EEG-based speaker-listener neural coupling reflects speech-selective

attentional mechanisms beyond the speech stimulus

Jiawei Li[1, 3, 5], Bo Hong[2, 3], Guido Nolte[4], Andreas K. Engel[4], Dan Zhang[1, 3*]

[1]Department of Psychology, School of Social Sciences, Tsinghua University, Beijing 100084,

China

[2]Department of Biomedical Engineering, School of Medicine, Tsinghua University, Beijing

100084, China

[3]Tsinghua Laboratory of Brain and Intelligence, Tsinghua University, Beijing 100084, China

[4]Department of Neurophysiology and Pathophysiology, University Medical Center Hamburg

Eppendorf, Hamburg 20246, Germany

[5]Department of Education and Psychology, Freie Universität Berlin, Habelschwerdter Allee,

Berlin 14195, Germany

[*]**Correspondence**:

Dan Zhang, Ph.D.

Room 334, Mingzhai Building, Tsinghua University, Beijing 100084, China

E-mail: dzhang@tsinghua.edu.cn; Tel: +86-10-62796737

A short running title (less than 40 characters):

inter-brain coupling beyond the stimulus

Three key points

1. Listener's EEG signals coupled to the speaker's 5 s before the speech onset,

which revealed a "beyond the stimulus" attentional modulation.

2. Speaker-listener attentional coupling is correlated to the listener's comprehension

performance, but the speech-listener's coupling didn't.

3. The implementation of temporal response function methods and the neural

language methods yielded novel perspectives to the analysis of the inter-brain

studies.

Abstract

When we pay attention to someone, do we focus only on the sound they make, the word they use, or do we form a mental space shared with the speaker we want to pay attention to? Some would argue that the human language is no other than a simple signal, but others claim that human beings understand each other not only by relying on the words that have been said but also formed a shared ground in the specific conversation. This debate was raised early, but the conclusion remains vague. Our study aimed to investigate how attention modulates the neural coupling between the speaker and the listener in a cocktail party paradigm. The temporal response function (TRF) method was employed to reveal how the listener was coupled to the speaker at the neural level. The results showed that the neural coupling between the listener and the attended speaker peaked 5 seconds before speech onset at the delta band over the left frontal region, and was correlated with speech comprehension performance. In contrast, the attentional processing of speech acoustics and semantics occurred primarily at a later stage after speech onset and was not significantly correlated with comprehension performance. These findings suggest that our human brain might have adopted a predictive mechanism to achieve speaker-listener neural coupling for successful speech comprehension.

Keywords: attention, electroencephalogram, temporal response function, natural speech, inter-brain

Introduction

"Men do not understand one another by actually exchanging signs for things…they do

it by striking the same note on their mental instruments."

Wilhelm von Humboldt

When we pay attention to someone in a cocktail party, what sort of information

from the speaker do the listeners exactly attend to? Are the listeners just perceiving

sound waves or process semantic meaning in the speech that the speaker generated? Or

do listeners and the speakers reach some common ground that is beyond the speech

stimulus itself? The question is about the core connection formed between the speaker

and the listener via speech. There are different perspectives on this issue: Some argued

the speech is just a type of signaling (Chomsky, 1959; Skinner, 1957); others suggested

that speakers and listeners need to be intentionally aligned, rather than relying solely

on speech (Gadamer, 1975; Searle, 1980). The discourse has been primarily conducted

at the theoretical level. However, empirical evidence, especially in neuroscience is still

lacking to settle the debate.

The cocktail party paradigm together with the emerging inter-brain approach

provides a unique and advantageous tool for examining this issue. The cocktail party

established an ideal benchmark for contrasting the information that the listener

deliberately focused on versus that which they disregarded (Cherry, 1953; McDermott,

2009; Middlebrooks et al., 2017; Shinn-Cunningham, 2008). In this situation, listeners

would receive the information from different speakers simultaneously, but the listener

only processes the speech that she or he attended in a cocktail party. The inter-brain

method entails the acquisition of neural data from both the speaker and the listener and

introduces the speaker's neural activity as a new reference in the speech comprehension

(Hasson et al., 2012; Jing Jiang et al., 2021; Z. Li et al., 2021; Pérez & Davis, 2023;

Redcay & Schilbach, 2019; Schoot et al., 2016; Stolk et al., 2016). Using this method,

the neural coupling with the speaker and the neural coupling with the speech could be

compared simultaneously. If the listener only perceives the information she or he

received in the speech as the behaviorism scientists suggested (Chomsky, 1959; Skinner,

1957, 1986), the speaker-listener coupling would be largely bounded with the neural

coupling with the speech stimuli itself. In other words, the neural coupling with the

speaker would occur at the similar time window or the similar frequency band with the

neural coupling with the speech stimuli. However, if the listener coupled with the

speaker "beyond the stimulus"(Hartley & Poeppel, 2020) or "extra-linguistic"(Schoot

et al., 2016), the time and frequency bands for the attention modulation of the speaker's

neural activity may reveal a different pattern comparing the speech-listener's coupling.

Previous inter-brain studies have provided potential evidence of the beyond the

speech attention. In the monologue situation, inter-brain studies revealed the distinct

pattern of the speaker-listener coupling comparing with the speech-listener coupling:

the speaker-coupling coupling occurs before the speech onset (Dai et al., 2018; Jing

Jiang et al., 2015; L. Liu et al., 2020; Stephens et al., 2010; Zheng et al., 2018) or on

the different frequency bands (Pérez et al., 2017). However, studies on speaker-listener

coupling in a cocktail party scenario remains sparse. There are few studies investigating

the attentional speaker-listener coupling or entrainment (Dai et al., 2018; Kuhlen et al.,

2012). Direct comparison of neural mechanisms underlying attention modulation in

speaker-listener coupling and listener-stimuli coupling is still insufficient. Previous

studies only compared the coupling between the speaker-listener and speaker-sound, or

the acoustic feature (Dai et al., 2018; Pérez et al., 2017). To our knowledge, there have

been no studies that directly compare the semantic features and the inter-brain features

in a cocktail party scenario. However, the meaning, or the semantic feature is vital to

understand the competing speech streams. Recent studies also revealed that processing

of semantic or linguistic features could also be modulated by attention, for the features

in the attended stream could also better understood than the unattended stream

(Broderick et al., 2018; Connolly et al., 1990; Dai et al., 2022; Har-shai Yahav & Zion

Golumbic, 2021; Heil et al., 2004). It still doesn't know whether the speaker and listener

aligned to each other sole rely on the meaning in the speech or have the extra-linguistic

attention modulation (Hartley & Poeppel, 2020; Hasson et al., 2012; Pérez & Davis,

2023; Schoot et al., 2016; Stolk et al., 2016).

The objective of this study was to compare the attention to the speech and the

speaker using a cocktail party paradigm. More specifically, we investigated the speaker-

listener coupling in contrast to the coupling between the listener's neural activity and

two types of features: the acoustic and semantic features. Naturalistic speech was used

as stimulus material, which contains much richer information than either the sound

sequence or single word stimulation employed in previous studies. The sound, the

meaning, and the speaker could appear at the same time in an ecological situation and

enables us to directly compare couplings to the different features simultaneously

(Broderick et al., 2018; Hamilton & Huth, 2018; Hartley & Poeppel, 2020; Nastase et

al., 2021; Sonkusare et al., 2019; Willems et al., 2020). A sequential dual-brain

approach(Redcay & Schilbach, 2019) was used, and the electroencephalogram (EEG)

of both speaker and listener was recorded. A temporal response function (TRF) method

was used to measure the difference between the coupling to attended and unattended

features. We also used the neural language processing (NLP) models to help us extract

the meaning in the text as the semantic feature. Based on previous studies, we

hypothesized that speaker-listener coupling was distinct from the speech-listener

coupling, which reveals beyond the speech attention modulation: for the time course,

The listener's neural activity is aligned with the speaker's neural activity in a broader

time range, may have a leading coupling (Dai et al., 2018; Kuhlen et al., 2012; Stephens

et al., 2010; Zheng et al., 2018). The attention modulation for the acoustic and semantic

feature occurs after the speech onset (Ding & Simon, 2012a, 2012b; Hillyard et al.,

1973; O'Sullivan et al., 2015; Power et al., 2011; Teoh et al., 2022), and the attention

effect of the semantic feature lasts longer (Broderick et al., 2018; Dai et al., 2022). For

the frequency bands, we hypothesize that the delta band reflects attention modulation

for the inter-brain feature and the semantic feature for they related to the comprehension

(Bai et al., 2022; Dai et al., 2022; Lu, Jin, Ding, et al., 2022; Teoh et al., 2022; Yu et al.,

2022; Zuk et al., 2021), and the theta band represents the acoustic feature (Ding et al.,

2014; Etard & Reichenbach, 2019).

