Temporal coherence of sound fluctuations across spectral channels is thought to aid auditory grouping and scene segregation. Although prior studies on the neural bases of temporal-coherence processing focused mostly on cortical contributions, neurophysiological evidence suggests that temporal-coherence-based scene analysis may start as early as the cochlear nucleus (i.e., the first auditory region supporting cross-channel processing over a wide frequency range). Accordingly, we hypothesized that aspects of temporal-coherence processing that could be realized in early auditory areas may shape speech understanding in noise. We then explored whether physiologically plausible computational models could account for results from a behavioral experiment that measured consonant categorization in different masking conditions. We tested whether within-channel masking of target-speech modulations predicted consonant confusions across the different conditions and whether predictions were improved by adding across-channel temporal-coherence processing mirroring the computations known to exist in the cochlear nucleus. Consonant confusions provide a rich characterization of error patterns in speech categorization, and are thus crucial for rigorously testing models of speech perception; however, to the best of our knowledge, they have not been used in prior studies of scene analysis. We find that within-channel modulation masking can reasonably account for category confusions, but that it fails when temporal fine structure cues are unavailable. However, the addition of across-channel temporal-coherence processing significantly improves confusion predictions across all tested conditions. Our results suggest that temporal-coherence processing strongly shapes speech understanding in noise and that physiological computations that exist early along the auditory pathway may contribute to this process.

Key words: cochlear nucleus; comodulation masking release; computational modeling; consonant confusions; cross-channel processing; wideband inhibition

Significance Statement
Temporal coherence of sound fluctuations across distinct frequency channels is thought to be important for auditory scene analysis. Prior studies on the neural bases of temporal-coherence processing focused mostly on cortical contributions, and it was unknown whether speech understanding in noise may be shaped by across-channel processing that exists in earlier auditory areas. Using physiologically plausible computational modeling to predict consonant confusions across different listening conditions, we find that across-channel temporal coherence contributes significantly to scene analysis and speech perception and that such processing may arise in the auditory pathway as early as the brainstem. By virtue of providing a richer characterization of error patterns not obtainable with just intelligibility scores, consonant confusions yield unique insight into scene analysis mechanisms.

Introduction
An accumulating body of evidence suggests that temporal-coherence processing is important for auditory scene analysis (Elhilali et al., 2009). Indeed, a rich psychophysical literature on grouping (Darwin, 1997), comodulation masking release (CMR; Schooneveldt and Moore, 1987), cross-channel modulation interference (Apoux and Bacon, 2008), and pitch-based masking release (Oxenham and Simonson, 2009) supports the theory that temporally coherent sound modulations can bind together sound elements across distinct spectral channels to form a perceptual object, which can help perceptually
segregate different sources in an acoustic mixture. This theory may help explain how we perform speech separation in a multisource environment (Krishnan et al., 2014), as speech naturally has common temporal fluctuations across channels, particularly in the syllabic (0–5 Hz), phonemic (5–64 Hz), and pitch (64–300 Hz) ranges (Crouzet and Ainsworth, 2001; Swaminathan and Heinz, 2011).

Object binding and scene segregation are perceptually defined phenomena, whose neural correlates are yet to be definitively established. These phenomena may in general be supported by a cascade of mechanisms throughout the auditory pathway (Pressnitzer et al., 2008; Shinn-Cunningham, 2020; Mishra et al., 2021). Prior studies on the neural bases of temporal-coherence processing mostly focused on cortical contributions (Elhilali et al., 2009; Teki et al., 2013; O’Sullivan et al., 2015). However, single-unit measurements and computational modeling of across-channel CMR effects suggest that temporal-coherence-based scene analysis may start early in the auditory pathway; for instance, the cochlear nucleus has the physiological mechanisms (e.g., wideband inhibition) needed to support such analysis (Pressnitzer et al., 2001; Meddis et al., 2002). Moreover, attention, which operates on segregated auditory objects (Shinn-Cunningham, 2008), affects responses in the early auditory cortex (Hillyard et al., 1973). Given this, binding and scene segregation likely start even earlier, such as in brainstem, and accumulate along the auditory pathway. However, no prior studies have directly tested the hypothesis that speech understanding in noise may be shaped by aspects of temporal-coherence processing that exist in early auditory areas.

Previous studies of temporal-coherence processing mostly used nonspeech stimuli (Elhilali et al., 2009; Teki et al., 2013; O’Sullivan et al., 2015). Moreover, a parallel literature on modeling speech-intelligibility mechanisms typically focused on overall intelligibility to test predictions of performance (Jørgensen et al., 2013; Relaño-Iborra et al., 2016). A detailed characterization of error patterns in speech categorization—crucial to rigorously examine any theory of speech perception—has not been previously used in studies of scene analysis. In contrast, confusion patterns in speech categorization, such as consonant confusion matrices (Miller and Nicely, 1955), have been widely used in the speech acoustics and cue-weighting literatures and can provide deeper insight into underlying mechanisms if used to test theories of scene analysis.

To address these gaps, we used a combination of online consonant identification experiments and computational modeling of temporal-coherence processing that is physiologically plausible in the cochlear nucleus (Pressnitzer et al., 2001), the first auditory area where cross-channel processing over a wide frequency range is supported. We asked whether the masking of target-speech envelopes by distracting maskermodulations (i.e., modulation masking: Bacon and Grantham, 1989; Stone and Moore, 2014) within individual frequency channels (as implemented in current speech-intelligibility models; Jørgensen et al., 2013; Relaño-Iborra et al., 2016) is sufficient to predict consonant categorization, or if across-channel temporal-coherence processing improves predictions by accounting for interference from masker elements that are temporally coherent with target elements but in different frequency channels. Crucially, instead of just trying to predict perceptual intelligibility measurements from model outputs, we predicted consonant confusion patterns in various listening conditions. Considering the error patterns in consonant categorization provided a richer characterization of the processes engaged during speech perception compared to looking only at percent correct scores. Our combined use of consonant confusions and physiologically plausible computational modeling provides independent evidence for the role of temporal-coherence processing in scene analysis and speech perception. Moreover, it suggests that this processing may start earlier in the auditory pathway than previously thought.

