Exploring the impact of social relevance on the cortical tracking of speech: viability and temporal response characterisation

Abbreviated title: Social relevance enhances cortical speech tracking

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Abstract

Human speech is inherently social. Yet our understanding of the neural substrates underlying continuous speech perception relies largely on neural responses to monologues, leaving substantial uncertainty about how social interactions shape the neural encoding of speech. Here, we bridge this gap by studying how EEG responses to speech change when the input includes a social element. In Experiment 1, we compared the neural encoding of synthesised undirected monologues, directed monologues, and dialogues. In Experiment 2, we extended this by using podcasts, addressing the additional challenges of real speech dialogue, such as dysfluency. Using temporal response function analyses, we show that the presence of a social component strengthens the cortical tracking of the speech envelope, despite identical acoustic properties. Neural responses to synthesised speech showed a strong correlation with those for real speech podcasts, with a stronger alignment emerging for more socially-relevant speech material. In addition, we demonstrate that robust neural indices of sound and lexical-level processing can be derived using real podcast recordings despite the presence of dysfluencies. Finally, we present a simulation to put to the test the robustness of temporal response function analyses under increasing levels of dysfluency. Together, these findings highlighting the impact of social elements in shaping auditory neural processing, providing a framework for future investigation and analysis of social speech listening and speech interaction.

Significance Statement

Human speech is rarely produced or processed in a social vacuum. Yet, our understanding of continuous speech neurophysiology mostly comes from experiments involving speech monologues. This study reveals how social context modulates the neural encoding of speech. We directly contrast neural signals recorded when participants listened to monologues and dialogues, using controlled material from speech synthesis and real podcast recordings. We found that the social element amplifies the neural encoding of speech features, reflecting greater engagement. We also show strong correlation between synthetic and real podcast neural responses, scaling with social relevance. Finally, we demonstrate that lexical processing can be measured robustly even amid natural dysfluencies. These insights advance our understanding of speech neurophysiology, informing future research on social speech.

Introduction

To understand speech, our brains must rapidly process sounds into linguistic meaning (Friederici, 2011; Poeppel et al., 2012). The neural basis of this phenomenon has been studied extensively using controlled paradigms involving targeted manipulations (Carreiras et al., 2005; Kutas & Federmeier, 2011; Osterhout & Holcomb, 1993), and more recently with listening tasks involving more realistic continuous streams of speech (Di Liberto & Ip, 2025; Di Liberto et al., 2021; Gwilliams et al., 2023). That work utilised a variety of neural recording technologies, including electro- and magneto-encephalography (EEG and MEG) (Défossez et al., 2023; Destoky et al., 2019), functional magnetic resonance imaging and functional near-infrared spectroscopy (Butler et al., 2020; Chen et al., 2020; Formisano et al., 2008), and intracranial electrical recordings (e.g., electrocorticography and stereotactic EEG) (Bernabei et al., 2021; Hamilton et al., 2021; Mesgarani et al., 2014). Overall, this research has led to a new understanding of the sound-to-meaning transformation, providing detailed insights into the spatial and temporal dynamics that unfold throughout that process (Kocagoncu et al., 2017; Uusvuori et al., 2008). However, the necessary experimental constraints have led to listening tasks centred around audiobook-style, pre-recorded speech material, while ignoring one of the key functions of speech: the social element.

Recent research has explored speech interaction, for example by measuring brain-to-brain synchrony using simultaneous neural recordings from interacting individuals (Kinreich et al., 2017; Schwartz et al., 2024). However, those interactive experiments are far removed from previous speech listening research, as other factors at play (e.g., social anxiety, familiarity, turn-taking) make it difficult to relate findings across the two domains. To achieve this and advance the study of social speech neurophysiology, we require a mechanistic understanding of how specific social factors impact the neural processing of speech, which requires careful experimental control. The present study takes a step in that direction, by extending the conventional audiobook-style listening paradigm to stimuli with progressively higher social relevance. In doing so, we investigate how challenges of real speech (e.g., dysfluencies) impact our ability to measure the neural processing of speech and language (Di Liberto & Ip, 2025; Mousavi et al., 2024).

We present results from two EEG experiments. The first experiment compares the
neural encoding of speech while participants listened to pre-recorded monologues and
dialogues (**Experiment 1**). *Dialogue* typically involves turn-taking, mutual feedback,
and shared communicative goals (Garrod & Pickering, 2004), while monologues lack
immediate (auditory) reciprocity. Nevertheless, monologues can also include a social
dimension by being *directed* to a listener, as opposed to *undirected monologues*, when
they are not aimed at any particular listener. Using the temporal response function
analysis (TRF), this study tests if social relevance impacts the neural processing of
speech during listening. This leads to our first hypothesis:

Hypothesis 1: Social relevance enhances the cortical encoding of speech,
leading to stronger encoding for directed than undirected monologues.

Synthesised speech was used to control for acoustic and lexical differences across
conditions, ensuring that differences in the EEG would be due to social relevance as
opposed to other factors such as prosody, word rate, or volume level (see **Methods**).
To generalise the results from Experiment 1 to the case of real human speech, we
also conducted a second experiment using real dialogue recordings from popular
podcasts (**Experiment 2**). The material included speech from different registers to
capture a broader range of speech types, increasing generalisability. Furthermore, real
spontaneous includes dysfluency(Shriberg, 2001), which can challenge natural
speech comprehension, for example by hampering syntactic parsing (Ferreira &
Bailey, 2004), contributing to processing delays that impact word predictability (Corley
et al., 2007). Here, we investigate the possibility of studying speech material including
dysfluencies, testing if robust EEG measurements of speech processing can be
derived at varying levels of dysfluency, with the following hypotheses:
118
119   **Hypothesis 2**: The social element of speech leads to a consistent neural
120   signature, which generalises across synthesised and real speech dialogue.
121   **Hypothesis 3**: Neural measurements of speech encoding are robust to
122   dysfluency in the speech material.
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Materials and Methods

EEG Experiment 1

Data acquisition and experimental setup. Twenty healthy individuals (10 male, 10

female; between 18 and 30 years old, median = 22) participated in the experiment. All

participants were native English speakers and reported no history of hearing

impairment or neurologic disorder, provided written informed consent, and were paid

for participating. The study was undertaken in accordance with the Declaration of

Helsinki and was approved by the ethics committee of the School of Computer Science

and Statistics at Trinity College Dublin. The experiment was carried out in a single

session for each participant. EEG data were recorded from 64 electrode positions,

digitised at 512 Hz using a BioSemi ActiveTwo system. Audio stimuli were presented

at a sampling rate of 48 kHz using Sennheiser HD 650 headphones and the

Psychtoolbox-3 MATLAB library (Brainard, 1997). Testing was conducted at Trinity

College Dublin in a dark room. Participants were instructed to maintain visual fixation

on a crosshair centred on the screen and to minimise motor activities during the

speech listening task.

