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1 2	Dimension-selective attention as a possible driver of dynamic,
3	context-dependent re-weighting in speech processing
4	Lori L. Holt 12
5	Adam T. Tierney <sup>34</sup>
6	Giada Guerra 34
7	Aeron Laffere
8	Frederic Dick 345
9 10	<sup>1</sup> Department of Psychology, Carnegie Mellon University, Pittsburgh, PA 15213
11	<sup>3</sup> Center for the Neural Basis of Cognition, Carnegie Mellon University, Pittsburgh, PA 15213
12	<sup>3</sup> Department of Psychological Sciences, Birkbeck College, University of London, London, WC1E 7HX
13	Centre for Brain and Cognitive Development, Birkbeck College, London, WC1E 7HX
14	<sup>3</sup> Department of Experimental Psychology, University College London, London, WC1H 0AP
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17	Corresponding Author:
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20	Lori L. Holt Professor, Department of Psychology
21	Carnegie Mellon University
22	5000 Forbes Avenue
23	Pittsburgh, PA 15213
24	loriholt@cmu.edu
25	Highlights:
26	Inginigitis.
27 28	Speech processing requires continuous reweighting across many acoustic dimensions
29	• This dynamic mapping may reflect the dynamics of auditory attentional mechanisms
30	• Animal neurobiological models can help to determine the putative role for attention
31	• We present results from a new attentional paradigm that ties together human and non-human research
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### Abstract

The contribution of acoustic dimensions to an auditory percept is dynamically adjusted and reweighted based on prior experience about how informative these dimensions are across the long-term and short-term environment. This is especially evident in speech perception, where listeners differentially weight information across multiple acoustic dimensions, and use this information selectively to update expectations about future sounds. The dynamic and selective adjustment of how acoustic input dimensions contribute to perception has made it tempting to conceive of this as a form of non-spatial auditory selective attention. Here, we review several human speech perception phenomena that might be consistent with auditory selective attention although, as of yet, the literature does not definitively support a mechanistic tie. We relate these human perceptual phenomena to illustrative nonhuman animal neurobiological findings that offer informative guideposts in how to test mechanistic connections. We next present a novel empirical approach that can serve as a methodological bridge from human research to animal neurobiological studies. Finally, we describe four preliminary results that demonstrate its utility in advancing understanding of human non-spatial dimension-based auditory selective attention. 

### 113 **1.1 Introduction**

Understanding a friend's speech in a crowded cafe, tracking the quality of a sick child's breathing through a 114 nursery monitor, and following the melody of a violin within an orchestra all require extracting the most 115 informative dimensions for the task at hand from a complex mix of acoustic signals. Each of these scenarios can 116 be conceived of as a variant of the classic 'cocktail party effect' (Cherry, 1953), whereby selective and sustained 117 endogenous attention is directed to a particular sound source as it evolves in time. Experimental paradigms 118 modeling these 'cocktail party' scenarios have often examined the contribution of acoustic dimensions 119 conveying sound sources' spatial position in disambiguating target signals from irrelevant background scenes. 120 This is appropriate given the importance of detecting and orienting to acoustic events in space. Yet, listeners 121 must employ selective auditory attention even when spatial cues are unavailable (such as over the telephone or 122 in listening to an orchestral recording over earbuds) or unreliable (as in listening within reverberant 123 environments). 124

Indeed, even in the absence of spatial cues, listeners appear to dynamically adjust and selectively weight the 125 contribution of multiple acoustic dimensions to an auditory percept based on prior experience about how 126 informative these dimensions are - individually and in concert – to behavior. Speech perception provides an 127 excellent case-in-point because individual speech sounds, phonemes like /b/ and /p/, are defined across 128 multiple acoustic dimensions. Typically, no one acoustic dimension is necessary or sufficient to unambiguously 129 signal a phoneme. Even more, factors like long-term and short-term acoustic distributional regularities and the 130 adjacent sound context can impact the effectiveness of specific acoustic dimensions in signaling speech sounds. 131 Even for a well-learned auditory skill like speech perception, the mapping of acoustics to percept remains 132 flexible. The dynamic and selective adjustment of how robustly different acoustic dimensions contribute to speech 133 recognition has made it tempting to conceive of this as a form of non-spatial auditory selective attention.

Our aim in this review is to explore this possibility. We first review some general background in the mapping of acoustics to speech. We next describe several speech perception phenomena to illustrate the highly dynamic nature of the mapping from acoustics to phoneme. We discuss how each of these phenomena resonates with a colloquial understanding of selective attention. But, we caution that it is important to recognize that *attention* may be best thought of as a cognitive placeholder that does not, in and of itself, point to a specific neurobiological mechanism (e.g., Cohen, Romero, Servan-Schreiber, & Farah, 1994).

We demonstrate this point by relating the human speech phenomena to illustrative neurobiological findings from nonhuman animal models. The neurobiological work offers informative guideposts in how we might make mechanistic connections back to human speech recognition. More specifically, it suggests that it would be unwise to be wholly satisfied with characterization of these speech phenomena as selective attention. There remains more explanatory work to be done, as a constellation of candidate neurobiological mechanisms exist that may support the dynamic nature of mapping acoustic input to behaviorally-relevant sounds like speech.