### Materials and Methods

**Stimulus generation.** The stimuli used in the present study draw from and expand on the materials and methods previously described in Viswanathan et al. (2021b). Twenty consonants from the Speech Test Video (STeVi) corpus (Sensimetrics) were used. The consonants were

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**Table 1. Rationale for the different stimulus conditions included in this study**

<table>
<thead>
<tr>
<th>No.</th>
<th>Stimulus condition</th>
<th>Rationale for inclusion in study</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>SiQuiet</td>
<td>Used as a control condition</td>
</tr>
<tr>
<td>2</td>
<td>SiSSN at −8 dB SNR</td>
<td>Widely used in the literature; used for calibration of prediction model</td>
</tr>
<tr>
<td>3</td>
<td>SiB at −8 dB SNR</td>
<td>Simulates ecologically relevant cocktail-party listening</td>
</tr>
<tr>
<td>4</td>
<td>SiDCmod at −18 dB SNR</td>
<td>To obtain a different modulation masking profile from stationary noise (which contains relatively more high-frequency modulation energy) and babble (which contains relatively more low-frequency modulation power; Viswanathan et al., 2021a)</td>
</tr>
<tr>
<td>5</td>
<td>SiB at 0 dB SNR subjected to 64-channel envelope vocoding (Vocoded SiB)</td>
<td>Used to compare performance across models that consider TFS and those that do not (as TFS can influence scene analysis and can convey consonant voicing information in noise; Viswanathan et al., 2021a,b)</td>
</tr>
</tbody>
</table>

The different listening conditions were chosen to span a range of modulation masking spectral profiles and TFS information, which allows for theories of scene analysis based on within-channel modulation masking and across-channel temporal coherence to be tested in a rigorous manner. Collectively, these conditions represent a diversity of scene acoustics, including important examples in our environment and clinical applications. The SNR levels were chosen to give approximately equal overall intelligibility across SiSSN, SiB, SiDCmod, and Vocoded SiB using a behavioral pilot study with three subjects who did not participate in the online consonant identification experiment. This was done to obtain roughly equal variance in the consonant confusion estimates for these conditions, which allows us to fairly compare confusion patterns across them. Equalizing intelligibility also maximizes the statistical power for detecting differences in the pattern of confusions. The overall intelligibility in each of these conditions was: −60%, which yielded a sufficient number of confusions for analysis.

**Table 2. Phonetic features of the 20 English consonants used in this study**

<table>
<thead>
<tr>
<th>Consonant</th>
<th>Voicing</th>
<th>MDA</th>
<th>PDA</th>
</tr>
</thead>
<tbody>
<tr>
<td>/b/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Bilabial</td>
</tr>
<tr>
<td>/d/</td>
<td>Unvoiced</td>
<td>Affricative</td>
<td>Palatal</td>
</tr>
<tr>
<td>/θ/</td>
<td>Unvoiced</td>
<td>Affricative</td>
<td>Velar</td>
</tr>
<tr>
<td>/s/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/z/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/s/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/m/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/n/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/p/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/f/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/v/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/θ/</td>
<td>Unvoiced</td>
<td>Affricative</td>
<td>Palatal</td>
</tr>
<tr>
<td>/s/</td>
<td>Unvoiced</td>
<td>Affricative</td>
<td>Labiodental</td>
</tr>
<tr>
<td>/l/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/r/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/z/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
<tr>
<td>/s/</td>
<td>Voiced</td>
<td>Stop</td>
<td>Velar</td>
</tr>
</tbody>
</table>

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Viswanathan et al. • Consonant Confusions and Temporal Coherence

Viswanathan et al. (2021b) Table 1

Mok et al., 2021 condition was found to be 1.75:1. The EI ratio was adjusted to obtain the best consonant confusion prediction accuracy for SiSSN (i.e., the calibration condition), and the optimal ratio for the calibration

2 ms. Note that our model simulations were rate based; that is, they used AN PSTHs rather than spikes. Thus, all outputs were half-wave rectified (i.e., firing rates were positive at every stage).

cochlear nucleus (CN). CN units at different CFs form the building blocks of this circuit. Each CN unit consists of a narrowband cell (NB) that receives narrow on-CF excitatory input from the auditory nerve (AN) and inhibitory input from a wideband inhibitor (WBI). The WBI in turn receives excitatory inputs from AN fibers tuned to CFs spanning 2 octaves below to 1 octave above the

cochlear nucleus (CN). AN fibers and inhibitory input from a wideband inhibitor (WBI). The WBI in turn receives excitatory inputs from AN fibers tuned to CFs spanning 2 octaves below to 1 octave above the

/ʃ/, /θ/, /r/, /s/, /z/, /ʒ/. The consonants were presented in consonant-vowel (CV) context, where the vowel was always /a/. Each consonant was spoken by two female and two male talkers (to reflect real-life talker variability).

channel was replaced with a noise carrier. The envelope in each channel was then used to modulate a random Gaussian white noise carrier; the result was band-pass filtered within the channel bandwidth and scaled to match the level of the original signal. We verified that the vocoding procedure did not significantly change envelopes at the cochlear level, as described in Viswanathan et al. (2021b). Table 1 describes the rationale behind including these different stimulus conditions in our study.

The stimulus used for online volume adjustment was running speech mixed with four-talkers babble. The speech and babble samples were obtained from the QuickSIN corpus (Killion et al., 2004); these were repeated over time to obtain a ~20 s total stimulus duration to give subjects sufficient time to adjust their computer volume with the instructions described in Experimental design. The root mean square value of this stimulus corresponded to 75% of the dB difference between the softest and loudest stimuli in the consonant identification experiment, which ensured that no stimulus was too loud for subjects once they had adjusted their computer volume to a comfortable level.

Participants. Full details of participant recruitment and screening are provided in Viswanathan et al. (2021b) and are only briefly reviewed here. Anonymous subjects were recruited for online data collection using Prolific.co. A three-part subject-screening protocol developed and validated by Mok et al. (2021) was used to restrict the subject pool. This protocol included a survey on age, native-speaker status, presence of persistent tinnitus, and history of hearing and neurologic diagnoses, followed by headphone/earphone checks and a speech-in-babble-based hearing screening. Subjects who passed this screening protocol were invited to participate in the consonant identification study, and when they returned, headphone/earphone checks were performed again. Only subjects who satisfied the following criteria passed the screening protocol: (1) 18–55 years old; (2) self-reported no hearing loss, neurologic disorders, or persistent tinnitus; (3) born and residing in the United States/Canada and a native speaker of North American English; (4) experienced Prolific subject; and (5) passed the headphone/earphone checks and speech-in-babble-based hearing screening (Mok et al., 2021).