Stimuli. Monologue scripts were generated with the large language model (LLM) GPT-

4 (OpenAI, 2024), using prompts such as "Could you create a monologue, with a

duration of approximately 2 minutes?". From 30 generated monologues, scripts with

highly similar content and those containing strongly emotional themes (e.g., scenarios

involving extreme sadness or anger) were excluded to avoid potential confounds in

the experiment. After this filtering process, 16 monologue scripts with sufficiently

varied topics were retained for use as stimuli. The selected monologues were further

processed with GPT-4, generating eight directed and eight undirected monologues.

That manipulation was achieved by using prompts such as: _"Turn this into a directed_

monologue that addresses the listener" and "_Turn this into an undirected monologue,_

where the speaker talks to themselves."

The 16 monologues (8 directed, 8 undirected) were then converted into dialogues

using the same approach, with the prompt: _"Convert the following monologue into a_

dialogue between two speakers while preserving the original content. Keep the

duration approximately the same and distribute the information naturally across the

speakers." The aim of this transformation was to create dialogue versions that
preserved the semantic content of the original monologues while introducing an
interlocutor. This approach allowed us to maintain comparable content across
monologue and dialogue conditions, thereby minimising potential confounds related
to topic or narrative structure.

Scripts were synthesised using the realistic AI-based voice generation software
PlayHT version 2.0[1]. The generated speech sounds were manually inspected
sentence-by-sentence to minimise generation artifacts. Speech segments with clear
artifacts or sounding unnatural were replaced by re-running the generation for the
affected segment, ensuring a consistent tone and emphasis. Note that this was
possible as each generation produces a different result. Altogether, this procedure
generated 32 sound files (16 monologues, 16 dialogues) synthesised using different
voices (16 voices: 8 male, 8 female). The final version of auditory stimuli has been
deposited in the OSF repository[2].

To control for potential acoustic and speech-structure confounds, several acoustic
properties of each stimulus, including duration, RMS amplitude, mean fundamental
frequency (F0), F0 variability (standard deviation), and word rate (number of words
per second) were quantified. Participants (N=15) participated in an online survey,
where they were asked to categorise whether speech recordings were undirected or
directed. Participants were presented with 16 speech recordings, each ~2-min of
duration. Acoustic measures were compared across conditions (undirected-directed
monologues, dialogues) using Kruskal-Wallis tests. No significant differences were
observed for any of the acoustic properties across the three conditions (all Kruskal-
Wallis tests, $p > 0.05$; see **Table S3**).

In addition, monologue stimuli (combination of directed and undirected conditions)
were compared with dialogue stimuli, showing no statistically significant differences
for any of the acoustic properties (all Wilcoxon rank-sum tests, $p > 0.05$; see **Table**
S3).

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[1] *https://app.play.ht/*
[2] *https://osf.io/hx3nz*

Experimental procedure. All thirty-two stimuli were presented in a single experimental

session in a randomised order, which was different across participants. Participants

were free to take breaks of arbitrary length in-between stimuli. To maintain

engagement throughout the experiment, participants were periodically (25% of the

trials) asked to rate how interesting they found the speech segments on a 5-point scale

(1: not at all interesting; 3: neutral; 5: very interesting).

EEG Experiment 2

Data acquisition and experimental setup. Twenty healthy individuals (10 male, 10

female; between 20 and 39 years old, median = 26), separate from Experiment 1,

participated in the second EEG experiment. All participants were native English

speakers and reported no history of hearing impairment or neurologic disorder,

provided written informed consent, and were paid for participating. The study was

undertaken in accordance with the Declaration of Helsinki and was approved by the

ethics committee of the School of Psychology at Trinity College Dublin.

The experiment was carried out in a single session for each participant. EEG data

were recorded from 64 electrode positions, digitised at 512 Hz using a BioSemi

ActiveTwo system. Audio stimuli were presented at a sampling rate of 44,100 Hz using

Sennheiser HD 280 Pro headphones and the PsychoPy Python library version

2022.2.5 (Peirce et al., 2019). Testing was conducted at Trinity College Dublin in a

dark room. Participants were instructed to maintain visual fixation on a crosshair

centred on the screen and minimise motor activities during the speech listening task.

Stimuli. Podcast recordings were selected from publicly available sources: 1)

American podcast shows called *Brains On!* and *Forever Ago;* 2) *WIRED 5 Levels of*

Difficulty series on YouTube; and 3) interviews with the same two hosts *Forever Ago,*

available online[3]. Dialogues contained general interest topics, such as questions about

science and math (e.g., why do we sweat, why memories as a young child are

forgotten). Altogether, forty-two podcast recordings involving 1-to-1 dialogues were

selected for this experiment (the same stimuli for all participants), totalling one hour in

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[3] https://laist.com/podcasts/servant-of-pod/kids-podcasts-a-true-alternative-to-screen-time;
https://creators.spotify.com/pod/profile/martine-severin/episodes/31--How-to-Become-an-Epic-Improv-Comic-with-Joy-Dolo-e1js9c2

duration. Sixteen of those recordings involved adult-adult (AA) dialogues, while the
remaining twenty-six were adult-child (AC) dialogues. The AA and AC dialogues
contained the same five adult speakers (podcast hosts), speaking to a range of
different children and other adults. The higher number of recordings for the latter
condition ensured that the two conditions had a comparable number of words, as child-
speech and child-directed speech are generally slower than adult-speech (Ferreira,
2019; Snow, 1972). To that end, the selected speech registers captured a range of
speaking styles for testing the robustness of our findings (e.g., the impact of
dysfluencies on the EEG encoding of speech) to variations in acoustic and lexical
properties of a speech input.

Experimental procedure. Participants were presented with the forty-two speech stimuli
(referred hereafter as trials) in a random order, separated by a short break. To
encourage attention, participants were asked to rate how interesting they found the
speech segment on a scale from 1-5 (1: not at all interesting; 3: neutral; 5: very
interesting) after each trial. Additionally, for a random subset of twenty percent of the
trials, the participants were asked a behavioural multiple-choice question to select the
set of keywords that best described the trial they just heard. This was also to keep the
participants engaged and on task throughout the experiment.