Figure 1 The stimuli, the experimental paradigm and the analysis process

(A) The paradigm and the different features. A "cocktail party" selective attention paradigm was used, in which the listener was asked to pay attention to one side of the speech stream and ignore the other. While listening to the speech stream, the listener's EEG signals were recorded. The speaker's EEG signals were defined as the "inter-brain" features in our study. The speaker-listener coupling is "beyond the stimulus". The acoustic and semantic features were included in the coupling between the speech stimuli and the listener. (B) The data analysis process. The encoding models were trained for different features separately. The Encoding r for the attended feature and the unattended feature was calculated time point by time point by applying the TRF method. The TRF of each feature was estimated separately in two frequency bands: delta and theta.

Materials and Methods

Ethics statement

The study was conducted in accordance with the Declaration of Helsinki and was approved by the local Ethics Committee of Tsinghua University. Written informed consent was obtained from all participants.

Participants

Two participants (both male, aged 26 and 24 years) were recruited for this study as speakers. Both speakers were from the broadcasting station of Tsinghua University and had experience related to broadcasting and hosting.

Twenty college students (10 females; mean age: 24.7 years; range: 20–43 years) from Tsinghua University participated in the study as paid volunteers for listeners. All participants were native Chinese speakers and reported having normal hearing and normal or corrected-to-normal vision.

The sample size (N = 20) was decided empirically following previous TRF-based studies on human speech processing (Broderick et al., 2018; Di Liberto et al., 2015; J. Li et al., 2022; Mirkovic et al., 2015).

Experimental procedure for the speakers

A sequential inter-brain approach was adopted by the present study (Redcay &

Schilbach, 2019), in which the neural activities of the speakers were recorded prior to

the listeners. The sequential design was more appropriate for this study than the real-

time interactive design because the speakers' audio and neural activity remained

consistent for all listeners (Leong et al., 2017; Y. Liu et al., 2017; Stephens et al., 2010).

In this experiment, each speaker participated in 30 trials, each of which was

approximately 51–76 seconds in length, while the speakers' audio signals and EEG

signals were recorded. The experimenter selected 28 trials for the listener's experiment,

excluding the two most unqualified trials.

The speaker first read the relevant material on the screen. There was a wide variety

of content to be covered, including one's hometown, a recent book, a fable, etc. The

speaker could decide how long they wanted to spend on preparation and start talking

when they were fully prepared (the length of preparation was usually 3 minutes). When

the speaker was prepared, they would press the spacebar on the computer keyboard,

and the recording would begin. When the spacebar was pressed, three 1000 Hz pure

tone cues were triggered (duration: 1000 ms; cue interval: 1500 ms). The cues were

presented as event markers, synchronized with the sound in the listener's experiment to

ensure that the neural signals of the speaker and listener remain aligned with the sound

stimuli. The speaker was asked to start speaking immediately after the end of the third

beep (within approximately 3 s). A fixation and a countdown timer appeared on the

screen during the talking part. The speaker was asked to stare at the fixation and to

complete the speaking as clearly, completely, and naturally as possible. During the

recording process, the experimenter listened to the speaker's narration simultaneously

and controlled the quality. The experimenter had the right to ask the speaker to retell

the clip if there was a reason for the lack of fluency, length, etc., that might affect the

listeners' perception. The materials of both speakers' content were varied. Between each

trial, the speakers were allowed to rest on their own. During the experiment, the

speakers were asked to control their head movements and facial muscles to obtain better

quality EEG signals.

The speech stimuli were recorded from two male speakers using the microphone

of an iPad2 mini (Apple Inc., Cupertino, CA) at a sampling rate of 44,100 Hz.

Experimental procedure for the listeners

The experiment consisted of four blocks, each containing seven trials. Two speech

streams were presented simultaneously in each trial, one to the left ear and the other to

the right ear. Two speech streams of the same trial matched the volume, i.e., the root

mean squared intensity of the amplitude of the speech streams in the same trial were

the same. The participants were instructed to attend to one spatial side according to the

hints on the screen ("Please pay attention to the [LEFT/RIGHT]"). Considering the

possible duration difference between the two audio streams, the researchers set the end

of the trial after the longer speech audio had ended. Each trial began when participants

pressed the SPACE key on the computer keyboard. A white fixation cross was also

displayed throughout the trial. The speech stimuli were played immediately after the

keypress and were preceded by the three beep sounds.

At the end of each trial, four multiple-choice questions (two for the attended story

and the other two for the unattended story) were presented sequentially in random order

on the computer screen. Each question had four options, and participants entered the

letter of the correct option as their answer. The listeners were not explicitly informed

about the correspondence between the questions and the stories. For instance, one

question following a story about one's hometown was, "What is the most dissatisfying

thing about the speaker's hometown? (推测讲述人对于家乡最不满意的地方在于?)",

and the four choices were A) There is no heating in winter; B) There are no hot springs

in summer; C) There is no fruit in autumn; D) There are no flowers in spring (A. 冬天

没暖气; B. 夏天没温泉; C. 秋天没水果; D. 春天没鲜花). The single-trial

comprehension accuracy could be 0% (two wrong answers), 50% (one correct answer),

or 100% (two correct answers) for both the attended and the unattended stories. No

feedback was given as to whether the questions were answered correctly.

After completing these questions, participants rated their concentration level of the

attended stream, the experienced difficulty performing the attention task, and the

familiarity with the attended material using three 10-point Likert scales. Throughout

the trial, participants were required to maintain visual fixation on the fixation cross

while listening to the speech. Meanwhile, they were asked to minimize eye blinks and

all other motor activities. The participants were recommended to take a short break

(around 1 min) after every trial within one block and a long break (no longer than 10

min) between blocks.

In each block, the side being attended to was fixed (two blocks for attending to the

left side and two for attending to the right side). Within each block, the identity of the

speaker is kept constant on the left and right sides. The to-be-attended spatial side and

the corresponding speaker identity were balanced within the participant, with seven

trials per side for both speakers. The assignment of the stories to the four blocks was

randomized across the participants.

The experiment was conducted in a sound-attenuated, dimly lit, and electrically

shielded room. The participants were seated in a comfortable chair in front of a 19.7-

inch LCD monitor (Lenovo LT2013s). The viewing distance was approximately 60 cm.

The experimental procedure was programmed in MATLAB using the Psychophysics

Toolbox 3.0 extensions (Brainard & Brainard, 1997). The speech stimuli were delivered

binaurally via an air-tube earphone (Etymotic ER2, Etymotic Research, Elk Grove

Village, IL, USA) to avoid possible electromagnetic interference from auditory devices.

The volume of the audio stimuli was adjusted to be at a comfortable level (~70 dB SPL)

that was well above the auditory threshold. The average presentation level was

measured with a B&K (Brüel & Kjær, Nærum, Denmark) Sound Level Meter (Type

2250 Investigator) with a 1-inch Free-field Microphone (Type 4144) and an Artificial

Ear (Type 4152).

