Behavioral/Cognitive

Cortical Tracking of a Background Speaker Modulates the Comprehension of a Foreground Speech Signal

Mahmoud Keshavarzi, Enrico Varano, and Tobias Reichenbach
Department of Bioengineering and Centre for Neurotechnology, Imperial College London, South Kensington Campus, London, SW7 2AZ, England

Understanding speech in background noise is a difficult task. The tracking of speech rhythms such as the rate of syllables and words by cortical activity has emerged as a key neural mechanism for speech-in-noise comprehension. In particular, recent investigations have used transcranial alternating current stimulation (tACS) with the envelope of a speech signal to influence the cortical speech tracking, demonstrating that this type of stimulation modulates comprehension and therefore providing evidence of a functional role of the cortical tracking in speech processing. Cortical activity has been found to track the rhythms of a background speaker as well, but the functional significance of this neural response remains unclear. Here we use a speech-comprehension task with a target speaker in the presence of a distractor voice to show that tACS with the speech envelope of the target voice as well as tACS with the envelope of the distractor speaker both modulate the comprehension of the target speech. Because the envelope of the distractor speech does not carry information about the target speech stream, the modulation of speech comprehension through tACS with this envelope provides evidence that the cortical tracking of the background speaker affects the comprehension of the foreground speech signal. The phase dependency of the resulting modulation of speech comprehension is, however, opposite to that obtained from tACS with the envelope of the target speech signal. This suggests that the cortical tracking of the ignored speech stream and that of the attended speech stream may compete for neural resources.

Key words: auditory; background speech; cortical tracking; selective attention; tACS

Significance Statement
Loud environments such as busy pubs or restaurants can make conversation difficult. However, they also allow us to eavesdrop into other conversations that occur in the background. In particular, we often notice when somebody else mentions our name, even if we have not been listening to that person. However, the neural mechanisms by which background speech is processed remain poorly understood. Here we use transcranial alternating current stimulation, a technique through which neural activity in the cerebral cortex can be influenced, to show that cortical responses to rhythms in the distractor speech modulate the comprehension of the target speaker. Our results provide evidence that the cortical tracking of background speech rhythms plays a functional role in speech processing.

Introduction
Speech is a fascinatingly complex signal, whose processing requires analysis of individual phonemes, syllables, and words to extract meaning (Ingram, 2007; Poeppel et al., 2012; Hickok and Small, 2015). Moreover, when confronted with background noise such as other people talking, traffic or music, our brain must first segregate the target speech signal from the background acoustics before the processing of the target speech stream can begin (Bregman, 1994; Snyder and Alain, 2007).

An ignored speech signal can, however, still affect behavior. Hearing one’s own name in the background can, for instance, shift our attention to the corresponding speaker (Moray, 1959; Wood and Cowan, 1995). Moreover, when attempting to listen to a target speaker in the presence of a distractor voice, a listener occasionally understands words from the distractor speaker (Brungart, 2001). A background speech signal in a listener’s native language is accordingly more distracting than speech in a foreign language or nonspeech sounds such as music or traffic (Brungart, 2001; Cooke et al., 2008; Cooke and Lu, 2010). However, the neural encoding of a background speech signal remains poorly understood.
An important aspect of speech processing in a computer is the parsing of the speech stream into functional constituents such as syllables and words. The brain presumably employs a similar parsing. Cortical activity, in particular in the δ (1–4 Hz) and theta (4–8 Hz) frequency ranges, has indeed been found to track speech rhythms, such as those set by the rate of syllables and words (Giraud and Poeppel, 2012; Ding and Simon, 2014; Di Liberto et al., 2015). A computational model shows that such tracking can yield effective online syllable parsing (Hyalì et al., 2015).

Cortical tracking of speech rhythms may contribute to the processing of both an attended and an ignored speech signal. When selectively attending to one of two competing speakers, cortical activity tracks the rhythms of the attended speech as well as those of the ignored voice, although the cortical tracking of the attended speaker is stronger than that of the unattended speaker (Ding and Simon, 2012; Mesgarani and Chang, 2012; Horton et al., 2013). Moreover, the cortical tracking of an attended talker is modulated by speech comprehension, such as when speech is noise-vocoded, presented in different levels of background noise or compared between a native and foreign language (Ding et al., 2014; Brodbeck et al., 2018; Broderick et al., 2018; Vanthornhout et al., 2018; Etard and Reichenbach, 2019; Weisbart et al., 2020).

The modulation of the cortical speech tracking by cognitive factors such as attention and comprehension are not merely side effects of other neural processes. Instead, a functional role of the cortical tracking of an attended speaker in speech processing has been shown through transcranial alternating current stimulation (tACS), a noninvasive technique to influence cortical activity in the frequency range of the stimulation (Antal and Paulus, 2013; Helfrich et al., 2014). Applying tACS with waveforms that were derived from the speech envelope, reflecting the speech rhythms, has demonstrated an effect on the cortical speech tracking (Zoefel et al., 2018). Moreover, this type of tACS has been found to modulate the comprehension of speech in noise, as well as the comprehension of noise-vocoded speech (Riecke et al., 2018; Wilsch et al., 2018; Kadir et al., 2020; Keshavarzi et al., 2020).