EEG data preprocessing
Neural data from both experiments were analysed offline using MATLAB software
(MathWorks), using the same preprocessing and analysis pipeline. EEG signals were
digitally filtered using Butterworth zero-phase filters with order 2, implemented with the
function *filtfilt*. The primary analyses were carried out on EEG data band-pass filtered
between 0.5 and 8 Hz, as that range has been shown to be particularly sensitive to
the cortical tracking of speech (Di Liberto et al., 2015; Doelling et al., 2014; O'Sullivan
et al., 2015; Vanthornhout et al., 2018). Low-frequencies from 0.5 Hz to 1 Hz were
included as they were previously shown to be important for measuring the neural
encoding of lexical predictions with EEG (Broderick et al., 2018; Carta et al., 2024).
Secondary analyses were carried out to pinpoint specific sub-bands driving effects of
interest. In Experiment 1, we considered the delta-band (0.5-4 Hz), theta-band (4-8
Hz), alpha-band power (8-12 Hz), beta-band power (12-30 Hz) and broadband (0.5-
30 Hz). In Experiment 2, we considered just the delta-band (0.5-4 Hz) and theta-band

(4-8 Hz), as they led to the highest EEG prediction correlation in Experiment 1 (**Figure
S1**). The power of alpha and beta bands was calculated as the instantaneous
magnitude of the Hilbert transform in those sub-bands. This selection is based on
evidence that alpha and beta rates reflect predictive processing, attention, and working
memory (Haegens et al., 2017; Samaha et al., 2015). EEG signals were down sampled
to 64 Hz, reducing computational and storage costs. EEG channels with a variance
exceeding three times that of the surrounding ones were replaced by a spherical spline
interpolation (Delorme & Makeig, 2004; Perrin et al., 1989). Signals were then re-
referenced to the average of the two mastoid channels.

Stimulus features
The present study examines the neural encoding of speech and language features
during speech listening. The same features and analyses were used across the two
experiments.

Sound envelope and derivative. The sound envelope was calculated as the
instantaneous magnitude of the Hilbert transform of the sound-pressure waveform.
The speech envelope captures the slow amplitude modulations of the speech signal,
which have been shown to be robustly reflected in the EEG signal (Crosse et al.,
2016a; Ding & Simon, 2012; Lalor & Foxe, 2010). The envelope-related features
assessed in this study comprised the sound envelope and half-way rectified envelope
derivative (calculated from the envelope at the original sampling rate with the MATLAB
diff function, and by replacing negative values with zero). Matching the processed EEG
signal, these acoustic features were then down sampled to 64 Hz.

Word onsets. The forced alignment software DARLA (Reddy & Stanford, 2015) was
used to produce a word-by-word time-alignment between each speech segment and
its transcript. The onset time of all words in the speech stimuli was then coded into
sparse vectors of zeros and ones with the same length and sampling rate as the other
features, where ones represent the onset time of a word, consistent with previous
studies (Broderick et al., 2018; Chalehchaleh et al., 2025; Di Liberto et al., 2021).

Lexical surprise and entropy. The second experiment in the present study investigates
the possibility of measuring neural signatures of lexical surprise and entropy in the

presence of dysfluencies, which are typically part of dialogues (Bosker et al., 2014;
Clark & Tree, 2002; Fox Tree, 2001; Tree, 1995). Next-word lexical surprise and
entropy values were extracted from Mistral-7B-Instruct-v0.1 (Jiang et al., 2023), an
open source state-of-the-art LLM that has been fine-tuned on conversation datasets.
For each trial, lexical surprise was calculated as the negative log contextual probability
of each word (Heilbron et al., 2022; Slaats & Martin, 2025). Additionally, for each trial
lexical entropy was calculated as the weighted sum over the surprise values for all
possible next words, and measures the level of uncertainty the model has for its next-
word prediction (Goldstein et al., 2022; Slaats & Martin, 2025). Lexical surprise and
entropy feature vectors were built by scaling the word onset vector by its
corresponding surprise and entropy value, respectively. Previous work employing
similar models (e.g., GPT-2) found lexical expectation values to reliably explain
EEG/MEG variance when considering speech listening tasks (Dou et al., 2024;
Heilbron et al., 2022). However, that work typically involved scripted speech
monologues without dysfluencies.

Dysfluencies. To investigate the impact of dysfluencies on lexical surprise and entropy
and their neural signatures, we built two lexical surprise and entropy vectors: (1) one
that included dysfluencies and (2) one that excluded dysfluencies in the speech
material (**Figure 4A**). Such dysfluencies were manually transcribed, annotated, and
removed (see the dysfluency types considered in **Table S1**). To balance the amount
of data in both versions of the surprise and entropy vectors, the annotated dysfluencies
were removed from both feature vectors. Therefore, the main difference in the two
versions of the surprise and entropy vectors lies primarily in if/how the presence of
dysfluencies impacts the predictions of upcoming words. In practice, this led to two
surprise and entropy feature vectors with the same word onsets, but different surprise
and entropy values.

Temporal response function analysis
A system identification approach was employed to determine how specific stimulus
features map onto EEG activity. This approach, known as the Temporal Response
Function (TRF) (Ding & Simon, 2012; Edmund C. Lalor, 2009), utilises regularised
ridge linear regression (Crosse et al., 2016) to estimate a filter, or model, that best
captures how the brain converts input stimulus features into the resulting neural

305 responses. One of the two metrics derived from the TRF is the weights of the model
306 over time. The brain's response to a speech stimulus is not instantaneous; rather, it
307 unfolds over time when speech is heard. Therefore, the brain's electrical activity is
308 affected not just at the moment the sound occurs, but also over a subsequent time
309 period. This delayed and extended influence is captured in the time lags of the TRF
310 weights between the stimulus and the EEG response. These time lags can be
311 analysed both spatially (across the scalp) and temporally (in terms of response
312 delays).

313 As is typical (Crosse et al., 2016b; Crosse et al., 2021), to evaluate the model's ability
314 to generalise and avoid overfitting, we used leave-one-out cross-validation across
315 trials. In this process, TRF models were trained on all but one trial and then used to
316 predict the EEG signal for the left-out trial. The TRF model performance is then
317 assessed by computing Pearson's correlation between the predicted EEG signal and
318 the actual EEG signal on the left-out fold at each scalp electrode. This constitutes the
319 second metric derived from the TRF: the prediction correlation, or the EEG variance
320 explained by a specific set of speech feature(s). The EEG variance captured by
321 speech features is interpreted as how well the model was able to map the encoding of
322 speech features in the brain signals. Fitting the model with one speech feature results
323 in a *univariate model,* while fitting the model while more than one speech feature
324 results in a *multivariate* model.

325 Using these objective metrics, univariate forward TRF models were used to assess
326 the corresponding neural response to each speech feature of interest. Three univariate
327 TRF models were fit in Experiments 1 and 2 for the following speech features: the
328 speech envelope, envelope derivative, and the word onset. In Experiment 1, the
329 strength of the neural response to the speech in each of the three conditions was
330 assessed through the TRF models' prediction correlations for any effect of social
331 relevance. In Experiment 2, the temporal dynamics of the EEG response to podcast
332 dialogues were examined from the TRF weights, expecting latencies consistent with
333 previous results on audiobook listening tasks. Despite variability in the acoustics of the
334 spontaneous dialogue stimuli across speech registers (Ferreira & Ferreira, 2019;
335 Snow, 1972) and, as a result, the neural responses, the TRF measured is designed to
336 remain robust and stable against such specific differences as it estimates the

consistent impulse response of the brain. This serves as an opportunity to build speech
dialogue TRFs that are stable across various types of speech and speech registers in
real dialogues. Notably, we expected to measure a negative deflection of the TRF at
latencies around 400 ms (i.e., the TRF-N400 response) for this podcast listening task
that, compatibly with previous monologue listening work (Broderick et al., 2018;
Broderick et al., 2021; Heilbron et al., 2022), would index word-level processing.