Data acquisition and pre-processing

EEG was recorded from 60 electrodes (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, and

O1/2), which were referenced to an electrode between Cz and CPz, with a forehead

ground at Fz. A NeuroScan amplifier (SynAmp II, NeuroScan, Compumedics, USA)

was used to record EEG at a sampling rate of 1000 Hz. Electrode impedances were kept

below ten kOhm for all electrodes.

The recorded EEG data were subjected to an artifact rejection procedure using

independent component analysis. Independent components (ICs) with large weights

over the frontal or temporal areas, together with a corresponding temporal course

showing eye movement or muscle movement activities, were removed. The remaining

ICs were then back-projected onto the scalp EEG channels, reconstructing the artifact-

free EEG signals. While the relatively long duration of the speech trials in the present

study (~1 minute per story, see Experimental procedure) has made it more difficult for

the participants to avoid inducing movement-related artifacts as compared to the

classical ERP-based studies, a temporally continuous, non-interrupted EEG segment

per trial was preferred for the employment of the ICA method. Therefore, any ICs with

artifact-like EEG activities for more than 20% of the trial time (i.e., ~12 sec) were

rejected, leading to around 4–11 ICs rejected per participant. The cleaned EEG data

were used for the mTRF analysis without any further artifact rejection procedures.

Next, the EEG data were segmented into 28 trials according to the markers

representing speech onsets. The analysis window for each trial was extended from 10

to 55 s (duration: 45 s) to avoid the onset and the offset of the stories.

The pre-processed EEG signals were re-referenced to the average of all scalp

channels and then downsampled to 128 Hz before the modeling. Then, the EEG data

were filtered in delta (1–4 Hz) and theta (4–8 Hz) (filter order: 64, one-pass forward

filter). The use of a causal FIR filter ensured that filtered EEG signals were decided

only by the current and previous data samples (de Cheveigné & Nelken, 2019), which

is essential for accurate time-course analysis. The filter order of 64 was chosen to keep

a balance of temporal resolution and filter performance: the filtered EEG signals were

therefore calculated based on the preceding 500 ms data (64 at 128 Hz).

Speaker-listener coupling: Inter-brain Features

288   The inter-brain method enables us to study how the listeners are aligned with the

289   attended speaker (Dai et al., 2018; Hasson et al., 2012; Jing Jiang et al., 2021; Stephens

290   et al., 2010). We used the speaker's neural activity as the representation of the inter-

291   brain feature. The speaker's EEG served as the inter-brain feature. It followed the same

292   pre-processing procedure as the listener's EEG.

Speech listener coupling: acoustic and semantic features

● *Acoustic Features*

The amplitude envelope of the speech represented the acoustic features of the

speech. It was obtained using a Hilbert transform and then down-sampled to the same

sampling rate of 128 Hz.

● *Semantic Features*

The original audio recorded by the speaker during the EEG recording was converted

to the text first automatedly by *Iflyrec* software (Iflytek Co., Ltd, Hefei, Anhui) and then

double checked manually. The onset time of every word was extracted during this

process.

The recent emergence of Natural Process Language (NLP) models has enabled the

description of the semantic features in speech (Brodbeck et al., 2022; Broderick et al.,

2018, 2021). Next word prediction is one of the fundamental NLP tasks using the

semantic information in the texts (Schrimpf et al., 2021; Vaswani et al., 2017). The goal

of the task is to predict the next word when given a sequence of words $W_1, W_2, ..., W_{t-1}$,

which was consistent with the human understanding process. The probability of the

next word is $P(W_t|W_1, ..., W_{t-1})$ and can be calculated by varied NLP models. The

surprisal of the word was defined as follow, which reflected how surprised the next

word (Willems et al., 2016):

$$surprisal(t) = -\log P(W_t|W_1, ..., W_{t-1}) \tag{1}$$

The index was calculated based on ADAM, a widely accepted classical natural

language process model (Bengio et al., 2003; Kingma & Ba, 2015). The model was

trained on a couple of *People's Daily.* There were 534,246 words involved in the model

training. 66,781 words were in the cross-validation set, and 66,781 words used as a test

set. The details of the model are described in Table 1.

After calculating the surprisal index of every word, we generated a "semantic vector"

at the same sampling rate as the EEG data (Broderick et al., 2018). The vectors

contained the time-aligned impulses at the start of each word of the surprisal value for

every audio clip.

Table 1 Details of Neural Language Processing Model

Type	Parameter
model type	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

Temporal response function modeling

The analysis workflow for the analysis related to the attended speech stream is

shown in Figure 1. The neural responses to the three different features were

characterized using a temporal response function (TRF)-based modeling method

(Crosse et al., 2016, 2021).

Three different features mentioned above are the input signal required by TRF. The

corresponding neural response $r(t, n)$ can be formulated as follows:

$$r(t,n) = \sum_{\tau} w(\tau,n)\, S(t-\tau) + \varepsilon(t,n) \qquad (2)$$

where $r(t,n)$ is the actual EEG response at every channel n; $t = 1 \ldots T$ is the time

point; $S(t-\tau)$ means the multivariate stimulus representation; $w(\tau,n)$ is the channel

specific TRF at lag τ and $\varepsilon(t,n)$ is the residual.

The TRF is estimated by minimizing the mean square error between the actual

neural response $r(t,n)$ and the neural response predicted by the model $\hat{r}(t,n)$. The

Pearson's correlation between the actual neural response and predicted neural response

was referred as Encoding *r*. The mTRF toolbox (Crosse et al., 2016) was used to

estimate the TRF(w) as fellow:

$$w = (S^T S + \lambda I)^{-1} S^T r \qquad (2)$$

where λ is the ridge regression parameter, *I* is the identity matrix, and the matrix

S is the stimulus matrix. The lambda varied from 10^{-1} to 10^{8} (lambda = 10^{-1}, 10^{0}, …,

10^{8}) to make the model optimal (Crosse et al., 2021; Teoh et al., 2022). The λ value,

which produces the highest encoding *r*, averaged across trials and channels, was

selected as the regularization parameter for all trials per participant (Broderick et al.,

2019). The cross-validation procedure was implemented in a leave-one-trial-out

manner: the TRFs were trained based on data from 27 trials and tested on the left-out

trial each time.

The TRF was trained at every individual time lags of -8 s to 8 s to investigate the

specific interval of attentional modulation for each feature. At a sampling rate of 128

Hz, there are 2049 individual time-lag intervals of 7.625 ms. The choice of the window

was aligned with the previous fMRI and fNIRS inter-brain studies.

The TRF calculation procedure was performed for the EEG signals from each EEG

channel filtered at the delta and theta frequency bands. Only attended features are used

as input to the model, and TRFs trained by the attended features and the neural response

were applied to the tests of attended and unattended features, referred to as Encoding

Encoding r_{att} and Encoding r_{unatt}, respectively. This approach was largely used in the

attention-related studies (Mirkovic et al., 2015; O'Sullivan et al., 2015).

Quantification and statistical analysis

The paired t-tests were performed to investigate the attentional modulation for

different features, contrasting the encoding r of the attended speech versus the

unattended speech at each time lag. The Fisher-z transformation was used to convert r

encoding into z-score (Corey et al., 1998). The transformation transformed the r values

into a normal distribution that can be compared and assessed for statistical significance.

A nonparametric cluster-based permutation analysis was applied to account for

multiple comparisons (Maris & Oostenveld, 2007). In this procedure, neighboring

channel-latency bins with uncorrected t-tests p-value below 0.05 were combined into

clusters, for which the sum of the correlational *t*-statistics corresponding to the *t*-tests

was obtained. The combining process was initially automated by the toolbox and then

manually double-checked. Two clusters were combined if they shared a similar spatial

distribution (e.g., the clusters have a large overlap of the electrons). A null distribution

was created through permutations of data across participants ($n = 1,000$ permutations),

which defined the maximum cluster-level test statistics and corrected *p*-values for each

cluster. Clusters with *p*-values below 0.01 based on clusters were selected for further

analysis.