However, it remains unclear what functional role the cortical tracking of an ignored speech signal has, and whether it can influence behavior. Here, we therefore set out to investigate the impact of tACS with the speech envelope of a distractor voice on speech comprehension.

Materials and Methods

Experimental design and statistical analysis. To investigate the functional role of the cortical tracking of a background speech signal, we used a simplified version of a cocktail party with two voices, male and female (Fig. 1A,B). Subjects were instructed to listen to a particular target speaker and to ignore the other distractor voice. Their auditory cortices were stimulated simultaneously and noninvasively with alternating current (Fig. 1A). The current waveforms were derived from the speech envelope of either the target or the distractor speech signal (Fig. 1B).

Most of the power spectral density of the speech envelope is contained in two frequency bands, the δ band (1–4 Hz) and the theta band (4–8 Hz). Previously we showed that the modulation of speech comprehension results from stimulation with the theta frequency portion but not from the δ frequency portion of the speech envelope (Keshavarzi et al., 2020) and that speech comprehension can be enhanced when the current has no temporal delay with respect to the speech stream (Keshavarzi and Reichenbach, 2020). In this study, we accordingly used current waveforms that represented the theta-band portion of the speech envelope and that were temporally aligned to the acoustic signal (Fig. 1C).

We determined the modulation of speech comprehension through tACS by shifting the current waveforms by four different phases: 0°, 90°, 180°, and 270°. We previously showed that tACS with the theta-band portion of the envelope of a target speech then leads to a modulation of speech comprehension that depends on the phase shifts in a cyclical manner (Keshavarzi et al., 2020).

To assess the potential dependence of the speech comprehension scores CS(φn) on the phase shifts φn with n = 0, 1, 2, 3, we first determined, for each subject, the mean comprehension score $\overline{CS}$ across the different phase shifts, $\overline{CS} = \frac{1}{4} \sum_{k=0}^{3} CS(\phi_k)$. We then focused on the differences of the speech comprehension scores $\Delta CS(\phi_n)$ of each subject from the mean value, $\Delta CS(\phi_n) = CS(\phi_n) - \overline{CS}$. Because of the cyclical nature of the phase shifts, the dependence of this change in the speech comprehension score on the phase shifts $\phi_n$ could be expressed through the discrete Fourier transform as follows:

$$\Delta CS(\phi_n) = \sum_{k=0}^{3} a_k e^{i k \pi \phi_n / 180°},$$

in which the complex coefficients $a_k$ were calculated as $a_k = \frac{1}{4} \sum_{n=0}^{3} \Delta CS(\phi_n) e^{-i k \pi \phi_n / 180°}$. Because the speech comprehension...
scores were real, it followed that the coefficients $a_0$ and $a_2$ were real as well and that $a_1 = a'_1$. Equation 1 for the dependence of $\Delta CS(\phi)$ on the phases $\phi$, could thus be rewritten as follows:

$$\Delta CS(\phi) = A_0 + A_1 \cos(\phi - \varphi_1) + A_2 \cos(2\phi),$$

(2)

in which $A_0 = a_0$ was a constant offset; $A_1 = 2(a_1 \sin \varphi_1)$ and $A_2 = a_2$, respectively, denoted the amplitude of the variation at the periods of 360° and 180°; and $\varphi_1 = \arg(a_1)$ was the phase offset at the first period.

To fit Equation 2 to the data, we first rewrote the middle term on the right-hand side, $A_1 \cos(\phi - \varphi_1)$, as a sum of a sine and a cosine function of $\phi$, to obtain the following:

$$\Delta CS(\phi) = A_0 + B_1 \sin(\phi) + B_2 \cos(\phi) + A_2 \cos(2\phi),$$

(3)

with the coefficients $B_1 = A_1 \cos(\varphi_1)$ and $B_2 = A_1 \sin(\varphi_1)$. We then determined the parameters $A_0$, $B_1$, $B_2$, and $A_2$ through multiple linear regression. The corresponding $p$ values were corrected for multiple comparisons using the false discovery rate (FDR) correction. The amplitude $A_1$ of the variation at the period of 360° followed as $A_1 = \sqrt{B_1^2 + B_2^2}$, and the corresponding phase offset as $\varphi_1 = \arctan(B_2/B_1)$.

Because the data may contain outliers, we also used robust regression to determine the parameters $A_0$, $B_1$, $B_2$, and $A_2$ in Equation 3 (Holland and Welsch, 1977). Robust regression employs an iteratively reweighted least-squares algorithm that assigns weights to each data point. Outlier data points thereby obtain a lower weight so that they contribute less to the parameter estimation.