Multivariate TRF (mTRF) analyses were also conducted in Experiment 2 to assess if
the cortical encoding of lexical surprise and entropy was more aligned with a next-
word prediction model including or disregarding dysfluencies. This enables us to
evaluate how much, if at all, dysfluencies impact lexical predictions during speech,
informing us on the most appropriate approach for measuring lexical expectations in
the human brain in contexts involving real speech (i.e., with dysfluencies) (Heilbron et
al., 2022; Mousavi et al., 2024). Sound envelope and word segmentation information
features (e.g., speech, speech derivative, word onset) were concatenated to represent
the stimulus, along with one of two versions of surprise: either dysfluent or non-
dysfluent surprise.

Statistical analyses
Parallel statistical procedures were applied to the datasets from the two experiments.
Statistical analyses were performed on EEG prediction-correlation values obtained
from the TRF models. For each participant and condition, prediction correlations were
averaged across all EEG electrodes to obtain subject-level values, and these values
were used in all statistical tests. To determine whether the TRF models captured
meaningful neural tracking of the speech features, prediction correlations were tested
against zero using one-sample *t*-tests.

To compare experimental conditions, repeated-measures statistical tests were used
because all conditions were recorded from the same participants. When three levels
of social relevance (undirected monologue, directed monologue, and dialogue) were
compared, a one-way repeated-measures ANOVA was applied to the subject-level
averages. When the omnibus test was significant, pairwise comparisons between
conditions were performed using Wilcoxon signed-rank tests.

For analyses of TRF weights across time lags, multiple comparisons were controlled
using the false discovery rate (FDR) procedure with $p = 0.05$, according to the
procedure set out by (Benjamini & Yekutieli, 2001) for dependence assumptions.

A cross-experiment comparison was carried out by correlating TRFs for the two
datasets. Considering that the two datasets differ in both stimuli and participants, we
carried out that comparison via a bootstrap resampling procedure on the group-
averaged TRF weights. Pearson's correlation scores were calculated for the TRF
weights corresponding to univariate envelope, envelope derivative, and word onset
features. The general assumption in averaging TRFs (as for ERPs) is that TRFs for
individual participants may vary, but that participants share certain consistent
dynamics, which would emerge by averaging. Here, the assumption was that
averaging TRFs from two distinct samples of participants (from the same population)
would lead to the same TRF (when enough participants are available), with higher
TRF correlation for more similar experimental tasks. Accordingly, we expected higher
inter-experiments TRF correlations when considering more similar tasks (i.e., dialogue
listening). This test was carried out via a bootstrap resampling procedure that derived
1,000 correlations between the average TRFs for each of the conditions in Experiment
1 and for Experiment 2, generated using a random sample with replacement on each
participant cohort. Because Mauchly's test indicated that the assumption of sphericity
was violated ($p < 0.05$), a nonparametric Friedman test was conducted to examine
differences in the correlations across conditions for each TRF feature using JASP
software (Love et al., 2019).

The dysfluency analysis in Experiment 2 included repeated measures ANOVAs. Post-
hoc tests using paired t-tests with the Holm correction were performed to examine the
effect of including/excluding dysfluencies on TRF model predictions, from which we
report the adjusted p-value. Matlab was used for statistical analysis in Experiment 1.
For Experiment 2, statistical analyses were performed using Matlab and JASP
software.

Code and data accessibility
The TRF analysis was conducted using the publicly available and open source mTRF-
toolbox version 2.6 (Crosse et al., 2016a; Crosse et al., 2021), which can be

downloaded from https://github.com/mickcrosse/mTRF-Toolbox or
https://sourceforge.net/projects/ aespa/. Stimulus features and EEG data were stored
according to the continuous-event neural data structure CND (Di Liberto et al., 2024).
Analyses were conducted using scripts shared via the GitHub channel[4] of the
Cognition and Natural Sensory Processing (CNSP) and Natural Sensory Processing
open science initiative[5]. Custom scripts were used for data visualisation and statistical
analyses. Datasets for Experiments 1 and 2 will be publicly shared via the open
science repository OSF (osf.io) at the time of publication and will also be included in
the list of publicly available datasets of the CNSP-initiative.

Results

Experiment 1

The survey indicated that participants could categorise undirected versus directed
speech sounds, achieving an accuracy of 0.94 (average across participants; ± 0.11
SD). Performance was significantly above chance level, as confirmed by one-sample
t-tests (overall: $t(14) = 15.9$, $p = 1.1 \times 10^{-32}$; undirected stimuli: $t(14) = 13.2$, $p = 3.1 \times$
10^{-21}; directed stimuli: $t(14) = 17.4$, $p = 2.4 \times 10^{-33}$; **Fig. S4**). Monologues were rated
as more interesting on average than dialogues ($M_{monologue} = 3.42$, $M_{dialogue} = 2.87$;
paired t-test; $t(18) = -2.26$, $p = 0.036$), while no statistically significant difference in
interest ratings emerged between undirected and directed monologues ($M_{undirected} =$
3.72; $M_{directed} = 2.89$; paired t-test: $t(8) = -2.13$, $p = 0.066$).

Univariate TRFs were fit to model the relationship between stimulus features (speech
envelope, envelope derivative, and word onset) and the broadband EEG signal (0.5-
30 Hz). Separate TRF models were fit for the three stimulus features. First, we report
results for the widely used sound envelope feature as in previous studies (Muncke et
al., 2022; Vanthornhout et al., 2018). EEG prediction correlation values calculated with
leave-one-out cross validation (see **Materials and Methods**) had magnitudes
compatible with envelope TRF analyses in previous research (r~0.2-0.4, after
averaging across all participants and EEG channels). Comparing EEG prediction
correlations for envelope TRFs across conditions (undirected monologue, directed

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[4] *https://github.com/CNSP-Workshop/CNSP-resources/tree/main/CNSP/ exampleScripts*
[5] *https://cnspworkshop.net/*

monologue, and dialogue) revealed a statistically significant effect of social relevance
(one-way repeated measures ANOVA, $F(2, 38) = 5.98$, $p = 0.006$), with more socially-
relevant stimuli producing larger EEG prediction correlation values. Post-hoc analyses
indicated lower prediction correlations in the undirected monologue condition
compared with both the directed monologue (Wilcoxon signed-rank tests, $p = 0.008$)
and dialogue conditions ($p = 0.003$; **Figure 1A**). No statistically significant differences
were observed between directed monologue and dialogue conditions ($p = 0.89$). No
statistically significant effects of social relevance were observed for the other stimulus
features considering broadband EEG (envelope derivative: $F(2, 38) = 2.16$, $p = 0.129$;
word onset: $F(2, 38) = 1.03$, $p = 0.366$).