The above statistical analysis followed the standard cluster-based permutation

procedure as employed in classical ERP and related studies (Arnal et al., 2011; Henry

& Obleser, 2012; Zhang et al., 2012). Note that the reported *p*-values were only

corrected for the tests performed within each frequency band by using cluster-based

permutation tests. No multiple comparison correction was employed across different

frequency bands.

Correlation between clusters

The *Spearman* correlation of Encoding r_{att} in each cluster was calculated for each

pair of clusters to analyze the correlation between them.

Partial correlation between the behavioral performance and the neural activity

The partial correlation between the behavioral performance and the coefficients in

was calculated to reveal the unique contribution of a certain feature to the

comprehension performance. Specifically, the correlations between the mean Encoding

r_{att} in the cluster and the accuracy of the questions were calculated while controlling

other features.

Data Availability Statement

Data generated in our study has been uploaded to Mendeley Data (Li, Jiawei (2022),

"Speaker-listener Cocktail Party", Mendeley Data, V3, doi: 10.17632/wkh75s27kn.3).

When existing toolboxes were used for data analysis, citations have been provided.

Results

Behavioral Performance of the Listeners

The average comprehension performance was significantly better for the 28 attended stories than for the 28 unattended stories (67.0±2.5% (standard error) vs. 36.0±1.6%, $t(19) = 10.948$, $p < .001$; the four-choice chance level: 25%). The participants reported a moderate level of attention (8.146±0.343 on a 10-point Likert scale) and attention difficulties (2.039±0.530 on a 10-point Likert scale). The accuracy for the attended story was significantly correlated with both the self-reported attention level ($r = .476$, $p = .043$) and attention difficulty ($r = -.677$, $p = .001$). The self-reported story familiarity level was low for all the participants (0.860±0.220 on a 10-point Likert scale) and was not correlated with comprehension performance ($r = -.224$, $p = .342$). These results suggest that participants' selective attention was effectively manipulated, and the measurement of comprehension performance was reliable. The response accuracy varied from 25.0 % to 51.8% for unattended stories.

The speaker-listener coupling precedes the onset of speech on delta band

A cluster-based permutation (Maris & Oostenveld, 2007) was conducted to reveal the difference between the encoding of the attended and the unattended features and control for multiple comparisons. As figure 2 shows, there was only one inter-brain cluster found in the delta band (cluster-based $p < .001$). The inter-brain cluster had a

wide time range of -4.836 to -0.539 s with the electrodes in the left frontal region. Figure

S1 displays results on other frequency bands.

Figure 2 The attention modulation in speaker-listener coupling. (A) The null distribution of the

cluster-based log(t-statistics) of the inter-brain feature. For illustration purposes, the t-statistics were

transformed into log(t). The label "count" for the x-axis shows the number of occurrences of the

corresponding t-value in the 1000 permutations. The gray lines indicate significant clusters. (B) The

time course and the topo-plot of the significant inter-brain cluster. The light pink line represents the

Encoding r_{att}, and the dark pink represents the Encoding r_{unatt}. The shaded region depicts a significant

difference in the time window. The topo-plot of the average difference between Encoding r_{att} and

Encoding r_{unatt} cluster. The black dots indicate the channels in the cluster.

The attention modulation for acoustic and semantic features occurs after the
speech onset

As shown in Figure 3, the *Acoustic_Early* cluster covered the left-lateralized fronto-

central and occipital electrodes (cluster-based permutation $p < .001$) at a latency of

0.219–0.359 s after the onset of speech. The *Acoustic_Late* cluster had a later latency

of 1.508–1.602 s with the electrodes in the right-frontal regions (cluster-based $p = .005$).

The clusters only appeared on the theta bands.

Figure 3 The attention modulation for the acoustic feature. (A) The null distribution of the cluster-based log(t-statistics) of the acoustic feature. (B) and (C) The time course and the topo-plot of the significant acoustic cluster. The dark blue line represents the Encoding r_{att}, and the light line represents the Encoding r_{unatt}. The shaded region depicts a significant difference in the time window. The topo-plot of the average difference between Encoding r_{att} and Encoding r_{unatt} cluster. The black dots indicate the channels in the cluster.

In contrast to the acoustic feature, semantic clusters were found in the delta band. As Figure 4 indicates, the *Semantic_Early* cluster occurred at 0.227−0.621 s covering the electrodes in frontal and central regions (cluster-based $p = .002$). The *Semantic_Late* cluster was found at 1.073−1.516 s involving the wide distribution of the electrodes (cluster-based $p < .001$).

Figure 4 The attention modulation for the semantic feature. (A) The null distribution of the cluster-based log(t-statistics) of the semantic feature. (B) and (C) The time course and the topo-plot of the significant semantic cluster. The dark brown line represents the Encoding r_{att}, and the light brown represents the Encoding r_{unatt}. The shaded region depicts a significant difference in the time window. The topo-plot of the average difference between Encoding r_{att} and Encoding r_{unatt} cluster. The black dots indicate the channels in the cluster.

The speaker-listener coupling was correlated to the comprehension performance

As Table 2 illustrated, only the inter-brain cluster has a significant partial correlation with the behavioral performance, partial correlation $r = -.769$, $p = .002$ (FDR corrected, Figure 5B). All the other clusters didn't reveal significant correlations. We further analyzed the Encoding r_{att} in the inter-brain cluster and difficulty. We found significant positive correlation, $r = .499$, $p = .025$ as shown in Figure 5C.

As Figure 5A indicates, the average Encoding r_{att} in *Semantic_Early* cluster and

the average Encoding r_{att} in *Acoustic_Early* clusters were highly correlated (r(18)

= .786, $p <$ .001, FDR_corrected). The average Encoding r_{att} in *Semantic_late* cluster

and the average Encoding r_{att} in *Acoustic_Early* cluster were also highly correlated

(r(18) = .565, $p =$ .045, FDR_corrected). There were no other significant correlations

between other clusters ($ps >$ .05).

Figure 5 The correlation between the neural activity and the behavioral result. (A) The pairwise
spearman correlations between Encoding r_{att} in different clusters. The acoustic features and the
semantic features correlated with each other. *A_early* stands for Acoustic_Early, *A_late* stands for
Acoustic_Late, *S_early* is short for Semantic_early, *S_late* stands for Semantic_Late, and *Inter* is short
for Inter-brain. **(B)** and **(C)** The correlation between the Encoding r_{att} in inter-brain clusters and the
accuracy of the comprehension questions and the self-rated difficulty.

Table 2 The partial correlation between the Encoding *r* and accuracy of the
comprehension performance

Clusters	*rho*	*p* values (FDR-corrected)
Acoustic-early	.512	.089
Acoustic-late	-.306	.291
Semantic-early	-.412	.170
Semantic-late	.098	.710
Inter-brain	-.769 **	.002

p < .05, ** *p* < .01 (FDR corrected)

Discussion

Our study directly compares the speaker-listener coupling and the stimulus-listener

coupling in the "cocktail party scene" using the high temporal resolution EEG inter-

brain recording method. Our result uncovered a distinctive pattern of neural alignment,

wherein the listener synchronized with the speaker and speech stimulus in disparate

ways: The listeners aligned to the speaker's neural activity 5 s before the speech onset,

which reveals a significant difference from the speech-listener coupling. Other two

types of couplings occur after the speech onset: The attention modulated the sound

200–350 ms after the speech onset in the theta band. The meaning of speech was

modulated later, in the delta band at 200–600 ms.