To determine if the robust regression yielded a better fit than the standard regression, we used bootstrapping with 10,000 samples. For each sample, we performed both the standard regression as well as the robust regression that started at the beginning of the target speech and lasted 500 ms. Smooth onsets and offsets were implemented through ramps with a duration of 100 ms. Four further waveforms were computed from the envelope of the target speech by bandpass filtering this signal in the theta frequency band (zero-phase IIR filter, low cutoff of 4 Hz, high cutoff of 8 Hz, order 6). The bandpass filtered envelopes were further processed so that all maxima and minima occurred at equal values and were then shifted by four different phases (0°, 90°, 180°, 270°). The four remaining types of tACS waveforms were computed from the envelope of the distractor speech; these computations were analogous to the computation of the waveforms from the envelope of the target speech.

**Experimental procedure.** Participants were seated in a soundproof room. Two rubber electrodes (35 cm²) covered by sponge pads and wetted with a 0.9% saline solution (5 ml per electrode) were then placed on the T7 and T8 locations, and two further ones were placed to the left and right of the Cz location of the participant’s head (International 10–10 system of electrode placement). The electrode placed on T7 and the one to the left of Cz were connected to one neurostimulation device and the two remaining ones to another device. The electrodes at the T7 and T8 locations functioned as the anodes and the ones at Cz as the cathodes. The distance between electrodes of each device was set to be almost equal and below 10 kΩ.

To determine the maximum magnitude of current stimulation for each subject, a 5 s sinusoidal signal with a frequency of 3 Hz was presented to her or him. The signal amplitude was initially 0.1 mA and was increased in steps of 0.1 mA up to a maximum of 1.5 mA. The increase in amplitude was stopped once the subject reported a skin sensation, and the amplitude used in the prior step was used as the maximum magnitude of current stimulation during the next parts of the study. Across the participants we thereby obtained a mean amplitude of 0.67 mA with an SD of 0.24 mA.

The participants listened to diotically presented acoustic stimuli through earphones. For each participant, we determined the signal-to-noise ratio (SNR) corresponding to the comprehension level of 50% using an adaptive procedure (Kollmeier et al., 1988; Kaernbach, 2001) while the sham stimulation was applied. To this end, the subject was asked to listen to the male voice and ignore a competing female voice and to then repeat what they heard. The procedure was started with a randomly selected SNR between -3 dB and 0 dB. If the participant understood no more than two keywords correctly, the SNR was then increased by 1 dB for the next trial; otherwise, it was decreased by 1 dB. The procedure was stopped after seven reversals in the SNR or after running 17 trials. This was done four times for each participant. The average SNRs obtained were then used to adjust the speech envelope according to 50% accuracy.
of the last three SNR values during the last three repetitions was considered as the 50% sentence reception threshold (SRT). Across subjects we thereby obtained a mean SRT of -6.9 dB and a SD of 1.6 dB. To save measurement time, the SRT was only determined for the male voice as the target speaker and not also for the female voice as the target of attention. Although the SRT for the female voice might differ slightly from the SRT for the male speaker, such differences could not bias the subsequent results, which were based on the comprehension of both the male and the female voice together.

For each subject we then used the subject-specific SRT to measure their speech comprehension under tACS with the speech envelope. For each type of tACS waveform, we presented subjects with 26 sentences in noise. Thirteen of these 26 sentences had the female voice as the target and a male voice as distracting talker, and vice versa for the other 13 sentences. The relative contribution of the male and of the female voice in the acoustic mixture therefore depended on whether the male or the female voice was the target of attention. In particular, the SRT was chosen per subject as determined from the adaptive procedure. Because the SRTs were negative for all subjects, this meant that when attention was directed toward the male voice, the male voice was fainter than the female voice. On the other hand, when the subject was instructed to attend to the female voice, the female voice had an amplitude below that of the male voice. Both target and distracting sentences were randomly selected in each trial and were unknown to both the experimenter and the subject. The order of attention was randomly determined, and subjects were informed over a display before each sentence whether they should attend to the male or the female speaker. After listening to a sentence, the participants repeated what they understood. The response was recorded by a microphone. The recorded responses were manually graded by the experimenter to determine the percentage of correctly understood words. The participant took a brief rest for 3 min after every 60 trials.

Data availability. Data and code are available on request.

Results

We first verified that the envelopes of the target and of the distract sentences were indeed unrelated. We found that Pearson’s correlation coefficient between the two envelopes was 0.003 ± 0.002 (mean and SEM, average over 100 pairs of sentences). The correlation coefficient was therefore small and not significantly different from zero (p = 0.98, two-tailed Student’s t-test).