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Figure 1. Social relevance impacts the cortical tracking of speech. (A) *Univariate*
forward TRFs were fitted for envelope, envelope derivative, and word onset features
to predict the on the broadband EEG in Experiment 1. EEG prediction correlations
(Pearson's r) were first averaged across electrodes within each participant, and group
means (± SEM across participants) are shown for the three experimental conditions:
undirected monologue, directed monologue, and dialogue.Envelope TRFs for the
socially-relevant conditions exhibited stronger EEG prediction correlations (one-way
repeated measures ANOVA with Wilcoxon signed-rank post hoc comparisons; *p < 0.05, **p
*< 0.01). **(B)** The same analysis was repeated for all EEG sub-bands of interest. F-value*
of the repeated measures ANOVA is reported for each sub-band, indicating the effect
of social relevance across features and EEG frequency bands. The strongest effects
of social relevance for the envelope TRF emerged in the delta-band, and for word-
*onset in the gamma-band. **(C)** Scalp topographies of the EEG prediction correlations*
for each condition and speech feature. Warmer colours indicate higher prediction
correlation values. All topographies showed qualitatively similar spatial patterns,
compatible with previous studies on continuous speech listening (Di Liberto et al.,

    *2015). **(D)** TRF weights averaged across all participants at channel Cz are shown for*
the three conditions and stimulus features. Shaded areas represent the standard error
of the mean (SEM) across participants.

TRF and ANOVA analyses were repeated for different EEG sub-bands to determine if the effect was specific to certain cortical rhythms. The envelope TRF analysis showed strongest effects in low-frequency EEG (**Figure 1B**), with a statistically significant effect in the delta-band (F(2, 38) = 5.80, p = 0.006); a trending effect in the alpha-band (F(2, 38) = 3.18, p = 0.053); and no statistically significant effects elsewhere (theta-band: $F_{(2, 38)} = 0.27$, $p = 0.763$; beta-band: $F_{(2, 38)} = 1.89$, $p = 0.166$). Conversely, the word onset TRF showed strongest effects in higher EEG frequencies, with a statistically significant effect in the beta-band (one-sample ANOVA, $F_{(2, 38)} = 6.38$, $p = 0.004$); a trending effect in the alpha-band ($F_{(2, 38)} = 2.95$, $p = 0.064$); and no statistically significant effects elsewhere (delta-band: $F_{(2, 38)} = 0.61$, $p = 0.551$; theta-band: $F_{(2, 38)} = 0.57$, $p = 0.559$). No statistically significant effects were measured for the envelope derivative TRF (delta-band: $F_{(2, 38)} = 2.75$, $p = 0.077$; theta-band: $F_{(2, 38)} = 0.27$, $p = 0.765$; alpha-band: $F_{(2, 38)} = 1.45$, $p = 0.248$; gamma-band: $F_{(2, 38)} = 2.17$, $p = 0.128$).

We also report TRF weights for the models in the EEG broadband. Based on previous research, we expected a P1-N1-P2 pattern with latencies around 40, 80, and 140ms respectively (Crosse et al., 2016a; Di Liberto et al., 2015; Lalor & Foxe, 2010), where longer latencies have been linked with factors such as selective attention (Power et al., 2012). The literature also informs us on the neural dynamics corresponding with higher order features like word onsets, such as a large slow negative deflection at a latency of approximately 400 ms (Broderick, 2018). Here, we found TRF patterns that were qualitatively similar to that literature in all those aspects.

Experiment 2

EEG signals were recorded as participants listened to spontaneous two-party dialogues from popular podcasts, each involving a podcast host and a guest. Forward univariate TRFs were derived for the two speech features showing an effect of social relevance: the sound envelope and word onset features (**Figure 2**; we also report the envelope derivative results in **Figure S2**). As in Experiment 1, EEG prediction correlations were calculated with leave-one-out cross validation for each stimulus

486 feature separately. EEG prediction correlations were above chance for both sound
487 envelope (**Figure 2A-left**; *t*-tests, *p* < 0.001) and word onsets (**Figure 2B-left**; *t*-tests,
488 *p* < 0.001), reflecting a statistically significant EEG encoding of the two features.

489
Figure 2. Characterising the cortical tracking of speech using real podcast
dialogues. *Univariate forward TRF models in Experiment 2 for envelope **(A)** and word*
*onset **(B)** features. These analyses aggregated data from both listening conditions (AA*
and AC). Scalp distributions of the average EEG prediction correlations are reported
(left panels). TRF weights at channels Fz, Cz and Pz (middle-panels) and the TRF
weights at three key time-lags exhibit patterns compatible with the literature
(Chalehchaleh et al., 2025; Di Liberto et al., 2015; Lalor, 2009).

In Experiment 2, the podcast host was always an adult speaker, while the podcast
guest was either an adult or a child. This led to two conditions (adult-adult
conversations - AA; adult-child conversations - AC) that we compared to determine if
their multi-faceted differences (e.g., prosody, lexis, word-rate) would impact the
envelope and word onset TRF model fits. While there may be other differences that
exist elsewhere between AA and AC speech, for the purposes of the current analyses,
we focused on just these two speech features. EEG prediction correlations averaged
across scalp locations did not differ significantly between AA and AC stimuli (Wilcoxon
paired signed-rank test (*p <0.05): envelope: p = 0.177; word onset: p = 0.165). The
effect of stimulus type (AA versus AC) upon TRF feature weights at channel Cz was
also examined for each time lag. No statistically significant differences emerged (FDR-
corrected Wilcoxon paired signed-rank test, *p > 0.05 across all time lags). On this
basis, the analyses that follow are carried out by combining AA and AC stimuli. While

no effects of speech register emerged in this investigation, further work could

investigate possible effects on other linguistic features (Piazza et al., 2025).

The TRF weights were qualitatively comparable with the previous literature (Di Liberto

et al., 2015; Lalor et al., 2009) and with the results from Experiment 1, with the

envelope TRF presenting the typical P1-N1-P2 characteristics, and the word onset

TRF exhibiting the typical dominant negative deflection at about 400 ms (**Figure 2-**

middle). We measured Pearson's correlations between the group TRF weights across

the two experiments. The resulting group-level correlations after averaging the TRF

weights of the 64 EEG channels across participants are shown in **Figure 3,** indicating

high correlations (up to 0.92) across experiments, with average correlation values

increasing with social relevant i.e., undirected monologue, directed monologue,

synthesised dialogue.

Figure 3. Speech podcast TRFs are more correlated with synthesised speech
TRFs involving socially-relevant speech than undirected monologues.
Correlations between group-level average TRF weights of the 64 EEG channels were
calculated on the original samples from the two experiments. A bootstrap resampling
procedure was performed to assess the robustness of this pattern. The distribution
and confidence intervals of the bootstrap, the bootstrap mean (dot), and the group-
level average (diamond) TRF correlation are plotted. TRF weights for the real podcast
dialogues in Experiment 2 shows high correlations with the three conditions in
Experiment 1 in the following descending order: synthesised dialogue, directed
monologue, undirected monologue. This pattern emerges for all speech features in
the analysis, with statistically significant differences emerging only for the word onset
features across all conditions.