Our study proved the distinct role of the attentional speaker-listener coupling in the

"cocktail party": the spatial distribution and the time range are different from the

couplings to the acoustic feature and the semantic feature. For the spatial distribution,

the central electrodes are primarily involved in the acoustic and semantic modulations,

only the left-frontal electrodes are recruited in the inter-brain modulation. The left

frontal regions play a critical region in the language process (Har-shai Yahav & Zion

Golumbic, 2021; Hickok & Poeppel, 2007) and are a 'high-order' area in attention

selection (Zion Golumbic et al., 2013). The left-frontal electrodes may indicate a unique

contribution of IFG when listeners are under adverse listening conditions in previous

inter-brain studies (Dai et al., 2018; Z. Li et al., 2021; L. Liu et al., 2020). For the time

range, the listeners are aligned to the speaker's neural activity 5 s before the speech

onset, where other two features occur after the speech. This result replicated previous

inter-brain studies using fMRI or fNIRS in the high temporal resolution EEG signals

recording method (Dai et al., 2018; J. Jiang et al., 2012; L. Liu et al., 2020; Stephens et

al., 2010). Our result further suggests that the speaker-listener coupling was different

from speech-listener coupling, which highlights the significance of the speaker in the

speech comprehension process (Dai et al., 2018; Pérez et al., 2017, 2019).

The correlation to the behavioral performance further highlights the unique role of

the speaker-listener coupling. In our study, the coupling to the acoustic feature and

semantic feature were correlated, but the coupling to the speaker's neural activity is

independent of the speech stimuli. The speaker-listener coupling had a negative

correlation with the comprehension performance (Figure 5B). We further found that the

neural response was positively correlated with the perceived task difficulty reported by

the listeners (Figure 5C). We successfully replicated the findings of previous inter-brain

studies that indicate a relationship between inter-brain coupling and comprehension

performance (Z. Li et al., 2021, 2022; Stephens et al., 2010). However, in contrast to

these studies, our results reveal a negative correlation. The discrepancy observed may

be attributed to variations in the recording methodology employed: the previous studies

using mainly the blood oxygen signals, like fNRIS and fMRI, but we used a high

temporal resolution EEG signal. The coupling in two types of signals may have

different interpretations. We called the negative correlation observed in our study a

"compensation" mechanism: when the listeners find it hard to complete the task, they

start to guess what the speaker may want to say. However, guessing is not always correct.

Therefore, the comprehension performance is decreasing. While speech itself may only

serve as a trigger and an entrainment signal, the listener could actively "understand"

the speaker not only through the speech itself but rely on the "beyond the stimulus"

grounding (Bashivan et al., 2019; Hartley & Poeppel, 2020; Hasson et al., 2012; Jing

Jiang et al., 2021; Redcay & Schilbach, 2019; Stolk et al., 2016).

Our study reveals the distinct roles of delta and theta bands in attentional

modulation. The theta band modulates with the acoustic feature, and the delta band

modulates with the semantic feature and inter-brain feature. This result possibly extends

our understanding of the functional roles of the two bands. It is consistent with the

previous finding that the theta band processes stimulus-linked features, like the acoustic

feature (Ding et al., 2014; Ding & Simon, 2012a, 2012b; Teng et al., 2017), and the

delta band reflects comprehension-related features, like the semantic feature (Ding et

al., 2014; Etard & Reichenbach, 2019; J. Li et al., 2022; Lu, Jin, Pan, et al., 2022). In

particular, the modulation for the inter-brain feature was also found in the delta band,

which may indicate that the delta bands reflect the comprehension-related functions.

Our study introduces novel analytical methods to the field of inter-brain studies.

We applied the temporal response function method (TRF) to depict the listener's neural

response to a high dimensional speaker's EEG signal. This method was widely used in

the studies of naturalistic speech (Crosse et al., 2016; Di Liberto et al., 2015; Ding &

Simon, 2012a, 2012b; O'Sullivan et al., 2015; Power et al., 2012), as well as the visual

studies (Jia et al., 2017, 2019) and memory studies (Huang et al., 2018). Using the TRF

method enables a direct comparison between the speaker-listener coupling and different

types of speech-listener coupling. We also used the NLP models (Armeni et al., 2019;

Brodbeck et al., 2022; Broderick et al., 2018; Kingma & Ba, 2015) to describe the

semantic meaning in the speech as the latest review suggested (Jing Jiang et al., 2021;

Pérez & Davis, 2023). The combined use of different lately methods enhance our

understanding of the attention modulation in a comprehensive way and provide a new

inspiration for the future studies underlying inter-brain coupling.

While this study sheds light on a new perspective in the "cocktail party effect",

there are several areas that warrant further exploration. Firstly, our study simplified the

semantic feature into the "surprisal index", which is a common choice in previous

studies(Aurnhammer & Frank, 2019; Brodbeck et al., 2022; Heilbron et al., 2022;

Mesik et al., 2021; Willems et al., 2016). However, there were alternative approaches

to extract the semantic feature in the text, such as the word embeddings. Subsequent

studies could undertake a comparative analysis of various semantic representations,

such as surprisal, entropy, or word embeddings, generated by diverse models including

word2vec (Broderick et al., 2018; Lu, Jin, Pan, et al., 2022; Mikolov et al., 2013), GPT

(Heilbron et al., 2022; Schramowski et al., 2022; Solaiman et al., 2019) or BERT

(Devlin et al., 2019). This would enable a comprehensive evaluation of the consistency

of their findings. Meanwhile, our studies used a sequential-dual-brain approach to use

the speaker-listener coupling. While this approach has the potential to reduce variance

originating from the speaker's neural activity (Leong et al., 2017; Z. Li et al., 2021;

Redcay & Schilbach, 2019), it may not fully capture the complexity of neural activity

present in real-life conversations (Dai et al., 2018; Jing Jiang et al., 2012; Pérez et al.,

2017). Further studies could also apply the simultaneous dual-brain approach (Redcay

& Schilbach, 2019) to further investigate the issue.

In conclusion, our study used the attention as a spotlight and revealed that the

listener would strike the same "neural notes" with the speaker before the speech onset,

while the acoustic note is struck on the theta bands after the speech onset firstly, and

the semantic note comes later and lasts longer on the delta band. Our study depicts the

temporal dynamics of the attention modulation and the functional roles of different

frequency bands, which contributes to the old "cocktail party" a new integrative

perspective. In an era of rising artificial intelligence, our study still highlights the value

and significance of human-to-human interaction between speakers and listeners, as the

result demonstrates the possibility of mutual understanding between the speaker and

listener "beyond the stimulus" in their neural activities (Yeshurun et al., 2021).

Acknowledgments

This work was supported by the National Science Foundation of China (NSFC) and the German Research Foundation (DFG) in project Crossmodal Learning (grant number: NSFC 62061136001/DFG TRR-169/C1, B1), the National Natural Science Foundation of China (grant number: 61977041) and a postdoctoral fellowship of the Humboldt Foundation to J.L.

The authors would like to thank Prof. Dr. Xiaoqin Wang and Dr. Yue Ding for providing the shielded room for the experiment as well as necessary technical support. The authors would like to express their gratitude towards Dr. Zhiyuan Liu and members of his lab for computing the NLP models.