We then proceeded to analyze the effect of tACS on speech comprehension. We found that tACS with the target envelope resulted indeed in a significant modulation (Fig. 2C; Table 1). Both standard multiple linear regression and robust multiple linear regression to fit Equation 3 to the data showed a dependence of the speech comprehension score on the phase shift of the stimulation. In particular, the parameter $B_3$ was significantly different from 0, although the parameters $B_1$ and $A_2$ were not. The dependence of the speech comprehension scores on the phase shift of the stimulation waveform emerged therefore at the longest possible period of 360° but not at the shorter period of 180°. Because of the insignificance of the parameter $B_1$, the amplitude $A_1$ of the variation at the period of 360° followed simply as the parameter $B_2$: $A_1 = B_2$. The corresponding phase shift was $\varphi_1 = 0°$.

Robust regression yielded a significantly higher $r^2$ value than the standard regression ($p < 2 \times 10^{-16}$, bootstrap with 10,000 samples and paired one-tailed Student’s t test). Stimulation with tACS with the target envelope thus modulated speech comprehension by ~3.0%, the value of $A_1 = B_2$ obtained from the robust regression.

Our results regarding the dependence of speech comprehension on the phase of tACS with the target envelope largely replicated our earlier findings (Keshavarzi et al., 2020). Indeed, we previously showed that this type of tACS yields a modulation of speech comprehension so that the highest comprehension score is observed at 0° phase shift and the lowest at a phase shift of ~240°. Here we observed similarly that no phase shift yielded the maximal speech comprehension, whereas we obtained the lowest comprehension scores at a phase shift of 180°. The moderate deviation in the phase shift of the lowest comprehension score might be explained by the different number of phase shifts that we used in the two studies, as well as by experimental uncertainties.

Armed with the ability to assess the influence of tACS on speech comprehension, we then analyzed the effect of tACS with the distractor envelope (Fig. 2D; Table 2). As before, we used multiple linear regression with FDR correction for multiple comparisons to fit Equation 3 to the data and to analyze the statistical significance of the different terms (Fig. 2D; Table 2). As for tACS with the target envelope, we found that the parameter $B_3$ was significant but not the parameters $B_1$ and $A_2$. Importantly, tACS...
with the distractor envelope did therefore lead to a significant modulation of speech comprehension.

The modulation emerged at the period of 360°. Because of the insignificance of the parameter $B_1$, and the negativity of the parameter $B_2$, the amplitude $A_1$ followed as $A_1 = -B_2$ and the corresponding phase shift as $\varphi_1 = 180^\circ$. The amplitude of the modulation of speech comprehension through stimulation with tACS with the distractor envelope was therefore $\approx 1.6\%$. As for stimulation with the target envelope, we found that the robust regression gave a significantly better fit to the data than the standard regression ($p < 2 \times 10^{-16}$, bootstrap with 10,000 samples and paired one-tailed Student's $t$ test).

Although tACS with the target envelope and tACS with the distractor envelope both modulated speech comprehension at the longest-possible period of 360°, there were two important differences between the two types of stimulation. First, the phase shifts that led to the best and worst speech comprehension scores were the opposite in the two cases. Although tACS with the speech envelope at a phase shift of 0° led to the best speech comprehension when based on the envelope of the target speech, this phase shift produced the worst speech comprehension when derived from the envelope of the distractor speech. Inversely, a phase shift of 180° gave the lowest comprehension scores for tACS with the target envelope but the highest comprehension score for tACS with the distractor envelope.

Second, the strength of the modulation of speech comprehension differed between tACS with the target envelope and tACS with the distractor envelope. Specifically, considering the results from the robust regression, tACS with the target envelope yielded a modulation amplitude of $A_1 = 3\%$, which was higher than the modulation amplitude of $A_1 = 1.6\%$ that we obtained from tACS with the distractor envelope ($p = 0.037$, two-tailed Student's $t$ test). We note, however, that the $p$ value was close to 0.05, and that the difference between the amplitudes $A_1$ became insignificant when considering the results from the standard regression ($p = 0.1$, two-tailed Student's $t$ test).

Previous research investigated phase shifts between tACS with the speech envelope and the acoustic waveform in the case of rhythmic speech, where syllables occurred at a specific and fixed frequency. These studies found that the tACS did modulate speech comprehension but that the best and worst phase shifts were highly variable between subjects (Riecke et al., 2018; Zoefel et al., 2018). Similarly, related work on temporal delays between the sound and the neurostimulation waveform found large variability among subjects, and in one case, no significant effect on the population level (Riecke et al., 2018; Wilsch et al., 2018; Erkens et al., 2020; Wang et al., 2020). In our previous work on the role of phase shifts in tACS with the speech envelope, we used, as in this study, natural speech in which syllables did not occur at a single fixed frequency and we found a phase dependency of the resulting speech comprehension that was largely consistent across the different subjects.