To assess whether this pattern was statistically reliable across participants, conducted

Friedman tests (due to Mauchly's violation of sphericity), followed by post-hoc pairwise

Wilcoxon signed-rank tests by considering each participant as a datapoint (N = 20).

This analysis revealed a statistically significant effect of condition on the inter-

experiment correlation when considering the word onset TRFs ($\chi^2(2)$ = 11.70, p =

0.003, Kendall's W = 0.29), while no statistical significant differences were detected

for envelope ($\chi^2(2)$ = 2.50, p = 0.29), and envelope derivative TRFs ($\chi^2(2)$ = 4.90, p =

0.09). Post-hoc tests on word onset TRFs indicated that synthesised dialogue TRFs

were significantly more correlated with real dialogue TRFs from Experiment 2 than

directed monologue TRFs (Wilcoxon signed-rank $r_{\text{synth-Dialog vs. real-Dialog}} > r_{\text{directed-M vs. real-}}$

$_{\text{Dialog}}$: $p < 0.001$), with the correlation further decreasing for monologue TRFs ($r_{\text{directed-M}}$

$_{\text{vs. real-Dialog}} > r_{\text{undirected-M vs. real-Dialog}}$: $p < 0.001$) from Experiment 1. This trend is consistent

with the correlation pattern synthesised dialogue > directed monologue > undirected

monologue as on the original sample.

Additional analyses were carried out to obtain a more detailed assessment of the

distribution and robustness of the cross-experiment correlations. Specifically, we

performed a bootstrap resampling procedure (with replacement) and generated 1,000

correlations between the group-averaged TRF in Experiment 2 to each of the

conditions in Experiment 1 for each feature. Compatibly with the participant-level

results, the bootstrap distributions indicate particularly robust inter-experiment

correlations for word onset TRFs, especially for synthesised dialogue.

Investigating the impact of dysfluencies on lexical expectation measurements

Experiment 2 involved real podcast recordings, which include dysfluencies by nature.

Indeed, the dysfluency types and frequencies can vary due to a variety of factors (e.g.,

speaker, social context, social anxiety, fatigue, familiarity with interlocutor, knowledge

of topic, speaking a native or non-native language). To investigate the impact of

dysfluencies on the lexical surprise and entropy feature and its corresponding brain

response, two lexical surprise and entropy vectors were constructed, including and

excluding dysfluencies in the speech material, respectively (**Figure 4A**). As such, we

carried out a characterisation of the dysfluency type and frequency for this specific

dataset split across conditions AA and AC (**Figure 4B**), finding a majority of filler-type

dysfluency and an overall frequency of dysfluent words of 3.36%, whereby ~5% of all
words were filler-type dysfluencies (e.g., "uh"/"like"/"you know"), ~1% were repetitions,
with less than 1% consisting of replacements and restarts (see **Materials and**
Methods and **Table S1**). The frequency distribution of these dysfluencies types was
consistent across podcast episodes, with a statistically significant main effect of
dysfluency type (repeated measures ANOVA: $F(4,60) = 28.49$, $p < 0.001$, $\eta^2 = 0.42$;
post hoc tests in **Table S2**). We found no statistically significant interaction of
dysfluency type and experimental condition (adult-adult podcast AA; adult-child
podcast AC) (repeated measures ANOVA: $F(4,60) = 1.04$, $p = 0.39$, $\eta^2 = 0.02$).

Figure 4. Investigating the impact of dysfluencies on LLMs. (A) *Audio transcripts*
were analysed using two different lexical surprise and entropy vectors. While both
vectors only included non-dysfluent words, for sake of comparison, lexical surprise
and entropy were calculated using Mistral by including or excluding dysfluent words
from the text input, producing the dysfluent and non-dysfluent feature vectors. ***(B)***
Dysfluencies were identified manually in the text material. We report the dysfluency
frequency organised according to dysfluency type and experimental condition (adult-
adult podcast AA; adult-child podcast AC). ***(C)*** *The correlations between Mistral's*
expectations when including or excluding dysfluent words in the calculation of lexical
surprise and entropy are shown, quantifying the potential impact of dysfluency on the
measurement of lexical expectations. Doubling the number of dysfluencies led to a
statistically significant effect of dysfluency. On this basis, future research should
measure dysfluencies in the speech material, since differences across participant
cohorts or conditions may reflect differences in the prevalence of dysfluent words in
the heard or produced speech.

Here, we investigated if the presence of dysfluent words in the speech material (and
its transcript) significantly impacts our ability to model the neural encoding of lexical

expectations. To that end, we introduced two additional stimulus features in
Experiment 2. These comprised word onset vectors modulated by lexical surprise and
entropy values calculated using Mistral (see **Materials and Methods**). The first vector
involved surprise values calculated when including all words, of which 3.3% of the
were dysfluent. The second vector only included non-dysfluent words (dysfluencies
were identified manually). However, dysfluent and non-dysfluent surprise values did
not differ in this dataset, with a Pearson's correlation between the two surprise vectors
of 0.98. Due to this high correlation, we also did not expect differences to report in the
coupling between these vectors and the corresponding EEG signals. While we could
measure a statistically significant neural encoding of lexical surprise (using either
dysfluent or non-dysfluent vectors) when compared to a baseline acoustic model
(**Figure S3**), this is not guaranteed to be the case in speech material with more
dysfluencies.

These virtually identical regressors suggest that at this low volume of dysfluencies,
using speech material that includes or excludes dysfluencies would render little to no
difference in the neural processing and encoding of speech. However, social speech
can widely vary in the number of dysfluencies depending on numerous factors (e.g.,
speaker, social context, anxiety, tiredness, familiarity with interlocutor, knowledge of
topic, speaking a native or non-native language). As a consequence, the neural
encoding of speech may be impacted when more dysfluencies are present. An
excessive number of dysfluencies (e.g., more dysfluent than non-dysfluent words) may
disrupt lexical prediction in both the LLM and the human brain. Hence, we then
assessed what level of dysfluency frequencies constitute an issue for lexical prediction
in LLMs and to what extent this sensitivity persists by manually increasing the number
of dysfluencies in the speech transcript. Specifically, we doubled and quadrupled the
existing dysfluencies in Experiment 2, calculating the correlation between the lexical
surprise and entropy values between the dysfluent and non-dysfluent vectors (**Figure
4C**). Lower correlations indicate a stronger impact of dysfluencies. We found a
significant impact of dysfluency frequency (FDR-corrected Wilcoxon paired sign-
ranked test: $p_{originalSur\ vs.\ doubledSur} < 0.001$, $p_{originalSur\ vs.\ quadSur} < 0.001$, $p_{originalEntro\ vs.}$
$_{doubledEntro} < 0.001$, $p_{doubledEntro\ vs.\ quadEntro} < 0.001$, $p_{originalEntro\ vs.\ quadEntro} < 0.001$), whereby
there was practically no sensitivity to dysfluencies for the frequency in the original
material (dysfluent-nondysfluent correlation ~ 1) and significant sensitivity to

dysfluencies once they were doubled, though this effect seems to level out.
Additionally, the strongest sensitivity overall emerged for entropy vectors (correlation
down to ~0.7).