Reference

Armeni, K., Willems, R. M., van den Bosch, A., & Schoffelen, J.-M. (2019). Frequency-specific brain dynamics related to prediction during language comprehension. *NeuroImage*, *198*, 283–295. https://doi.org/10.1016/j.neuroimage.2019.04.083

Arnal, L. H., Wyart, V., & Giraud, A. L. (2011). Transitions in neural oscillations reflect prediction errors generated in audiovisual speech. *Nature Neuroscience*, *14*(6), 797–801. https://doi.org/10.1038/nn.2810

Aurnhammer, C., & Frank, S. L. (2019). Evaluating information-theoretic measures of word prediction in naturalistic sentence reading. *Neuropsychologia*, *134*(August), 107198. https://doi.org/10.1016/j.neuropsychologia.2019.107198

Bai, F., Meyer, A. S., & Martin, A. E. (2022). Neural dynamics differentially encode phrases and sentences during spoken language comprehension. In *PLoS Biology* (Vol. 20, Issue 7). https://doi.org/10.1371/journal.pbio.3001713

Bashivan, P., Kar, K., & DiCarlo, J. J. (2019). Neural population control via deep image synthesis. *Science*, *364*(6439). https://doi.org/10.1126/science.aav9436

Bengio, Y., Ducharme, R., & Vincent, P. (2003). A neural probabilistic language model. *Journal of Machine Learning Research*, *3*, 1137–1155.

Brainard, D. H., & Brainard, D. H. (1997). The Psychophysics Toolbox. In *Spatial vision* (pp. 433–436).

Brodbeck, C., Bhattasali, S., Cruz Heredia, A. A. L., Resnik, P., Simon, J. Z., & Lau, E. (2022). Parallel processing in speech perception with local and global representations of linguistic context. *ELife*, *11*, 1–28. https://doi.org/10.7554/eLife.72056

Broderick, M. P., Anderson, A. J., Di Liberto, G. M., Crosse, M. J., & Lalor, E. C. (2018). Electrophysiological Correlates of Semantic Dissimilarity Reflect the Comprehension of Natural, Narrative Speech. *Current Biology*, *28*(5), 803-809.e3. https://doi.org/10.1016/j.cub.2018.01.080

Broderick, M. P., Anderson, A. J., & Lalor, E. C. (2019). Semantic Context Enhances the Early Auditory Encoding of Natural Speech. *The Journal of Neuroscience*, *39*(38), 7564–7575. https://doi.org/10.1523/JNEUROSCI.0584-19.2019

Broderick, M. P., Di Liberto, G. M., Anderson, A. J., Rofes, A., & Lalor, E. C. (2021). Dissociable electrophysiological measures of natural language processing reveal differences in speech comprehension strategy in healthy ageing. *Scientific Reports*, *11*(1), 1–12. https://doi.org/10.1038/s41598-021-84597-9

Cherry, E. C. (1953). Some experiments on the recognition of speech, with one and with two ears. *The Journal of the Acoustical Society of America*, *25*(5), 975–979.

Chomsky, N. (1959). Verbal behavior by B. F. Skinner. *Language*, *35*(1), 26–58.

Connolly, J. F., Stewart, S. H., & Phillips, N. A. (1990). The effects of processing
requirements on neurophysiological responses to spoken sentences. *Brain and*
Language, *39*(2), 302–318.

Crosse, M. J., Di Liberto, G. M., Bednar, A., & Lalor, E. C. (2016). The Multivariate
Temporal Response Function (mTRF) Toolbox: A MATLAB Toolbox for
Relating Neural Signals to Continuous Stimuli. *Frontiers in Human*
Neuroscience, *10*, 604. https://doi.org/10.3389/fnhum.2016.00604

Crosse, M. J., Zuk, N. J., Di Liberto, G. M., Nidiffer, A. R., Molholm, S., & Lalor, E.
C. (2021). Linear Modeling of Neurophysiological Responses to Speech and
Other Continuous Stimuli: Methodological Considerations for Applied Research.
Frontiers in Neuroscience, *15*. https://doi.org/10.3389/fnins.2021.705621

Dai, B., Chen, C., Long, Y., Zheng, L., Zhao, H., Bai, X., Liu, W., Zhang, Y., Liu, L.,
Guo, T., Ding, G., & Lu, C. (2018). Neural mechanisms for selectively tuning in
to the target speaker in a naturalistic noisy situation. *Nature Communications*,
9(1), 1–12. https://doi.org/10.1038/s41467-018-04819-z

Dai, B., McQueen, J. M., Terporten, R., Hagoort, P., & Kösem, A. (2022). Distracting
linguistic information impairs neural tracking of attended speech. *Current*
Research in Neurobiology, *3*, 100043.
https://doi.org/10.1016/j.crneur.2022.100043

de Cheveigné, A., & Nelken, I. (2019). Filters: When, Why, and How (Not) to Use
Them. *Neuron*, *102*(2), 280–293. https://doi.org/10.1016/j.neuron.2019.02.039

Devlin, J., Chang, M. W., Lee, K., & Toutanova, K. (2019). BERT: Pre-training of
deep bidirectional transformers for language understanding. *NAACL HLT 2019 -*
2019 Conference of the North American Chapter of the Association for
Computational Linguistics: Human Language Technologies - Proceedings of the
Conference, *1*(Mlm), 4171–4186.

Di Liberto, G. M., O'Sullivan, J. A., & Lalor, E. C. (2015). Low-Frequency Cortical
Entrainment to Speech Reflects Phoneme-Level Processing. *Current Biology*,
25(19), 2457–2465. https://doi.org/10.1016/j.cub.2015.08.030

Ding, N., Chatterjee, M., & Simon, J. Z. (2014). Robust cortical entrainment to the
speech envelope relies on the spectro-temporal fine structure. *NeuroImage*, *88*,
41–46. https://doi.org/10.1016/j.neuroimage.2013.10.054

Ding, N., & Simon, J. Z. (2012a). Emergence of neural encoding of auditory objects
while listening to competing speakers. *Proceedings of the National Academy of*
Sciences of the United States of America, *109*(29), 11854–11859.
https://doi.org/10.1073/pnas.1205381109

Ding, N., & Simon, J. Z. (2012b). Neural coding of continuous speech in auditory
cortex during monaural and dichotic listening. *Journal of Neurophysiology*,
107(1), 78–89. https://doi.org/10.1152/jn.00297.2011

Etard, O., & Reichenbach, T. (2019). Neural Speech Tracking in the Theta and in the
Delta Frequency Band Differentially Encode Clarity and Comprehension of
Speech in Noise. *The Journal of Neuroscience*, *39*(29), 5750–5759.
https://doi.org/10.1523/JNEUROSCI.1828-18.2019
Gadamer, H. G. (1975). *Truth and Method*. Seabury Press.
https://books.google.co.jp/books?id=zQnXAAAAMAAJ
Hamilton, L. S., & Huth, A. G. (2018). The revolution will not be controlled: natural
stimuli in speech neuroscience. *Language, Cognition and Neuroscience*, 1–10.
https://doi.org/10.1080/23273798.2018.1499946
Har-shai Yahav, P., & Zion Golumbic, E. (2021). Linguistic processing of task-
irrelevant speech at a cocktail party. *ELife*, *10*, 1–24.
https://doi.org/10.7554/eLife.65096
Hartley, C. A., & Poeppel, D. (2020). Beyond the Stimulus: A Neurohumanities
Approach to Language, Music, and Emotion. *Neuron*, *108*(4), 597–599.
https://doi.org/10.1016/j.neuron.2020.10.021
Hasson, U., Ghazanfar, A. A., Galantucci, B., Garrod, S., & Keysers, C. (2012).
Brain-to-brain coupling: A mechanism for creating and sharing a social world.
Trends in Cognitive Sciences, *16*(2), 114–121.
https://doi.org/10.1016/j.tics.2011.12.007
Heil, M., Rolke, B., Pecchinenda, A., Heil, M., Rolke, B., & Pecchinenda, A. (2004).
Automatic Semantic Activation Is No Myth. *Psychological Science*, *15*(12),
852–857. https://doi.org/doi.org/10.1111
Heilbron, M., Armeni, K., Schoffelen, J.-M., Hagoort, P., & de Lange, F. P. (2022). A
hierarchy of linguistic predictions during natural language comprehension.
Proceedings of the National Academy of Sciences, *119*(32), 2020.12.03.410399.
https://doi.org/10.1073/pnas.2201968119
Henry, M. J., & Obleser, J. (2012). Frequency modulation entrains slow neural
oscillations and optimizes human listening behavior. *Proceedings of the National*
Academy of Sciences, *109*(49), 20095–20100.
https://doi.org/10.1073/pnas.1213390109
Hickok, G., & Poeppel, D. (2007). The cortical organization of speech processing.
Nature Reviews Neuroscience, *8*(5), 393–402. https://doi.org/10.1038/nrn2113
Hillyard, S. A., Hink, R. F., Schwent, V. L., & Picton, T. W. (1973). Electrical signs
of selective attention in the human brain. *Science*, *182*(4108), 177–180.
https://doi.org/10.1126/science.182.4108.177
Huang, Q., Jia, J., Han, Q., & Luo, H. (2018). Fast-backward replay of sequentially
memorized items in humans. *ELife*, *7*, e35164.
https://doi.org/10.7554/eLife.35164
707