Experimental design. The online consonant identification experiment was previously described in Viswanathan et al. (2021b). Subjects performed the experiment using their personal computers

Figure 1. CMR circuit based on wideband inhibition in the cochlear nucleus. This physiologically plausible circuit was proposed by Pressnitzer et al. (2001) to model CMR effects seen in the cochlear nucleus (CN). CN units at different CFs form the building blocks of this circuit. Each CN unit consists of a narrowband cell (NB) that receives narrow on-CF excitatory input from the auditory nerve (AN) and inhibitory input from a wideband inhibitor (WBI). The WBI in turn receives excitatory inputs from AN fibers tuned to CFs spanning 2 octaves below to 1 octave above the CF of the NB that it inhibits. The time constants for the excitatory and inhibitory synapses are 5 ms and 1 ms, respectively. The WBI input to the NB is delayed with respect to the AN input by 2 ms. Note that our model simulations were rate based; that is, they used AN PSTHs rather than spikes. Thus, all outputs were half-wave rectified (i.e., firing rates were positive at every stage). All synaptic filters were initially normalized to have unit gain, then the gain of the inhibitory input was allowed to vary parametrically to implement different excitation-to-inhibition (EI) ratios between 3:1 and 1:1. The EI ratio was adjusted to obtain the best consonant confusion prediction accuracy for SiSSN (i.e., the calibration condition), and the optimal ratio for the calibration condition was found to be 1.75:1.
and headphones/earphones. Our online infrastructure included checks to prevent the use of mobile devices. The experiment consisted of the following parts: (1) headphone/earphone checks, (2) demonstration (Demo), and (3) Test. Each of these three parts had a volume-adjustment task at the beginning. In this task, subjects were asked to make sure that they were in a quiet room and wearing wired (not wireless) headphones or earphones. They were instructed not to use desktop/laptop speakers. Headphone/earphone use was checked using the procedures in Mok et al. (2021). They were then asked to set their computer volume to 10–20% of the full volume, after which they were played a speech-in-babble stimulus and asked to adjust their volume up to a comfortable but not too loud level. Once subjects had adjusted their computer volume, they were instructed not to adjust the volume during the experiment, as that could lead to sounds being too loud or soft.

The Demo stage consisted of a short training task designed to familiarize subjects with how each consonant sounds and with the consonant identification paradigm. Subjects were instructed that in each trial they would hear a voice say, “You will mark *something* please.” They were told that they would be given a set of options for “something” at the end of the trial, and that they should click on the corresponding option. After subjects had heard all consonants sequentially (i.e., the same order as the response choices) in quiet, they were tasked with identifying consonants presented in random order and spanning the same set of listening conditions as the Test stage. Subjects were instructed to ignore any background noise and only listen to the particular voice saying, “You will mark *something* please.” To ensure that all subjects understood and were able to perform the task, only those subjects who scored ≥85% in the Demo’s SiQuiet control condition were selected for the Test stage.

Subjects were given instructions in the Test stage similar to those in the Demo but told to expect trials with background noise from the beginning. The Test stage presented, in random order, the 20 consonants (with one stimulus repetition per consonant) across all four talkers and all five experimental conditions; the only exception to this rule was in the first subpart of the Test stage. Subjects were not told what consonant was presented to avoid overtraining to the acoustics of how each consonant sounded across the different conditions; the only exception to this rule was in the first subpart of the Demo, where subjects heard all consonants in quiet in sequential order.

Separate studies were posted on Prolific.co for the different talkers. When a subject performed a particular study, they would be presented with the speech stimuli for one specific talker consistently over all trials. Thus, each subject was not just trained with one talker but tested with all talkers to avoid training-testing disparities. To obtain results that are generalizable, we used 50 subjects per talker (subject overlap between talkers was not controlled); with four talkers, this yielded 200 subject-talker pairs or data samples. Within each talker and condition, all subjects performed the task with the same stimuli. Moreover, all condition effect contrasts were computed on a within-subject basis and averaged across subjects.

**Data preprocessing.** Only samples with intelligibility scores ≥85% for the SiQuiet condition in the Test stage were included in results reported here. All conditions for the remaining samples were excluded from further analyses as a data quality control measure. This yielded a final N = 191 samples.

**Quantifying confusion matrices from perceptual measurements.** The 20 English consonants used in this study were assigned the phonetic features described in Table 2. The identification data collected in the Test stage were used to construct consonant confusion matrices (pooled over samples) for the different conditions; these matrices in turn were used to construct voicing, place of articulation (POA), and manner of articulation (MOA) confusion matrices by pooling over all consonants.

Given that all psychophysical data were collected online, we performed data quality checks; the analyses performed and the results are described in detail in Mok et al. (2021) and Viswanathan et al. (2021b), and are only briefly presented here. We compared consonant confusions for SiSSN, a commonly used condition in the literature, with previous lab-based findings. Phatak and Allen (2007) found that for a given overall intelligibility, recognition scores vary across consonants. They identified three groups of consonants, C1, C2, and C3 with low, high, and intermediate recognition scores, respectively, in speech-shaped noise. The SiSSN data that we collected online closely replicated that key trend for the groups they identified. Moreover, based on a graphical analysis of confusion patterns in speech-shaped noise, Phatak and Allen (2007) identified perceptual clusters (i.e., sets where one consonant is confused most with another in the same set). In the current study too, we identified perceptual clusters for SiSSN by subjecting the consonant confusion matrix to a hierarchical clustering analysis (Ward, 1963); our results closely replicated the lab-based clustering results of Phatak and Allen (2007). As previous lab-based results were not readily available for the
remaining masking conditions in our study, we instead examined whether subjects randomly chose a different consonant from what was presented when they made an error, or if there was more structure in the data. The percent errors in our data fell outside the distributions expected from random confusions, suggesting that the error patterns have a nonrandom structure. Together, these results support the validity of our online-collected data.