Discussion

Speech communication is an ideal testbed for investigating human cognition, as it
involves a multitude of processes involved in the transformation of sound to meaning.
Such processes encompass mechanisms such as statistical prediction, attention and
cognitive effort. The possibility of extending that work to social speech opens up
opportunities for probing a whole range of processes that are typically excluded from
monologue listening research, such as social role and speaker familiarity,
communication accommodation, social anxiety, and many others (Di Liberto & Ip,
2025). However, this opportunity comes with the important challenge of having to
isolate the phenomena of interests from a much richer and harder-to-control set of
behaviours and neural processes. One key for disentangling those processes and
phenomena is to pinpoint how exactly these socially-relevant factors impact the spatial
and temporal patterns reflecting the neural processing of speech and language. Here,
we take a first step in that direction by means of a principled step-by-step approach,
determining the impact of including a social element in speech listening tasks
(excluding interaction in our scope of investigation).

The results from Experiment 1 indicate that social relevance substantially enhances
the neural tracking of speech. Importantly, this effect cannot be explained by
differences in low-level acoustic or speech-structure properties across stimulus
categories. To assess this possibility, we quantified several stimulus characteristics,
including duration, root-mean-square (RMS) amplitude, mean fundamental frequency
(F0), F0 variability, and word rate. No systematic differences were observed between
conditions, nor between monologue and dialogue stimuli. These findings indicate that
the observed neural effects are unlikely to be driven by basic acoustic or speech-rate
differences and instead reflect modulation by socially relevant aspects of the stimuli.
The increased encoding emerged specifically for the speech envelope, but not for
word-level processing (EEG prediction correlation metric; **Figure 1A**). Previous work
on selective attention has also shown similar increases in envelope tracking.

Interestingly, however, the increased speech tracking in selective attention tasks also
corresponded with changes in the TRF patterns, especially with a modulation of the
TRF-P2 component (Carta et al., 2024; Power et al., 2012; Vanthornhout et al., 2018),
while the increased envelope tracking here does not correspond to changes in the
temporal or spatial dynamics of the TRF (**Figure 1D**). As such, our results are
compatible with a general increased engagement with the task and stimulus, but with
important differences from the widely studied selective attention phenomenon. It
should be noted that we also detected sensitivity to the social relevance of speech in
the beta band, specifically for word onset encoding. This result supports the potential
and relevance of measuring cortical tracking at higher EEG rates (even though
stronger speech tracking typically emerges in the delta and theta bands). While we did
not have specific hypotheses pertaining to the EEG beta band in this study, we note
that studying speech tracking at different frequency rates may provide an additional
dimension for more clearly disentangling the cortical tracking of distinct speech
features, and should be considered (without forgetting about the risks that come with
filtering operations) (De Cheveigné & Nelken, 2019).

While Experiment 1 isolated the impact of social relevance on speech tracking, our
design in Experiment 2 takes us a step further by measuring neural dynamics in more
naturalistic settings, wherein spontaneous speech introduces additional complexities.
The results in Experiment 2 (**Figure 2**) demonstrated temporal response
characterisations that are not only consistent with previous monologue-listening
literature (Broderick et al., 2018; Di Liberto et al., 2021; Heilbron et al., 2022), but also
with the neural speech encoding results obtained in Experiment 1 (**Figure 3**). Our
results indicated similar spatial and temporal patterns for synthesised and real speech
recordings; the EEG encoding of speech features, including the speech envelope and
word onset, was highly correlated across synthesised and real speech recordings,
achieving correlations of up to 0.92 for word onset TRFs. This is quite remarkable,
considering that the two experiments involved completely different speech material
and distinct participant cohorts. The fact that we observed a consistent increase in that
similarity with social relevance is also a strong indication that the impact of social
relevance on the spatial and temporal patterns of speech is consistent across material
and participants. Further work is needed to better characterise and test the boundaries
of consistency in neural tracking of speech features across synthesised and real

speech and to characterise the sensitivity of TRFs to variations in synthesiser model
and configuration. Nonetheless, our findings suggest that the similarity between
synthesised and real speech TRFs and their modulation by the social relevance of
speech are remarkably high.

Experiment 2 demonstrates the feasibility of employing the TRF using naturalistic and
socially-relevant speech material. Our dysfluency analysis indicated that the presence
of dysfluencies observed in the podcast stimuli utilised does not impact the ability to
derive indices of surprise and entropy, which can, in turn, be used to improve TRF
models. This is promising for future research, as it suggests the possibility of studying
speech tracking even in presence of dysfluencies. However, note here that the LLM
dysfluency simulation analysis highlights that when a larger portion of the speech
material includes dysfluencies, they can exert a notable impact on lexical expectation
indices.

From this analysis, two key conclusions emerge. First, dysfluencies do not pose a
problem up to a certain threshold, which may eliminate the need for manual transcript
cleaning when using LLMs. Second, it is essential to quantify and report the dysfluency
profile - both type and frequency - used in a given experiment, particularly when
generating new stimuli. This task is relatively straightforward in listening experiments,
where all participants are exposed to identical stimuli. In contrast, speech interaction
paradigms, such as hyperscanning studies, introduce greater methodological
complexity. A central challenge is that individual participants often exhibit unique
dysfluency profiles (Befi-Lopes et al., 2014; Pistono et al., 2024), which can
differentially affect measures of lexical expectations - both in terms of LLM-based
estimation and neural encoding (Goldstein et al., 2022). This variability reinforces the
need for systematic dysfluency measurement and individualised reporting to ensure
robust and interpretable findings. Finally, it is worth noting that the observed sensitivity
of LLMs to dysfluency opens new avenues for research focused on its impact on the
neural encoding of speech. Future investigations could specifically examine this effect
in participant cohorts with speech production challenges, such as individuals who
stutter or those experiencing social anxiety. Such studies would not only deepen our
understanding of speech processing under naturalistic conditions but also offer
insights into the neural mechanisms underlying communicative difficulties.