We therefore sought to investigate the subject-to-subject variability of speech comprehension for tACS with both target and distractor envelopes. To this end, we first determined for every individual subject and for each of the two stimulation types the phase that yielded the highest comprehension score. We then measured the phase relative to this best phase for that particular subject (Fig. 3A,B).

We used multiple linear regression as well as robust regression to fit Equation 5 to the data and to determine the statistical significance of the parameters. We found, however, no significant modulation of speech comprehension by the relative phases, neither for tACS with the target envelope (Table 3) nor for tACS with the distractor envelope (Table 4). These results showed that the alignment to the best phase increased rather than decreased the noise in the data, indicating that the best phase did not vary substantially between participants.

To further investigate the intersubject variability of the modulation of speech comprehension through tACS, we computed histograms of the best phases of each subject (Fig. 3C,D). For tACS with the target envelope, we found that the distribution of the best phases was significantly different from a uniform distribution ($p = 2 \times 10^{-5},$ Rayleigh test). The distribution had a

### Table 1. Multiple linear regression for the dependence of the speech comprehension scores on the stimulation phase, for stimulation with the target envelope

| Variable | Value | CI        | $p$ value |
|----------|-------|-----------|-----------|
| $A_0$    | $-6.7 \times 10^{-11}$% | $[-0.81, 0.81]$% | 1         |
| $B_1$    | 0.21% | $[-0.93, 1.4]$% | 0.9       |
| $B_2$    | 2.8%*** | $[1.6, 3.9]$% | $3 \times 10^{-5}$ |
| $A_1$    | 0.22% | $[-0.59, 1.03]$% | 0.9       |

### Table 2. Multiple linear regression for the dependence of the speech comprehension scores on the stimulation phase, for stimulation with the distractor envelope

| Variable | Value | CI        | $p$ value |
|----------|-------|-----------|-----------|
| $A_0$    | $3.5 \times 10^{-14}$% | $[-0.5, 0.5]$% | 1         |
| $B_1$    | -0.06% | $[-0.77, 0.65]$% | 1         |
| $B_2$    | -1.6%*** | $[-2.3, -0.9]$% | $9 \times 10^{-5}$ |
| $A_1$    | -0.14% | $[-0.64, 0.36]$% | 1         |
maximum at a phase shift of 0° and a mean phase shift of 4.4°, with an angular deviation of 0.74 and a resultant vector length of 0.72. The distribution could be fitted well with a unimodal von Mises distribution ($p = 0.005$, Watson’s test). This was in line with our finding above that the phase shift of 0° led to the highest speech comprehension scores for this type of tACS (Fig. 2A).

The distribution of the best phases for tACS with the distractor envelope was significantly nonuniform as well ($p = 0.002$, Rayleigh test). However, the best phases occurred around phase shifts of 180°; the maximum of the distribution was at 180°, and its mean was 169°, with an angular deviation of 0.93 and a resultant vector length of 0.57. The unimodal nature of the distribution was confirmed by fitting a von Mises distribution ($p = 0.005$, Watson’s test). The peak in the distribution coincided with the maximum of the speech comprehension scores at this phase shift (Fig. 2B).

Although the angular deviation of the distribution for the best phases obtained for tACS with the distractor envelope was larger than the one for tACS with the target envelope, the difference was not significant ($p = 0.32$, concentration homogeneity test).

We also investigated whether tACS with the target or distractor envelope could improve speech comprehension beyond sham stimulation. We therefore compared the speech comprehension scores at the best phase shift of 0° for tACS with the target envelope as well as at the best phase shift of 180° for tACS with the distractor envelope, to the scores under sham stimulation (Fig. 4). We found that there was no statistically significant difference between the different comprehension scores (one-way ANOVA, $df = 2, F = 1.72, p = 0.2$).

### Discussion

Together, our results showed that tACS with the distractor envelope influenced speech comprehension, providing evidence that the cortical tracking of the ignored speaker plays a functional role in speech processing. Indeed, because the target and the distractor speech signals that we used were unrelated, the envelope of the distractor speech carried no information about the target signal. tACS with the distractor envelope could not therefore influence the cortical tracking of the target speech but only that of the distractor speech.

TACS with the target envelope influenced speech comprehension in a very similar manner to tACS with the distractor envelope. In particular, both types of stimulation led to a modulation of speech comprehension that varied sinusoidally, at the longest possible period, with the applied phase shifts. Moreover, both types of tACS led to a modulation of speech comprehension that had a largely consistent phase dependency across the different subjects. In particular, although earlier studies showed that stimulation parameters such as delay or phase shift could influence speech comprehension in a manner that was highly variable from subject to subject (Riecke et al., 2018; Wilsch et al., 2018; Zoefel et al., 2018), our results did not show such variability. Indeed, when adjusting the phase shift to the best value per subject, we no longer obtained a significant modulation of speech comprehension, indicating that the adjustment increased rather than decreased the noise in the data. This result was in line with our previous studies on the effects of tACS with the speech envelope on speech comprehension (Kadir et al., 2020; Keshavarzi and Reichenbach, 2020; Keshavarzi et al., 2020).