We wished to test whether there are any significant differences in consonant confusion patterns across the different masking conditions, namely, SiSSN, SiB, SiDCmod, and Vocoded SiB. If so, these differences could then be predicted by computational modeling to test our hypothesis about the role of temporal-coherence-based across-channel masking of target speech by noise fluctuations. As the SiQuiet condition was intended to primarily be used as a control condition to ensure data quality (see Data preprocessing), SiQuiet data were not subjected to this analysis. To test whether confusion patterns differed across the masking conditions, we first normalized the overall intelligibility for these conditions to 60% by scaling the consonant confusion matrices such that the sum of the diagonal entries was the desired intelligibility (note that overall intelligibility was not normalized for the main modeling analyses of this study). By matching intelligibility in this manner, differences in consonant categorization and category errors rather than differences in overall error counts because of one condition being inherently easier at a particular SNR. Because overall intelligibility was similar across the masking conditions to start with (see Fig. 5), small condition differences in intelligibility could be normalized without loss of statistical power. Confusion-matrix differences between the intelligibility-matched conditions and talkers. These responses were then used as inputs to the auditory-periphery model. The parameters of this model were set as follows. Thirty cochlear filters with characteristic frequencies (CFs) equally spaced on an ERB-number scale (Glasberg and Moore, 1990) between 125 and 8000 Hz were used. Normal function was chosen for the outer and inner hair cells. The species was chosen to be human with the Shera et al. (2002) cochlear tuning at low sound levels; however, with suppression, the Glasberg and Moore (1990) tuning is effectively obtained for our broadband, moderate-level stimuli (Heinz et al., 2002; Oxenham and Shera, 2003). The noise type parameter for the inner-hair-cell synapse model was set to fixed fractional Gaussian noise to yield a constant spontaneous auditory-nerve firing rate. To avoid single-fiber saturation effects, the spontaneous rate of the auditory-nerve fiber was set to 10, corresponding to that of a medium-spontaneous-rate fiber. An approximate implementation of the power-law adaptation dynamics in the synapse was used. The absolute and relative refractory periods were set to 0.6 ms.

The periphery model was simulated with the same speech stimuli used in our psychophysical experiment (i.e., CV utterances that spanned 20 consonants, four talkers, and five conditions, and were embedded in a carrier phrase) as input. The level for the target speech was set to 60 dB SPL across all stimuli, as this produced sufficient (i.e., firing rate greater than spontaneous rate) model auditory-nerve responses for consonants in quiet and also did not saturate the response to the loudest stimulus. The periphery model was provided with just one audio channel input for all conditions except SiDCmod, as that was the only condition that was dichotic rather than diotic. Instead, for SiDCmod, the model was separately simulated for each of the two audio channels. Two hundred stimulus repetitions were used to derive peristimulus time histograms (PSTHs) from model auditory-nerve outputs. The model was simulated for the full duration of each stimulus, as opposed to just the time period when the target consonant was presented. A PSTH bin width of 1 ms (i.e., a sampling rate of 1 kHz) was used. This was done to capture fine-structure phase locking up to and including the typical frequency range of human pitch for voiced sounds. In the case of the SiDCmod condition, a separate PSTH was computed for each of the two dichotic audio channels.

Although the full speech stimuli (including the carrier phrase and CV utterances) were used as inputs to the periphery model, the responses to the target consonants were segmented out from the model PSTHs before being input into the scene analysis models. This segmentation had to be performed manually because the duration of the carrier phrase varied across consonants and talkers, and the start and end times corresponding to any given target consonant were unknown a priori. The time segment corresponding to when the target consonant was presented was calculated for each speech-in-quiet stimulus by visualizing speech spectrograms computed by gammatone filtering (Patterson et al., 1987) followed by Hilbert-envelope extraction (Hilbert, 1906). One hundred twenty-eight gammatone filters were used for this purpose, with center frequencies between 100 and 8000 Hz and equally spaced on an ERB-number scale (Glasberg and Moore, 1990). A fixed duration of 104.2 ms was used for each consonant segment. Segmentation accuracy was verified by listening to the segmented consonant utterances. The time segments thus derived were used to extract model auditory-nerve responses to the different target consonants across the different conditions and talkers. These responses were then used as inputs to the scene analysis models described below.

Figure 3. Stimuli used to validate the CMR circuit model. The stimuli used were from Pressnitzer et al. (2001), and consisted of a target signal in a 10 Hz 100% SAM tonal complex masker. The masker differed depending on the experimental condition. In the Reference condition, the masker was a 1.1-kHz-carrier SAM tone (referred to as the OFC). In the Comodulated and Codeviant conditions, six flanking components were presented in addition to the OFC. The flanking components were SAM tones at the same level as the OFC. The flanking components were separated from the OFC by —800, —600, —400, 400, 600, and 800 Hz, respectively. The modulation of each flanking component was in phase with the OFC modulation in the Comodulated condition, but 180° out of phase with the OFC modulation in the Codeviant condition. The target signal was a 50-ms-long 1.1 kHz tone pip that was presented in the dips of the OFC modulation during the last 0.3 s of the stimulus period (i.e., in the last 3 dips) at different values of SCR (defined as the signal maximum amplitude over the amplitude of the OFC before modulation).
Scene analysis modeling to predict consonant confusions. To study the contribution of across-channel temporal-coherence processing to consonant categorization, we constructed two different scene analysis models. The first is a within-channel modulation-masking-based scene analysis model inspired by Relano-Iborra et al. (2016), and the second is a simple across-channel temporal coherence model mirroring the physiological computations that are known to exist in the cochlear nucleus (Pressnitzer et al., 2001).

