.02$, $p = .981$), peak latencies (paired-
929 samples t -test, $t(18) = -0.21$, $p = .834$), and reconstruction accuracies (paired-samples t -test, $t(18) = 0.95$, p
930 = .357). Grey dots indicate data points from each participant. Error bar denotes the standard error. n.s.: not
931 significant, *: $p < .05$, **: $p < .01$, ***: $p < .001$.

932 **Table S1.** Details of the Natural Language Processing model used to extract semantic features.

Type	Parameter
Model type	Long-Short Term Memory (LSTM)
Embedding size	200
Hidden units per layer	200
Number of layers	2
Initial learning rate	3
Gradient clipping	0.25
Sequence length	35
Drop out	0.2
Epoch	50
Batch size	3

933 **Table S2.** Descriptive statistics of semantic features. Std. means standard deviation.

	Number of words	Word entropy			Word surprisal		
		Mean	Range	Std.	Mean	Range	Std.
Story1	213	6.56	9.31	2.17	6.12	16.98	3.22
Story2	245	6.41	9.77	2.16	5.56	17.02	3.38
Story3	247	6.53	9.77	2.24	5.49	13.37	2.96
Story4	249	6.68	10.28	2.32	6.17	15.25	3.31
Story5	240	6.90	10.27	2.24	6.05	13.68	3.23
Story6	228	6.59	10.00	2.16	6.08	15.80	3.18
Story7	281	6.52	9.96	2.41	5.84	17.05	3.04
Story8	245	6.53	9.47	2.31	5.86	13.34	3.13
Story9	254	6.48	9.88	2.39	6.35	16.62	3.27
Story10	244	6.56	9.84	2.35	5.82	17.03	3.26
Story11	231	6.41	9.60	2.24	6.53	18.55	3.81
Story12	271	6.41	10.01	2.29	6.23	17.65	3.59
Story13	317	6.57	10.22	2.30	5.91	17.74	3.49
Story14	243	6.93	10.10	2.35	6.47	16.95	3.42
Story15	305	6.24	9.45	2.17	5.66	17.37	3.1
Story16	303	6.57	9.70	2.32	6.33	17.19	3.36
Story17	344	6.53	9.46	2.09	5.98	15.66	3.06
Story18	251	6.69	9.71	2.01	6.06	14.75	3.33
Story19	266	6.35	9.58	2.14	5.67	13.77	3.04
Story20	274	6.45	9.69	2.12	5.87	14.36	3.08
Story21	248	6.27	9.54	2.36	5.34	15.96	3.11
Story22	274	6.40	10.15	2.45	5.82	17.36	3.45
Story23	272	6.32	9.75	2.29	5.57	15.25	2.93
Story24	264	6.67	9.88	2.23	6.04	16.43	3.29
Story25	271	6.53	9.53	2.16	6.02	17.32	3.14
Story26	253	6.54	9.62	2.17	6.41	15.85	3.19
Story27	261	6.44	9.41	2.03	6.17	17.82	3.15
Story28	236	6.75	10.02	2.41	6.03	13.41	3.36
Story29	282	6.65	9.89	2.04	6.27	15.88	3.09
Story30	235	6.71	9.78	2.28	6.31	18.16	3.48

934 **Table S3.** Time lags of interests (TOIs) of each cluster in the acoustic-level TRF responses.

Cluster	Time lags of interests (TOIs)
A1	383 ~ 483 ms
A2a	406 ~ 453 ms
A2b	484 ~ 539 ms
A3a	344 ~ 383 ms
A3b	414 ~ 469 ms
A4a	406 ~ 445 ms
A4b	484 ~ 531 ms
A5a	336 ~ 383 ms
A5b	414 ~ 469 ms

935 **Table S4.** Time lags of interests (TOIs) of each cluster in the semantic-level TRF responses.

Cluster	Time lags of interests (TOIs)
E1	-297 ~ -195 ms
E2	-398 ~ -250 ms
E3	-625 ~ -359 ms
E4	-8 ~ 141 ms
E5	250 ~ 406 ms
E6	273 ~ 398 ms
E7	602 ~ 961 ms
E8	563 ~ 727 ms

936 **Table S5.** Spearman's correlation between the reconstruction accuracies of each cluster in acoustic-level
937 TRF responses and the behavioral performance. Significant correlation results are bolded (uncorrected).

	Comprehension	Clarity	Intelligibility
	Performance	Ratings	Ratings
A1	$r = .11, p = .647$	$r = -.11, p = .662$	$r = .10, p = .694$
A2a	$r = .05, p = .831$	$r = -.46, p = .047$	$r = -.04, p = .858$
A2b	$r = .25, p = .309$	$r = -.50, p = .032$	$r = .02, p = .926$
A3a	$r = .09, p = .702$	$r = -.52, p = .025$	$r = .05, p = .854$
A3b	$r = -.002, p = .994$	$r = -.51, p = .028$	$r = -.06, p = .798$
A4a	$r = .08, p = .757$	$r = -.47, p = .043$	$r = -.08, p = .734$
A4b	$r = .18, p = .465$	$r = -.42, p = .071$	$r = .004, p = .986$
A5a	$r = .03, p = .918$	$r = -.46, p = .048$	$r = .002, p = .997$
A5b	$r = .22, p = .361$	$r = -.52, p = .026$	$r = .15, p = .541$

938 **Table S6.** Spearman's correlation between the reconstruction accuracies of each cluster in semantic-level
939 TRF responses and the behavioral performance. Significant correlation results are bolded (uncorrected).

	Comprehension	Clarity	Intelligibility
	Performance	Ratings	Ratings
E1	$r = -.09, p = .714$	$r = -.21, p = .394$	$r = .14, p = .560$
E2	$r = .11, p = .668$	$r = .23, p = .339$	$r = -.07, p = .762$
E3	$r = -.01, p = .997$	$r = -.30, p = .206$	$r = -.31, p = .203$
E4	$r = -.02, p = .928$	$r = -.24, p = .332$	$r = .09, p = .721$
E5	$r = .51, p = .027$	$r = .15, p = .527$	$r = .37, p = .121$
E6	$r = .13, p = .594$	$r = -.22, p = .365$	$r = -.30, p = .215$
E7	$r = .20, p = .404$	$r = .04, p = .881$	$r = .15, p = .551$
E8	$r = .20, p = .411$	$r = -.28, p = .237$	$r = -.14, p = .575$