Jia, J., Fang, F., & Luo, H. (2019). Selective spatial attention involves two alpha-band
components associated with distinct spatiotemporal and functional
characteristics. *NeuroImage*, *199*, 228–236.
https://doi.org/10.1016/j.neuroimage.2019.05.079

Jia, J., Liu, L., Fang, F., & Luo, H. (2017). Sequential sampling of visual objects
during sustained attention. *PLoS Biology*, *15*(6), 1–19.
https://doi.org/10.1371/journal.pbio.2001903

Jiang, J., Dai, B., Peng, D., Zhu, C., Liu, L., & Lu, C. (2012). Neural Synchronization
during Face-to-Face Communication. *Journal of Neuroscience*, *32*(45), 16064–
16069. https://doi.org/10.1523/JNEUROSCI.2926-12.2012

Jiang, Jing, Chen, C., Dai, B., Shi, G., Ding, G., Liu, L., & Lu, C. (2015). Leader
emergence through interpersonal neural synchronization. *Proceedings of the*
National Academy of Sciences, *112*(14), 4274–4279.
https://doi.org/10.1073/pnas.1422930112

Jiang, Jing, Dai, B., Peng, D., Zhu, C., Liu, L., & Lu, C. (2012). Neural
synchronization during face-to-face communication. *Journal of Neuroscience*,
32(45), 16064–16069. https://doi.org/10.1523/JNEUROSCI.2926-12.2012

Jiang, Jing, Zheng, L., & Lu, C. (2021). A hierarchical model for interpersonal verbal
communication. *Social Cognitive and Affective Neuroscience*, *16*(1–10), 246–
255. https://doi.org/10.1093/scan/nsaa151

Kingma, D. P., & Ba, J. L. (2015). Adam: A method for stochastic optimization. *3rd*
International Conference on Learning Representations, ICLR 2015 - Conference
Track Proceedings, 1–15.

Kuhlen, A. K., Allefeld, C., & Haynes, J.-D. (2012). Content-specific coordination of
listeners' to speakers' EEG during communication. *Frontiers in Human*
Neuroscience, *6*(October), 1–15. https://doi.org/10.3389/fnhum.2012.00266

Leong, V., Byrne, E., Clackson, K., Georgieva, S., Lam, S., & Wass, S. (2017).
Speaker gaze increases information coupling between infant and adult brains.
Proceedings of the National Academy of Sciences of the United States of
America, *114*(50), 13290–13295. https://doi.org/10.1073/pnas.1702493114

Li, J., Hong, B., Nolte, G., Engel, A. K., & Zhang, D. (2022). Preparatory delta phase
response is correlated with naturalistic speech comprehension performance.
Cognitive Neurodynamics, *16*(2), 337–352. https://doi.org/10.1007/s11571-021-
09711-z

Li, Z., Hong, B., Wang, D., Nolte, G., Engel, A. K., & Zhang, D. (2022). Speaker-
listener neural coupling reveals a right-lateralized mechanism for non-native
speech-in-noise comprehension. *Cerebral Cortex*, 1–14.
https://doi.org/10.1093/cercor/bhac302

Li, Z., Li, J., Hong, B., Nolte, G., Engel, A. K., & Zhang, D. (2021). Speaker–Listener
Neural Coupling Reveals an Adaptive Mechanism for Speech Comprehension in
a Noisy Environment. *Cerebral Cortex*, *31*(10), 4719–4729.
https://doi.org/10.1093/cercor/bhab118
Liu, L., Zhang, Y., Zhou, Q., Garrett, D. D., Lu, C., Chen, A., Qiu, J., & Ding, G.
(2020). Auditory-Articulatory Neural Alignment between Listener and Speaker
during Verbal Communication. *Cerebral Cortex*, *30*(3), 942–951.
https://doi.org/10.1093/cercor/bhz138
Liu, Y., Piazza, E. A., Simony, E., Shewokis, P. A., Onaral, B., Hasson, U., & Ayaz,
H. (2017). Measuring speaker–listener neural coupling with functional near
infrared spectroscopy. *Scientific Reports*, *7*(1), 43293.
https://doi.org/10.1038/srep43293
Lu, Y., Jin, P., Ding, N., & Tian, X. (2022). Delta-band neural tracking primarily
reflects rule-based chunking instead of semantic relatedness between words.
Cerebral Cortex, 1–11. https://doi.org/10.1093/cercor/bhac354
Lu, Y., Jin, P., Pan, X., & Ding, N. (2022). Delta-band neural activity primarily tracks
sentences instead of semantic properties of words. *NeuroImage*, *251*(January),
118979. https://doi.org/10.1016/j.neuroimage.2022.118979
Maris, E., & Oostenveld, R. (2007). Nonparametric statistical testing of EEG- and
MEG-data. *Journal of Neuroscience Methods*, *164*(1), 177–190.
https://doi.org/10.1016/j.jneumeth.2007.03.024
McDermott, J. H. (2009). The cocktail party problem. *Current Biology*, *19*(22), 1024–
1027. https://doi.org/10.1016/j.cub.2009.09.005
Mesik, J., Ray, L., & Wojtczak, M. (2021). Effects of Age on Cortical Tracking of
Word-Level Features of Continuous Competing Speech. *Frontiers in*
Neuroscience, *15*(April), 1–21. https://doi.org/10.3389/fnins.2021.635126
Middlebrooks, J. C., Simon, J. Z., Popper, A. N., & Fay, R. R. (2017). The Auditory
System at the Cocktail Party. In J. C. Middlebrooks, J. Z. Simon, A. N. Popper,
& R. R. Fay (Eds.), *Springer International Publishing*. Springer International
Publishing. https://doi.org/10.1007/978-3-319-51662-2
Mikolov, T., Chen, K., Corrado, G., & Dean, J. (2013). Efficient Estimation ofWord
Representations in Vector Space Tomas. *1st International Conference on*
Learning Representations, ICLR 2013 - Workshop Track Proceedings, 1–12.
Mirkovic, B., Debener, S., Jaeger, M., & De Vos, M. (2015). Decoding the attended
speech stream with multi-channel EEG: implications for online, daily-life
applications. *Journal of Neural Engineering*, *12*(4), 046007.
https://doi.org/10.1088/1741-2560/12/4/046007
784
785