In the within-channel modulation-masking-based model, the auditory-nerve PSTHs (i.e., the outputs from the periphery model; see Auditory periphery modeling) corresponding to the different consonants, conditions, and talkers were filtered within a 1 ERB bandwidth (Glasberg and Moore, 1990) to extract band-specific envelopes. Note that the envelopes extracted from auditory-nerve outputs may contain some TFS converted to envelopes via inner-hair-cell rectification (assuming envelope and TFS are defined at the output of the cochlea), but that is the processing that is naturally performed by the auditory system as well. Pairwise dynamic time warping (Rabiner, 1993) was performed to align the results for each pair of consonants across time. Dynamic time warping can help compensate for variations in speaking rate across consonants. A modulation filterbank (Ewert and Dau, 2000; Jørgensen et al., 2013) was then used to decompose the results at each CF into different modulation frequency (MF) bands. This filterbank consists of a low-pass filter with a cutoff frequency of 1 Hz in parallel with eight bandpass filters with octave spacing, a quality factor of 1, and center frequencies ranging from 2 to 256 Hz. For each condition, talker, CF, MF, and consonant, Pearson correlation coefficients were computed between the filterbank output for that consonant in that particular condition and the output for each of all 20 consonants in quiet. Each of the individual correlations was squared to obtain the variance explained; the
We wanted to test whether across-channel temporal-coherence processing of input fluctuations could better predict consonant categorization than a purely within-channel modulation masking model. To simulate across-channel temporal-coherence processing, we modeled a physiologically plausible wideband-inhibition-based temporal-coherence processing circuit proposed by Pressnitzer et al. (2001) to account for physiological correlates of CMR in the cochlear nucleus. A schematic of this circuit is provided in Figure 1. Note that the circuit model parameter corresponding to the excitation-to-inhibition ratio cannot be readily compared to its physiological correlate because the model is rate based and lacks important membrane conductance properties that spiking models can be endowed with. The overall across-channel scene analysis model is similar to the within-channel model, except that the envelope extraction stage of the within-channel model is replaced with the CMR circuit model in the across-channel model. Thus, the across-channel model can account for both within-channel modulation masking effects as well as across-channel temporal-coherence processing. Figure 2 shows schematics of both the within- and across-channel models.

To verify that the CMR circuit model (Fig. 1) produced physiological correlates of CMR similar to those reported by Pressnitzer et al. (2001), we used the same complex stimuli that they used (Fig. 3). The stimuli consisted of a target signal in a 100% sinusoidally amplitude-modulated (SAM) tonal complex masker. There were three experimental conditions: Reference, Comodulated, and Codeviant. In the Reference condition, the masker had just one component, a SAM tone with a carrier frequency of 1.1 kHz (to allow comparison to data from Pressnitzer et al., 2001); this masking component is also referred to as the on-frequency component (OFC). The Comodulated and Codeviant conditions presented the OFC along with six flanking components that were SAM tones at the same level as the OFC. The carrier frequency separation between the different flanking components and the OFC were −800, −600, −400, 400, 600, and 800 Hz, respectively. The flanking components were modulated in phase with the OFC in the Comodulated condition, and 180° out of phase with the OFC in the Codeviant condition. A 10 Hz modulation rate was used for all SAM tones. The target signal consisted of a 50-ms-long (i.e., half of the modulation time period) tone pip at 1.1 kHz that was presented in the dips of the OFC modulation during the last 0.3 s of the stimulus period (i.e., in the last three dips) at different values of signal-to-component ratio (SCR; defined as the signal maximum amplitude over the amplitude of the OFC before modulation). These stimuli were presented to the periphery model, and the corresponding model outputs were passed into the CMR circuit model.

The rate-level function at the output of the CMR circuit model (Fig. 4D) closely matches physiological data for chopper units in the ventral cochlear nucleus (Winter and Palmer, 1990) and was used to set the masker level for the CMR stimuli. The firing-rate threshold was 0 dB SPL for pure-tone inputs at CF; thus, a fixed level of 40 dB SPL (i.e., 40 dB SL) was used for the OFC. The PST outputs from the CMR circuit model (at 1.1 kHz CF) are shown in Figure 4A. The time-averaged statistics of the firing rate during the last 0.3 s of the stimulus period and in the absence of the target signal were used as the null distribution against which the neurometric sensitivity, d′, was calculated; a separate null distribution was derived for each condition. The average firing rate during the target signal periods was compared with the corresponding null distribution to estimate a separate d′ for each SCR threshold. The d′ values cannot be interpreted in a conventional manner given that the choice of window used to estimate the null-distribution parameters introduces an arbitrary scaling; thus, our choice of the d′ criterion to calculate CMR was instead based on avoiding floor and ceiling effects. Results indicate that the CMR circuit model shows a CMR effect consistent with actual cochlear nucleus data in that signal detectability is best in the Comodulated condition, followed by the Reference and Codeviant conditions (compare Figs. 4A and B with Figs. 2 and 6A from Pressnitzer et al., 2001).

The size of the predicted CMR effect is also consistent with perceptual measurements (Mok et al., 2021). As expected, no CMR effect is seen at the level of the auditory nerve. Thus, the CMR circuit model accounts for the improved signal representation in the Comodulated condition where the masker is more easily separable from the target signal, an advantage that derives from the fact that the different masking components are temporally coherent with one another. In addition, it also accounts for the greater cross-channel interference in the Codeviant condition, where the flanking components are temporally coherent with the target signal that is presented in the dips of the OFC. Finally, when the modulation rate of the input SAM tones was varied, CMR effects were still seen (Fig. 4C) and followed the same low-pass trend as human perceptual data (Carlyon et al., 1989).

Each scene analysis model was separately calibrated by fitting a logistic/sigmoid function mapping the neural consonant confusion matrix entries from that model for the SISN condition to corresponding perceptual measurements. The mapping derived from this calibration was used to predict perceptual consonant confusion matrices from the corresponding neural confusion matrices for unseen conditions. Voicing, POA, and MOA confusion matrices were then derived by pooling over all consonants. Finally, the Pearson correlation coefficient was used to compare model predictions to perceptual measurements across the voicing, POA, and MOA categories. The prediction accuracy for the different models is reported in Results.