Despite these similarities between tACS with the target and with the distractor envelope, there were two major differences. First, the phase shift that led to the highest speech comprehension score was 0° for tACS with the target envelope and 180° for tACS with the distractor envelope. This difference suggests that tACS with the envelope of a speech signal may act on the early perceptual separability of that signal from an acoustic background. At a phase shift of 0°, tACS with a speech envelope appeared to enhance the neural representation of that speech
stream, tACS with a speech envelope shifted by 180°, in contrast, presumably led to a suppression of the neural encoding of the corresponding speech stream.

This interpretation suggests that the cortical tracking of a speech signal plays, at least partly, a role in auditory stream formation, both for an attended and for an ignored acoustic source (Petersen et al., 2017; Hausfeld et al., 2018; Fiedler et al., 2019). Our finding that tACS with the distractor envelope modulates the comprehension of the target speech then implies that the formation of the two auditory streams of the target and the distractor speech are not independent of each other. Instead, an enhanced representation of one acoustic stream appears to cause a reduced representation of the other stream. Further investigation of such a competition between the cortical tracking of the foreground and of the background acoustic signals may use neuroimaging to quantify the neural responses, for instance paired to tACS and behavioral assessments as explored here.

As the second major difference between tACS with the target and with the distractor envelope, the former type of stimulation led to a stronger modulation of speech comprehension than the latter type. The difference in the modulation amplitudes emerged when analyzing the data with robust regression, accounting for outliers in the data but not when using standard regression. We found indeed that robust regression gave significantly better fits to the data than standard regression. The resulting higher certainty in the fit parameters obtained from robust regression compared with standard regression presumably allowed us to obtain a statistically significant difference in the modulation of speech comprehension between stimulation with the target and with the distractor envelope.

This stronger modulation may point to an effect of tACS with the target envelope that goes beyond auditory stream formation, such as by aiding the parsing of the attended speech stream into syllables and words. Previous research has indeed reported that the comprehension of noise-vocoded speech, in the absence of background noise, can be modulated by tACS with the speech envelope, proving that this type of tACS can influence speech processing beyond acoustic stream formation (Riecke et al., 2018). Because we found that the modulation of speech comprehension through tACS with the target envelope was ~3.0%, almost twice the corresponding value of 1.6% found for tACS with the distractor envelope, our results suggest that tACS with the envelope of a speech signal influenced predominantly auditory stream formation but also had a sizable impact on the further neural processing of that speech stream.

Our results therefore suggest that cortical tracking of an ignored speaker plays a functional role in auditory stream formation, whereas the cortical tracking of an attended speaker likely goes beyond that to encompass further semantic and syntactic processing. These findings are in line with recent results on the encoding of multiple talkers in the auditory cortex. Cortical responses in Heschel’s gyrus have indeed been found to be able to encode an ignored speaker, whereas cortical activity in the superior temporal gyrus responds more to an attended talker (O’Sullivan et al., 2019). In addition, early cortical tracking of acoustic onsets in mixtures of speech signals can occur even for onsets of an ignored speaker that are masked by acoustic activity of a target speaker, suggesting that the activity of the auditory cortex can recover such masked ignored signals, whereas later activity is restricted to features of an attended speaker (Brodbeck et al., 2020). Furthermore, cortical tracking of speech rhythms has been shown to encode information on the lexical as well as on the semantic level for an attended speech stream but not for an ignored one (Brodbeck et al., 2018; Broderick et al., 2018). In addition, behavioral studies have found accordingly that listeners can notice simple acoustic changes in an ignored voice but fail to understand it (Cherry, 1953; Deutsch and Deutsch, 1963; Treisman, 1964; Kidd et al., 2005). However, we also note that speech can be more distracting when intelligible, suggesting that some linguistic information is processed even in ignored speech (Dai et al., 2017).

We found previously that tACS with the target envelope could improve the comprehension of speech in speech-shaped noise beyond sham stimulation (Keshavarzi and Reichenbach, 2020; Keshavarzi et al., 2020). In our present study, however, the speech comprehension scores obtained during tACS with the best phase shifts were not significantly higher than those obtained under sham stimulation. This might be because of the different background noise that we used here, namely a competing talker rather than the previously used speech-shaped noise. The competing talker provided informational masking, whereas speech-shaped noise contributed only energetic masking (Pollack, 1975; Kidd et al., 2008). Informational and energetic masking affect speech comprehension differently (Brungart, 2001; Freyman et al., 2004; Kidd et al., 2016). Moreover, most of the recent studies on tACS with the speech envelope found a detrimental rather than beneficial impact of the stimulation on speech comprehension (Riecke et al., 2018; Wilsch et al., 2018; Kadir et al., 2020). In addition, as opposed
to speech-shaped noise, the amplitude of the competing talker fluctuated in time, allowing for dip listening (Miller and
Licklider, 1950; Bacon and Grantham, 1989; Gustafsson and Arlinger, 1994; Rosen et al., 2013). Both factors may influence
how TACS with the speech envelope affects speech comprehension. Disentangling these aspects will further clarify how
cortical tracking of speech rhythms can affect speech processing and potentially lead to the development of optimized
tACS waveforms for speech-in-noise enhancement.