Nastase, S. A., Liu, Y. F., Hillman, H., Zadbood, A., Hasenfratz, L., Keshavarzian,
N., Chen, J., Honey, C. J., Yeshurun, Y., Regev, M., Nguyen, M., Chang, C. H.
C., Baldassano, C., Lositsky, O., Simony, E., Chow, M. A., Leong, Y. C.,
Brooks, P. P., Micciche, E., … Hasson, U. (2021). The "Narratives" fMRI
dataset for evaluating models of naturalistic language comprehension. *Scientific*
Data, *8*(1), 1–22. https://doi.org/10.1038/s41597-021-01033-3
O'Sullivan, J. A., Power, A. J., Mesgarani, N., Rajaram, S., Foxe, J. J., Shinn-
Cunningham, B. G., Slaney, M., Shamma, S. A., & Lalor, E. C. (2015).
Attentional Selection in a Cocktail Party Environment Can Be Decoded from
Single-Trial EEG. *Cerebral Cortex*, *25*(7), 1697–1706.
https://doi.org/10.1093/cercor/bht355
Pérez, A., Carreiras, M., & Duñabeitia, J. A. (2017). Brain-To-brain entrainment:
EEG interbrain synchronization while speaking and listening. *Scientific Reports*,
7(1), 1–12. https://doi.org/10.1038/s41598-017-04464-4
Pérez, A., & Davis, M. H. (2023). Speaking and listening to inter-brain relationships.
Cortex, *159*, 54–63. https://doi.org/10.1016/j.cortex.2022.12.002
Pérez, A., Dumas, G., Karadag, M., & Duñabeitia, J. A. (2019). Differential brain-to-
brain entrainment while speaking and listening in native and foreign languages.
Cortex, *111*, 303–315. https://doi.org/10.1016/j.cortex.2018.11.026
Power, A. J., Foxe, J. J., Forde, E. J., Reilly, R. B., & Lalor, E. C. (2012). At what
time is the cocktail party? A late locus of selective attention to natural speech.
European Journal of Neuroscience, *35*(9), 1497–1503.
https://doi.org/10.1111/j.1460-9568.2012.08060.x
Power, A. J., Lalor, E. C., & Reilly, R. B. (2011). Endogenous auditory spatial
attention modulates obligatory sensory activity in auditory cortex. *Cerebral*
Cortex, *21*(6), 1223–1230. https://doi.org/10.1093/cercor/bhq233
Redcay, E., & Schilbach, L. (2019). Using second-person neuroscience to elucidate
the mechanisms of social interaction. *Nature Reviews Neuroscience*, *20*(8), 495–
505. https://doi.org/10.1038/s41583-019-0179-4
Schoot, L., Hagoort, P., & Segaert, K. (2016). What can we learn from a two-brain
approach to verbal interaction? *Neuroscience and Biobehavioral Reviews*,
68(December), 454–459. https://doi.org/10.1016/j.neubiorev.2016.06.009
Schramowski, P., Turan, C., Andersen, N., Rothkopf, C. A., & Kersting, K. (2022).
Large pre-trained language models contain human-like biases of what is right
and wrong to do. *Nature Machine Intelligence*, *4*(3), 258–268.
https://doi.org/10.1038/s42256-022-00458-8
822
823
824

825 Schrimpf, M., Blank, I. A., Tuckute, G., Kauf, C., Hosseini, E. A., Kanwisher, N.,
826   Tenenbaum, J. B., & Fedorenko, E. (2021). The neural architecture of language:
827   Integrative modeling converges on predictive processing. *Proceedings of the*
828   *National Academy of Sciences*, *118*(45), e2105646118.
829   https://doi.org/10.1073/pnas.2105646118

830 Searle, J. R. (1980). Minds, brains, and programs. In *the MIT press* (Vol. 3).
831   https://doi.org/10.1.1.83.5248

832 Shinn-Cunningham, B. G. (2008). Object-based auditory and visual attention. *Trends*
833   *in Cognitive Sciences*, *12*(5), 182–186. https://doi.org/10.1016/j.tics.2008.02.003

834 Skinner, B. F. (1957). Verbal behavior / B. F. Skinner. In *Verbal behavior*. Appleton-
835   Century-Crofts.

836 Skinner, B. F. (1986). Signs Through. *Evolution*, *1*(1), 115–122.

837 Solaiman, I., Brundage, M., Clark, J., Askell, A., Herbert-Voss, A., Wu, J., Radford,
838   A., & Wang, J. (2019). *Release Strategies and the Social Impacts of Language*
839   *Models*. http://arxiv.org/abs/1908.09203

840 Sonkusare, S., Breakspear, M., & Guo, C. (2019). Naturalistic Stimuli in
841   Neuroscience: Critically Acclaimed. *Trends in Cognitive Sciences*, *23*(8), 699–
842   714. https://doi.org/10.1016/j.tics.2019.05.004

843 Stephens, G. J., Silbert, L. J., & Hasson, U. (2010). Speaker–listener neural coupling
844   underlies successful communication. *Proceedings of the National Academy of*
845   *Sciences*, *107*(32), 14425–14430. https://doi.org/10.1073/pnas.1008662107

846 Stolk, A., Verhagen, L., & Toni, I. (2016). Conceptual Alignment: How Brains
847   Achieve Mutual Understanding. *Trends in Cognitive Sciences*, *20*(3), 180–191.
848   https://doi.org/10.1016/j.tics.2015.11.007

849 Teng, X., Tian, X., Rowland, J., & Poeppel, D. (2017). Concurrent temporal channels
850   for auditory processing: Oscillatory neural entrainment reveals segregation of
851   function at different scales. *PLoS Biology*, *15*(11), 1–29.
852   https://doi.org/10.1371/journal.pbio.2000812

853 Teoh, E. S., Ahmed, F., & Lalor, E. C. (2022). Attention Differentially Affects
854   Acoustic and Phonetic Feature Encoding in a Multispeaker Environment. *The*
855   *Journal of Neuroscience*, *42*(4), 682–691.
856   https://doi.org/10.1523/JNEUROSCI.1455-20.2021

857 Vaswani, A., Shazeer, N., Parmar, N., Uszkoreit, J., Jones, L., Gomez, A. N., Kaiser,
858   Ł., & Polosukhin, I. (2017). Attention is all you need. *Advances in Neural*
859   *Information Processing Systems*, *2017-Decem*(Nips), 5999–6009.

860
861
862

Willems, R. M., Frank, S. L., Nijhof, A. D., Hagoort, P., & Van Den Bosch, A. (2016). Prediction during Natural Language Comprehension. *Cerebral Cortex*, *26*(6), 2506–2516. https://doi.org/10.1093/cercor/bhv075

Willems, R. M., Nastase, S. A., & Milivojevic, B. (2020). Narratives for Neuroscience. *Trends in Neurosciences*, *43*(5), 271–273. https://doi.org/10.1016/j.tins.2020.03.003

Yeshurun, Y., Nguyen, M., & Hasson, U. (2021). The default mode network: where the idiosyncratic self meets the shared social world. *Nature Reviews Neuroscience*, *22*(3), 181–192. https://doi.org/10.1038/s41583-020-00420-w

Yu, W., Sun, W., & Ding, N. (2022). Asymmetrical cross-modal influence on neural encoding of auditory and visual features in natural scenes. *NeuroImage*, *255*, 119182. https://doi.org/10.1016/j.neuroimage.2022.119182

Zhang, Z. G., Hu, L., Hung, Y. S., Mouraux, A., & Iannetti, G. D. (2012). Gamma-band oscillations in the primary somatosensory cortex-A direct and obligatory correlate of subjective pain intensity. *Journal of Neuroscience*, *32*(22), 7429–7438. https://doi.org/10.1523/JNEUROSCI.5877-11.2012

Zheng, L., Chen, C., Liu, W., Long, Y., Zhao, H., Bai, X., Zhang, Z., Han, Z., Liu, L., Guo, T., Chen, B., Ding, G., Lu, C., & Chunming Lu, C. (2018). *Enhancement of teaching outcome through neural prediction of the students' knowledge state*. *January*, 1–12. https://doi.org/10.1002/hbm.24059

Zuk, N. J., Murphy, J. W., Reilly, R. B., & Lalor, E. C. (2021). Envelope reconstruction of speech and music highlights stronger tracking of speech at low frequencies. *PLoS Computational Biology*, *17*(9), 1–32. https://doi.org/10.1371/journal.pcbi.1009358