Statistical analysis. Permutation testing (Nichols and Holmes, 2002) with multiple-comparisons correction at 5% false discovery rate (FDR; Benjamini and Hochberg, 1995) was used to extract significant differences in the SISN, SiB, SiDCmod, and Voiced SiB consonant confusion matrices (across-channel perceptual measurements). The null distributions for permutation testing were obtained using a nonparametric shuffling procedure, which

---

### Table 3. Pearson correlation coefficients between within-channel model predictions and perceptual measurements

<table>
<thead>
<tr>
<th>Condition</th>
<th>Diagonal entries</th>
<th>Off-diagonal entries</th>
<th>All entries</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Correlation</td>
<td>p-value</td>
<td>Correlation</td>
</tr>
<tr>
<td>SiSSN</td>
<td>83%</td>
<td>0.0002***</td>
<td>67%</td>
</tr>
<tr>
<td>SiB</td>
<td>72%</td>
<td>0.0026***</td>
<td>64%</td>
</tr>
<tr>
<td>SiDCmod</td>
<td>66%</td>
<td>0.0072**</td>
<td>64%</td>
</tr>
<tr>
<td>Vocoder SiB</td>
<td>4%</td>
<td>0.4445</td>
<td>40%</td>
</tr>
</tbody>
</table>

Results are listed separately for the diagonal entries of the confusion matrix (i.e., proportion correct for the different consonant phonetic categories), off-diagonal entries (i.e., true confusions), and across all entries. Note that p-value ranges are mapped to symbols as follows: *** indicates 0 ≤ p < 0.001, ** indicates 0.001 ≤ p < 0.01, and * indicates 0.01 ≤ p < 0.05.

### Table 4. Pearson correlation coefficients between across-channel model predictions and perceptual measurements

<table>
<thead>
<tr>
<th>Condition</th>
<th>Diagonal entries</th>
<th>Off-diagonal entries</th>
<th>All entries</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Correlation</td>
<td>p-value</td>
<td>Correlation</td>
</tr>
<tr>
<td>SiSSN</td>
<td>83%</td>
<td>10^-12***</td>
<td>81%</td>
</tr>
<tr>
<td>SiB</td>
<td>85%</td>
<td>0.0001***</td>
<td>73%</td>
</tr>
<tr>
<td>SiDCmod</td>
<td>86%</td>
<td>10^-12***</td>
<td>72%</td>
</tr>
<tr>
<td>Vocoder SiB</td>
<td>63%</td>
<td>0.0131***</td>
<td>70%</td>
</tr>
</tbody>
</table>

Results are listed separately for the diagonal entries of the confusion matrix (i.e., proportion correct for the different consonant phonetic categories), off-diagonal entries (i.e., true confusions), and across all entries. Note that p-value ranges are mapped to symbols as follows: *** indicates 0 ≤ p < 0.001, ** indicates 0.001 ≤ p < 0.01, and * indicates 0.01 ≤ p < 0.05.
Table 5. Improvement in prediction accuracy offered by the across-channel model compared to the within-channel model

<table>
<thead>
<tr>
<th>Condition</th>
<th>Diagonal entries</th>
<th>Off-diagonal entries</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Improvement</td>
<td>Uncorrected p-value</td>
</tr>
<tr>
<td>SiB</td>
<td>12%</td>
<td>0.0225</td>
</tr>
<tr>
<td>SiDCmod</td>
<td>22%</td>
<td>&lt;10^-5</td>
</tr>
<tr>
<td>Vocoder SiB</td>
<td>59%</td>
<td>&lt;10^-5</td>
</tr>
<tr>
<td></td>
<td>8%</td>
<td>0.0406</td>
</tr>
<tr>
<td></td>
<td>8%</td>
<td>0.1006</td>
</tr>
<tr>
<td></td>
<td>30%</td>
<td>&lt;10^-5</td>
</tr>
</tbody>
</table>

The across-channel model showed improved correlations between model predictions and perceptual measurements for all the unseen conditions, with the largest improvement apparent for Vocoder SiB.

Results

Our aim was to test the hypothesis that speech understanding in noise is shaped by aspects of temporal-coherence processing that exist in early auditory areas. For this, we used a combination of online consonant identification experiments and computational modeling. In particular, we compared consonant confusion predictions from a model of across-channel temporal-coherence processing that is physiologically plausible in the cochlear nucleus with predictions from a purely within-channel model inspired by current speech-intelligibility models (see Scene analysis modeling to predict consonant confusions).

Figure 5 shows speech intelligibility measurements from the online consonant identification study. Approximately equal overall intelligibility was achieved for SiSSN, SiDCmod, SiB, and Vocoder SiB because of our careful choice of SNRs for these conditions based on pilotting (Table 1). This was done to obtain roughly equal variance in the consonant confusion estimates for these conditions, which allows us to fairly compare confusion patterns across them. Equalizing intelligibility also maximizes the statistical power for detecting differences in the pattern of confusions. ~60% overall intelligibility was obtained in each condition, which yielded a sufficient number of confusions for analysis.

The identification data collected in the online experiment were used to construct a consonant confusion matrix for each condition, then statistically significant differences in these matrices across conditions were extracted (see Quantifying confusion matrices from perceptual measurements and Statistical analysis). Results (Fig. 6) show significant differences in the confusion patterns across (1) conditions with different masker modulation statistics, and (2) stimuli with intact versus degraded TFS information. Computational modeling was then used to predict these differences across conditions to test our hypothesis about the role of temporal-coherence processing in scene analysis.

We constructed the following different models of scene analysis: (1) a within-channel model, which simulates masking of target-speech envelopes by distracting masker modulations within individual frequency channels, and (2) an across-channel model, which simulates across-channel temporal-coherence processing to account for interference from masker elements that are temporally coherent with target elements but in different frequency channels (see Scene analysis modeling to predict consonant confusions). We derived a separate neural confusion matrix for each model and listening condition. Then, each scene analysis model was separately calibrated by fitting a nonlinear mapping relating the neural consonant confusion matrix entries derived from that model for the SiSSN condition to corresponding perceptual measurements. Once fit, this mapping was used to quantitatively predict perceptual consonant confusions for novel conditions not used in calibration. Figure 7 shows results from the calibration step. In this figure, the different entries of the measured

equivalent in their predictive power, the individual frequency channels, and (2) an across-channel model, which simulates across-channel temporal-coherence processing to account for interference from masker elements that are temporally coherent with target elements but in different frequency channels (see Scene analysis modeling to predict consonant confusions). We derived a separate neural confusion matrix for each model and listening condition. Then, each scene analysis model was separately calibrated by fitting a nonlinear mapping relating the neural consonant confusion matrix entries derived from that model for the SiSSN condition to corresponding perceptual measurements. Once fit, this mapping was used to quantitatively predict perceptual consonant confusions for novel conditions not used in calibration. Figure 7 shows results from the calibration step. In this figure, the different entries of the measured
perceptual confusion matrix for SiSSN are plotted against the corresponding entries of the neural confusion matrix from each model for SiSSN. From this figure, it can be seen that the data show floor and ceiling effects, that is, as the neural metric increases (or decreases), the perceptual metric concomitantly increases (or decreases) but only up to a point, after which it saturates. This phenomenon is common to psychometric measurements. We fit this nonlinear relationship between the neural and perceptual data for SiSSN using a sigmoid/logistic function (Fig. 7; commonly used in the literature to obtain psychometric curves) separately for each model.