In summary, this study demonstrates that incorporating social relevance into speech
listening tasks enhances the neural tracking of speech and reveals remarkable
consistent spatial-temporal patterns across varied speech materials and participant
cohorts. By extending analysis to spontaneous speech and dysfluency, we
demonstrate the possibility of modelling the neural tracking of speech, as well as
lexical prediction mechanisms even in presence of dysfluencies. Based on our
simulation, we provide evidence for the importance of considering the issue of
dysfluency when studying social speech, when listening to speech interactions and
especially in experiments where participants engage in dialogue. Altogether, these
findings lay the groundwork for future investigations into the neural encoding of socially
complex and dysfluent speech, offering a principled path toward understanding
communication in real-world contexts.

Supplementary

Figure S1. EEG prediction correlations for different frequency bands (Experiment 1). *Univariate forward TRFs were fitted for envelope, envelope derivative, and word onset features to predict the EEG signal at frequency bands delta (A), theta (B), alpha (C), and beta (D). EEG prediction correlations (Pearson's r), averaged over all channels and participants, are presented as mean ± SEM, comparing the three experimental conditions: undirected monologue, directed monologue, and dialogue. One-way ANOVA and Wilcoxon signed-rank post hoc tests were conducted (*p < 0.05, **p < 0.01). This data extends the results in Figure 1.*

743 ***Figure S2.*** *TRF weights for the envelope derivative feature from Experiment 2 at three*
744 *selected EEG channels. This data extends the results in Figure 2.*

Dysfluency Type	Definition
Fillers	uh / um, you know, sort of, like, I'll say, I don't know
Repetitions	I-I-I
Restarts (also coined *false start* in (Shriberg, 1999))	*The speaker abandons completely the initial utterance and restarts it. This type of dysfluency does not actually involve a repair as the speaker begins a completely new utterance(Passali et al., 2022).*
Replacements (also coined *repair* in (Shriberg, 1999))	*The speaker replaces the dysfluent word(s) or phrase with the fluent one. In replacements, the reparandum is replaced by the repair (Passali et al., 2022)*

Table S1. Dysfluency types. *The dysfluency analyses in Figures 4 and 5 constitute*
a preliminary approach for handling dysfluencies that are a natural part of social
speech. This table indicates the four dysfluency types that were identified – fillers,
repetitions, restarts (also known as false starts), and replacements (also known as
repairs) (Shriberg, 1999). See Figure 4B for the distribution of these dysfluency types
in the speech material for Experiment 2. Note that Experiment 1 did not include any
dysfluencies.

		Mean Difference	SE	t	Cohen's d	p_{holm}
Fillers	**Repetitions**	5.812500	0.801358	7.253313	1.671350	7.394361×10^{-9}
	Replacements	6.718750	0.801358	8.384206	1.931937	9.777329×10^{-11}
	Restart	6.968750	0.801358	8.696177	2.003823	3.213573×10^{-11}

752 **Table S2. Post Hoc Comparisons of dysfluency types**. *A statistically significant*
753 *main effect of dysfluency type emerged when comparing the frequency distribution of*
754 *dysfluencies across trials in both AA and AC conditions with a repeated measures*
755 *ANOVA. Post hoc tests show how fillers significantly outnumber any other dysfluency*
756 *produced in real podcast dialogues. This data extends the results in Figure 4B.*

757
Figure S3. Investigating the impact of dysfluencies on speech TRFs. (C)
Multivariate TRFs were computed to predict the EEG signal using the dysfluent or non-
dysfluent vectors, concatenated with nuisance regressors capturing speech acoustics
and word segmentation (envelope, envelope derivative, and word onset). We report
the TRF weights corresponding to the dysfluent and non-dysfluent surprise features
at electrode Cz. (D) Both versions of the models that included acoustics and dysfluent
surprises returned a significant gain in the average EEG prediction correlations in
comparison to the acoustics-only mTRF, reflecting a statistically significant neural
encoding of lexical surprise. These data extend the results in Figure 4.

Forward multivariate TRFs were fit for each of the two vectors and including the three
features used in the earlier analysis as nuisance regressors (envelope, envelope
derivative, and word onsets), leading to a *non-dysfluent surprise TRF* and a *dysfluent*
surprise TRF. Note that, for sake of comparison, dysfluent and non-dysfluent vectors
were built to be identical in everything but the surprise values i.e., they had the same
word onsets. Finally, a third TRF was fit for the nuisance regressors without any lexical
surprise information (*baseline* TRF). When comparing the model weights between the
non-dysfluent and dysfluent surprise TRFs (Wilcoxon paired signed-rank test: p =
0.951; **Figure S3a**), no statistically significant differences emerged. In addition, EEG
prediction correlations were compared across the three TRF models, showing a
statistically significant differences across the three TRF models (repeated measures
ANOVA: $F(1, 19)$ = 7.96, p = 0.001, η^2 = 0.30; **Figure S3b**). *Post hoc* comparisons
indicated larger EEG prediction correlations when including either surprise vector (*t*-
tests; $r_{non-dysfluent\ surprise}$ > $r_{no-surprise}$, $t(19)$ = -3.54, p_{holm} = 0.003, Cohen's d = -0.16; $r_{dysfluent}$
$_{surprise}$ > $r_{no-surprise}$, $t(19)$ = -3.36, p_{holm} = 0.004, Cohen's d = -0.16). As expected, no
statistically significant differences were measured between non-dysfluent and
dysfluent surprise TRF prediction correlations (*t*-test: $t(19)$ = -0.18, p_{holm} = 0.861,
Cohen's d = -0.01).

Figure S4. Behavioural validation of stimulus categorisation into directed vs.
undirected. An independent group of participants (*N* = 15) was asked to categorise
speech recordings as directed or undirected. 16 recordings, each ~2-minute long,
were used in this experiment. The confusion matrix reports correct and incorrect
classification rates.

Measure	Comparison	Test	*p*-value	Effect size
Duration	Undirected vs Directed vs Dialogue	Kruskal–Wallis	0.57	0.01 (ε^2)
RMS amplitude	Undirected vs Directed vs Dialogue	Kruskal–Wallis	0.17	0.08 (ε^2)
Mean F0	Undirected vs Directed vs Dialogue	Kruskal–Wallis	0.93	0.01 (ε^2)
F0 variability	Undirected vs Directed vs Dialogue	Kruskal–Wallis	0.40	0.02 (ε^2)
Word rate	Undirected vs Directed vs Dialogue	Kruskal–Wallis	0.15	0.09 (ε^2)
Duration	Monologue vs Dialogue	Wilcoxon rank-sum	0.30	0.18 (r)
RMS amplitude	Monologue vs Dialogue	Wilcoxon rank-sum	0.51	0.12 (r)
Mean F0	Monologue vs Dialogue	Wilcoxon rank-sum	0.72	0.06 (r)
F0 variability	Monologue vs Dialogue	Wilcoxon rank-sum	0.51	0.11 (r)
Word rate	Monologue vs Dialogue	Wilcoxon rank-sum	0.32	0.18 (r)

791   ***Table S3. Acoustic and speech-structure properties across conditions***. No
792   significant differences were observed between undirected monologue, directed
793   monologue, and dialogue conditions, nor between monologue and dialogue stimuli.

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