References
Antal A, Paulus W (2013) Transcranial alternating current stimulation
(TACS). Front Hum Neurosci 7:317.
Bacon SP, Grantham DW (1989) Modulation masking: effects of modulation
frequency, depth, and phase. J Acoust Soc Am 85: 2575–2580.
Beyistol T IL (2018) Applied natural language processing with python. San
Francisco, CA: Apress.
Bird S, Klein E, Loper E (2009) Natural language processing with python:
analyzing text with the natural language toolkit. Sebastopol, CA: O’Reilly
Media, Inc.
Bregman AS (1994) Auditory scene analysis: The perceptual organization of
sound. Cambridge, MA: MIT.
Brodbbeck C, Hong LE, Simon JZ (2018) Rapid transformation from auditory
to linguistic representations of continuous speech. Curr Biol 28:3976–
3983.e5.
Brodbbeck C, Jiao A, Hong LE, Simon JZ (2020) Neural speech restoration at
the cocktail party: auditory cortex recovers masked speech of both
attended and ignored speakers. PLoS Biol 18:e3000883.
Broderick MP, Anderson AJ, Di Liberto GM, Crosse MJ, Lalor EC
(2018) Electrophysiological correlates of semantic dissimilarity
reflect the comprehension of natural, narrative speech. Curr Biol
28:803–809.e3.
Brunhart DS (2001) Informational and energetic masking effects in the
perception of two simultaneous talkers. J Acoust Soc Am 109:
1101–1109.
Cherry EC (1953) Some experiments on the recognition of speech, with one
and with two ears. J. Acoust. Soc. Am. 25: 975–979.
Cooke M, Lu Y (2010) Spectral and temporal changes to speech produced in
the presence of energetic and informational maskers. J Acoust Soc Am
128: 2059–2069.
Coe A, Garcia M, Lecumberri ML, Barker J (2008) The foreign language
cocktail party problem: energetic and informational masking effects in
non-native speech perception. J Acoust Soc Am 123: 414–427.
Dai B, McQueen JM, Hagoort P, Kösem A (2017) Pure linguistic interference
during comprehension of competing speech signals. J Acoust Soc Am
141:EL249–EL254.
Deutsch JA, Deutsch D (1963) Attention: some theoretical considerations.
Psychol Rev 70: 80–90.
Di Liberto GM, O’Sullivan JA, Lalor EC (2015) Low-frequency cortical
entrainment to speech reflects phoneme-level processing. Curr. Biol 25:
2457–2465.
Ding N, Simon JZ (2012) Emergence of neural encoding of auditory objects
while listening to competing speakers. Proc Natl Acad Sci U S A 109:
11854–11859.
Ding N, Simon JZ (2014) Cortical entrainment to continuous speech: func-
tional roles and interpretations. Front Hum Neurosci 8:311.
Ding N, Chatterjee M, Simon JZ (2014) Robust cortical entrainment to the
speech envelope relies on the spectro-temporal fine structure.
Neuroimage 88:41–46.
Erkens J, Schulte M, Vormann M, Herrmann CS (2020) Lacking effects of en-
velope transcranial alternating current stimulation indicate the need to
revisit envelope transcranial alternating current stimulation methods.
Neurosci Letters 15:2633105520936623.
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. J Neurosci 39: 5750–5759.
Fiedler L, Wöstmann M, Herbst SK, Obleser J (2019) Late cortical tracking of
ignored speech facilitates neural selectivity in acoustically challenging
conditions. Neuroimage 186:33–42.
Freyman RL, Balakrishnan U, Helfer KS (2004) Effect of number of masking
talkers and auditory priming on informational masking in speech recogni-
tion. J Acoust Soc Am 115: 2246–2256.
Giraud AL, Poeppel D (2012) Cortical oscillations and speech processing:
emerging computational principles and operations. Nat Neurosci 15:
511–517.
Gustafsson HA, Arlinger SD (1994) Masking of speech by amplitude-
modulated noise. J of the Acoustical Society of America 95: 518–
529.
Hausfeld L, Riecke L, Valente G, Formisano E (2018) Cortical tracking of
multiple streams outside the focus of attention in naturalistic auditory
scenes. Neuroimage 181:617–626.
Helfrich RF, Schneider TR, Rach S, Trautmann-Lengsfeld SA, Engel AK,
Herrmann CS (2014) Entrainment of brain oscillations by transcranial
alternating current stimulation. Curr Biol 24: 333–339.
Hickok G, Small SI (2015) Neurobiology of language. Cambridge, MA:
Academic.
Holland PW, Welsch RE (1977) Robust regression using iteratively
reweighted least-squares. Commun Stat Theory Methods 6:813–
827.
Horton C, D’Zmura M, Srinivasan R (2013) Suppression of competing
speech through entrainment of cortical oscillations. J Neurophysiol
109:3082–3093.
Hyafil A, Fontolan L, Kabelecon C, Guitkin B, Giraud A-L (2015) Speech
encoding by coupled cortical theta and gamma oscillations. Elife 4p.
e06213.
Ingram JC (2007) Neurolinguistics. An introduction to spoken language
processing and its disorders. Cambridge: Cambridge University
Press.
Kadir S, Kaza C, Weissbart H, Reichenbach T (2020) Modulation of speech-
in-noise comprehension through transcranial current stimulation with the
phase-shifted speech envelope. IEEE Trans Neural Syst Rehabil Eng
28:23–31.
Kaernbach C (2001) Adaptive threshold estimation with unforced-choice
tasks. Percept Psychophys 63:1377–1388.
Keshavarzi M, Reichenbach T (2020) Transcranial alternating current
stimulation with the theta-band portion of the temporally-aligned
speech envelope improves speech-in-noise comprehension. Front
Hum Neurosci 14:187.
Keshavarzi M, Kegler M, Kadir S, Reichenbach T (2020) Transcranial
current stimulation in the theta band but not in the delta band
modulates the comprehension of naturalistic speech in noise.
Neuroimage 210:116557.
Kidd G, Mason CR, Richards VM, Gallun FJ, Durlach NI (2008)
Informational masking. In: Auditory perception of sound sources. pp.
143–189. Boston, MA: Springer.
Kidd G, Arborg TL, Mason CR, Gallun FJ (2005) The advantage of know-
where to listen. J Acoust Soc Am 118: 3804–3815.
Kidd G, Mason CR, Swaminathan J, Roverud E, Clayton KK, Best V (2016)
Determining the energetic and informational components of speech-on-
speech masking. J Acoust Soc Am 140: 132–144.
Kollmeier B, Gilkey RH, Sieben UK (1988) Adaptive staircase techni-
ques in psychoacoustics: a comparison of human data and a mathe-
matical model. J of the Acoustical Society of America 83:1852–
1862.
Mesgarani N, Chang EF (2012) Selective cortical representation of
attended speaker in multi-talker speech perception. Nature 485:
233–236.
Miller GA, Licklider JCR (1950) The intelligibility of interrupted speech.
S. J. Neurosci., June 9, 2021 • 41(12):5093–5101
Keshavarzi et al. • Cortical Tracking of Background Speech