The model-specific mapping derived in the calibration step was used to predict perceptual consonant confusion matrices for each of the scene analysis models from the neural confusion matrices for unseen conditions (not used in calibration). Then, voicing, POA, and MOA confusion matrices were derived by pooling over all consonants (see Figs. 9, 10, 11). Finally, model predictions were compared with perceptual measurements for the different confusion matrix entries across the voicing, POA, and MOA categories. The results are shown in Figure 8 for SiB, SiDCmod, and Vocoder SiB. Visual comparison of the plotted data against the line of equality in Figure 8 suggests that there is a prediction bias for the SiDCmod and Vocoder SiB conditions for both the within- and across-channel models. This bias likely arises from our choice of calibration function (i.e., the sigmoid function) and the fact that calibration parameters were fitted to SiSSN, which may be suboptimal for the other conditions. Nonetheless, it can be seen that the cluster of points is less dispersed for the across-channel model compared to the within-channel model, indicating greater predictive accuracy for the across-channel model. The SiQuiet condition is not visualized, as there were ceiling effects in the intelligibility measurements (i.e., the diagonal entries of the confusion matrix were dominant) and very few confusions (i.e., off-diagonal entries were rare), which made it infeasible to meaningfully evaluate the quality of predictions for this condition (as there was no variance across either the on- or off-diagonal entries). But overall, across all entries for SiQuiet, both models predicted diagonal entries...
close to one and off-diagonal entries close to zero, in line with perceptual measurements. Pearson correlation coefficients were computed between the model predictions and perceptual measurements for the unseen conditions (Fig. 8) as well as for SiSSN (i.e., the calibration condition); the results are given in Tables 3 and 4 for the within- and across-channel models, respectively. Because the range of confusion matrix entries spanned three orders of magnitude, all comparisons were performed with log-transformed values. The correlations were statistically significant across all nonvocoded conditions for the within-channel model and across all conditions for the across-channel model (see Statistical analysis). The
strong correlation of the within-channel model predictions with perceptual data in the nonvocoded conditions (where TFS cues are preserved) provides independent evidence that speech understanding is strongly influenced by modulation masking when TFS cues are available (Viswanathan et al., 2021a); moreover, this result also suggests that modulations are used differently by the brain in the absence of natural TFS.

The across-channel model produced stronger correlation values compared to the within-channel model for all conditions, and the improvements were statistically significant across all conditions even after correcting for multiple comparisons (Table 5; see Statistical analysis). Thus, a simple physiologically plausible model of across-channel cochlear nucleus processing that shows CMR (Fig. 4) also yields category confusion predictions that match behavioral data and more specifically improves predictions compared to a within-channel model. Note that our within-channel model assumes perfect segregability of target-masker components that are separated in CF and MF (in line with current speech-intelligibility models; Jørgensen et al., 2013; Relaño-Iborra et al., 2016), and only models within-channel modulation masking. Specifically, within a particular channel (i.e., CF) and MF, masker modulations that are not in phase with the target are the only components that mask the target. However, our across-channel model simulates both within-channel modulation masking and cross-channel temporal-coherence-based interference. Specifically, masker components that are in a different channel from the target but that are temporally coherent with the target can interfere with target coding and perception. We implemented this interference via the CMR circuit model (Fig. 1), where temporally coherent pieces of the target and masker, even across distinct cochlear channels, coherently drive the wideband inhibitor, thereby enhancing outputs of the narrowband cell (which is inhibited by the wideband inhibitor) that are incoherent with the masker. Thus, our finding that model predictions are improved when cross-channel processing is added is consistent with the theory that across-channel temporal coherence shapes scene analysis (Elhilali et al., 2009). Moreover, this result also suggests that physiological computations that exist as early as the cochlear nucleus can contribute significantly to temporal-coherence-based scene analysis. Note that improvements to confusion predictions are apparent with the across-channel model for the same range of model parameters for which the CMR effect is also apparent.

Another key result from Table 5 is that the condition that showed the greatest improvement in confusion matrix predictions between the within- and across-channel models is Vocoded SiB. The masker in Vocoded SiB produces both within-channel modulation masking and cross-channel interference (as described above). These masking and interference effects are partially mitigated in SiB (and other nonvocoded conditions) compared to Vocoded SiB because the brain can use the pitch cue supplied by natural TFS to better separate the target and masker (Darwin, 1997; Oxenham and Simonson, 2009). The across-channel model is a better fit to perceptual data for all conditions, which suggests that cross-channel interference affects perceptual data. Thus, the improvement offered by this model is likely most apparent for Vocoded SiB because cross-channel interference effects contribute most to perception in this condition.

Note that while the main difference between the two scene analysis models tested in the current study is the exclusion/inclusion of cross-channel processing, another difference is that the within-channel model discards TFS, whereas the across-channel model uses the full simulated auditory-nerve output to drive the CMR circuit model. This raises the possibility that part of the
Figure 10. Full set of measured (top row) and model-predicted (middle, bottom rows) POA confusion matrices.

Figure 11. Full set of measured (top row) and model-predicted (middle, bottom rows) MOA confusion matrices.
improvement offered by the across-channel model could come simply from the inclusion of TFS information within each channel independently. To investigate whether the poorer performance of the within-channel model was partly because of discarding TFS, we reran the within-channel model by retaining the full auditory-nerve output (data not shown). We found that the predictions from the modified within-channel model were not significantly better than those of the original within-channel model. This confirms that the improvement in predictions given by the across-channel model comes largely from across-channel CMR effects, suggesting that categorical perception is sensitive to the temporal coherence across channels. Moreover, these CMR effects were restricted to low rates (<80 Hz or so; Fig. 4C), consistent with perceptual data (Carlyon et al., 1989). This suggests that the cross-channel processing did not benefit much from the TFS information included in driving the CMR circuit model.

For completeness, the full set of model-predicted and measured perceptual confusion matrices are shown for the voicing, POA, and MOA categories (Figs. 9, 10, 11); results are shown only for the SiB, SiDCmod, and Vocoder SiB conditions (i.e., the conditions unseen by the calibration step and having a sufficient number of confusions for prediction). In addition, the raw consonant confusion matrix measurements for all conditions are shown in Figure 12.

Discussion

To probe the contribution of temporal-coherence processing to speech understanding in noise, the present study used a behavioral experiment to measure consonant identification in different masking conditions in conjunction with physiologically plausible computational modeling. To the best of our knowledge, this is the first study to use confusion patterns in speech categorization to test theories of auditory scene analysis. The use of confusion data provides independent constraints on our understanding of scene analysis mechanisms beyond what overall intelligibility can provide. This is because percent correct data only convey binary information about whether target coding was intact, whereas consonant categorization and confusion data provide richer information about what sound elements received perceptual weighting.

We constructed computational models simulating (1) purely within-channel modulation masking (in line with current speech-intelligibility models; Relaño-Iborra et al., 2016), and (2) a combination of within-channel modulation masking and across-channel temporal-coherence processing mirroring physiological computations that are known to exist in the cochlear nucleus (Pressnitzer et al., 2001). Our across-channel temporal-coherence circuit produced a CMR effect (Fig. 4) that is consistent with actual cochlear nucleus data (Pressnitzer et al., 2001) and perceptual measurements (Mok et al., 2021). Moreover, consonant confusion pattern predictions were significantly improved for all tested conditions with the addition of this cross-channel processing (Table 5), which suggests that temporal-coherence processing strongly shapes speech categorization when listening in noise. This result is consistent with the theory that comodulated features of a sound source are perceptually grouped together and that masker elements that are temporally coherent with target speech but in a different channel from the target perceptually interfere (Schoonerveldt and Moore, 1987; Darwin, 1997; Apoux and Bacon, 2008). The only case where the within- and across-channel models were statistically equivalent was in predicting the off-diagonal entries (i.e., true confusions) for the SiDCmod condition; this may be because this condition has little coherent cross-channel interference from the masker since the masker is unmodulated (Stone et al., 2012).

An important difference between the cross- and within-channel masking simulated in our models is that while the cross-channel interference was produced by masker fluctuations that
were temporally coherent with the target, the within-channel masking was produced by masker components that were matched in both CF and MF with target components. While current speech-intelligibility models simulate the latter type of masking (Jørgensen et al., 2013; Relaño-Iborra et al., 2016), they do not account for cross-channel temporal-coherence-based masking as we have done here. This may explain why these models fail in certain conditions, including for vocoded stimuli (Steinmetzger et al., 2019). Indeed, even in the present study, although our within-channel masking model reasonably accounted for category confusions, it failed when TFS cues were unavailable (Table 3). One explanation for this is that because pitch-based masking release is poorer in the vocoded condition due to degraded TFS information (Oxenham and Simonson, 2009), the effects of cross-channel interference are more salient. This may also be the reason why the Vocoder SiB condition showed the greatest improvement in confusion pattern predictions after adding cross-channel processing (Table 5), which models these interference effects.

Although the lateral inhibition network used in Elhilali et al. (2003) bears some similarities to the across-channel CMR circuit model used in the current study, the CMR circuit model was explicitly based on physiological computations present in the cochlear nucleus and their CMR properties. Thus, another implication of the results of the present study is that physiological computations that exist as early as the cochlear nucleus can contribute significantly to temporal-coherence-based scene analysis. Such effects likely accumulate as we ascend along the auditory pathway (Elhilali et al., 2009; Teki et al., 2013; O’Sullivan et al., 2015). Indeed, scene analysis may in general be supported by a cascade of mechanisms throughout the auditory pathway, including both early-stage processing as well as cortical mechanisms. For example, there are circuits in the brainstem, midbrain, and cortex that exhibit sensitivity and selectivity to different spectrotemporal regularities in the input (Nelken et al., 1999; Pressnitzer et al., 2008; Kondo and Kashino, 2009; D[108]ienbrock et al., 2017; Mishra et al., 2021), these properties may in turn support auditory-object formation and scene segregation. Moreover, top-down cognitive processes such as attention can also contribute to scene analysis, especially when sound elements are otherwise perceptually similar (Shinn-Cunningham, 2020). Thus, future studies should explore the contributions of scene analysis mechanisms at different levels of the hierarchy of auditory processing.

The CMR circuit model used in the current study does not perform pitch-range temporal-coherence processing, and no CMR effect was seen at high modulation rates (Fig. 4C), consistent with perceptual data in the literature (Carlyon et al., 1989). Despite this, our across-channel model significantly improved predictions of category confusions compared to the within-channel model, which suggests that temporal-coherence processing at lower modulation rates is perceptually important. A future research direction is to extend the modeling framework proposed here to study the contributions of scene analysis mechanisms beyond the specific aspects of temporal-coherence processing studied here. One such extension could be to account for pitch-based source segregation (Bregman, 1990), perhaps by modeling a combined temporal-place code for pitch processing (Shamma and Klein, 2000; Oxenham et al., 2004; Oxenham and Simonson, 2009).

One limitation of the periphery model that we used (Bruce et al., 2018) is that it was developed to match nerve responses to simple stimuli. However, this family of periphery models has been successfully used to account for complex phenomena such as synchrony capture (Delgutte and Kiang, 1984), formant coding in the midbrain (Carney et al., 2015), and qualitative aspects of evoked potentials such as auditory brainstem responses and frequency-following responses (Shinn-Cunningham et al., 2013). Although a debate exists regarding the spatiotemporal properties of different periphery models in cochlear responses (Verhulst et al., 2015; Vecchi et al., 2021), those differences are subtle compared to the slower CMR effects that are important for the present study. A more general limitation of the models used in this study is that they are simple and do not incorporate many aspects of speech perception (e.g., context effects; Dubno and Levitt, 1981) because the goal here is to test specific theories of scene analysis. Nevertheless, the contrast between the models would be unaffected by these higher-order effects.

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