Freedman 109:311.
Pollack I (1975) Auditory informational masking. J Acoust Soc Am 57: S5–S5.
Riecke L, Formisano E, Sorger B, Baškent D, Gaudrain E (2018) Neural entrainment to speech modulates speech intelligibility. Curr Biol 28:161–169.
Rosen S, Souza P, Ekelund C, Majeed AA (2013) Listening to speech in a background of other talkers: effects of talker number and noise vocoding. J Acoust Soc Am 133: 2431–2443.
Snyder JS, Alain C (2007) Toward a neurophysiological theory of auditory stream segregation. Psychol Bull 133: 780–799.
Treisman AM (1964) Verbal cues, language and meaning in selective attention. Am J Psychol 77: 206–219.
Vanthorhout J, Decruy L, Wouters J, Simon JZ, Francart T (2018) Speech intelligibility predicted from neural entrainment of the speech envelope. J Assoc Res Otolaryngol 19:181–191.
Wang Y, Shi L, Dong G, Zhang Z, Chen R (2020) Effects of transcranial electrical stimulation on human auditory processing and behavior—a review. Brain Sciences 10:531.
Weisbart H, Kandylaki KD, Reichenbach T (2020) Cortical tracking of surprisal during continuous speech comprehension. J Cogn Neurosci 32:155–166.
Wilsch A, Neuling T, Obleser J, Herrmann CS (2018) Transcranial alternating current stimulation with speech envelopes modulates speech comprehension. Neuroimage 172:766–774.
Wood N, Cowan N (1995) The cocktail party phenomenon revisited: how frequent are attention shifts to one’s name in an irrelevant auditory channel? J Exp Psychol Learn Mem Cogn 21: 255–260.
Zoefel B, Archer-Boyd A, Davis MH (2018) Phase entrainment of brain oscillations causally modulates neural responses to intelligible speech. Curr Biol 28:401–408.e5.