146 But, how might we make progress in advancing dimension-based selective attention from a cognitive 147 placeholder to a real mechanistic understanding of human auditory behavior, including speech perception? 148 After all, there remains a substantial distance between speech perception and approaches from nonhuman 149 animal neurobiology. In the final section of the paper, we outline a novel empirical approach to human 150 dimension-based selective attention that may serve as a methodological bridge between human and nonhuman 151 animal literatures. By more closely aligning human experimental approaches with those that have been 152 successful in nonhuman animal neurobiology, it may be possible to draw from the vital interpretive frameworks 153 provided by neurobiological research. We briefly describe four empirical results to demonstrate the utility of 154 this approach in advancing understanding of human dimension-based auditory selective attention with the ultimate aim of achieving a more nuanced model of the multiple mechanisms potentially at play in phenomena 155 156 for which we use dimension-based selective attention as a cognitive placeholder.

## <sup>157</sup> 1.2 Examples from Speech Processing

158 To situate our examples, it is useful to begin with some common background in speech acoustics. Consider the 159 simple act of deciding whether your conversation partner has uttered /b/ or /p/, as in *beach* versus *peach*. If you 160 know of one acoustic dimension related to speech communication, there is a very good chance it is voice onset 161 time (VOT). The superstar of acoustic speech dimensions, VOT is defined in articulatory terms as the length of 162 time between the release of a stop-consonant like /b/or/p/and the onset of voicing, the vibration of the vocal 163 folds (Stevens, 2000). If you hold your fingers to your larynx while uttering *beach* and *peach* you will notice that 164 the delay from when your lips release the consonant and your vocal folds being to vibrate is a bit longer for the 165 'voiceless' consonant /p/ than the 'voiced' consonant /b/. This has multiple acoustic outcomes (Lisker, 1986). 166

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Chief among them, there is a greater temporal lag from the acoustic release burst associated with opening the mouth and the onset of a periodic acoustic signal originating from vibration of the vocal folds. Accordingly, it is rather easy to morph from voiced to voiceless consonants by parametrically lengthening this delay to create a series of speech sounds varying across VOT. At least in part as a result of this ease, the significance of VOT as an acoustic dimension in signaling voicing category distinctions like /b/-/p/, /d/-/t/, and /g/-/k/ has been studied across 100s, perhaps 1000s, of experiments spanning many languages (Abramson & Whalen, 2017).

Recent neurobiological research has very elegantly demonstrated that it is possible to recover a voicing code in human superior temporal cortex (Mesgarani, Cheung, Johnson, & Chang, 2014). These very exciting results can give the impression that we have discovered the neural code that supports categorization of an utterance as *beach* or *peach*. And, according to the classic 'textbook' understanding of the mapping from acoustics to phonetic categories, this would be true. But, contemporary research on the mapping of complex speech acoustics to phonetic categories makes clear that this textbook understanding is in need of an update. The situation is, in fact, more complex.

### 181 182 **1.2.1** The Textbook Understanding of Speech Processing, With Contemporary Updates

183 To situate the examples that follow below, it is important to recognize that theory and research directed at human speech processing have long grappled with the issue of how the complex acoustic dimensions that vary 184 across speech signals relate to phonemes, the linguistically distinct units of sound that differentiate meaning in 185 a language such as /b/ versus /p/ in *beach* versus *peach*. Chances are very good that in opening an introductory 186 perception or cognition textbook you will find a figure characterizing the *categorical perception* of speech (e.g., 187 Wolfe, Kluender, Levi, Bartoshuk, & Herz, 2015). Perhaps the best-known phenomenon of speech perception 188 (often demonstrated across the superstar dimension, VOT), categorical perception refers to the observation that 189 listeners' identification of speech sounds does not vary gradually across incremental changes in an acoustic 190 speech dimension. Instead, there is an abrupt shift across a restricted range of acoustic change. Endpoint stimuli 191 are identified as one phoneme with near-ceiling performance that transitions sharply to near-ceiling 192 identification of another phoneme. This categorical response appeared to be consistent with a mapping of speech 193 acoustics to discrete, symbolic phonemic representations (Liberman, Harris, Hoffman, & Griffith, 1957). By this 194 view, the subtle details of acoustic dimensions are unavailable once they are mapped discretely to a phoneme. 195 Additionally, this view emphasized the mapping of individual dimensions to phonemes, as in VOT to  $\frac{b}{-p}$ , 196 and led to a long (and ultimately somewhat fruitless, (Blumstein & Stevens, 1985; Lisker, 1985) search for 197 invariant acoustic cues that map to phonemes.

198 Contemporary research suggests that it is more productive to characterize speech as *categorized* rather than 199 *categorical* (Holt & Lotto, 2010). The mapping looks much less discrete when speech perception is studied using 200 more continuous methods. Listeners consistently rate some speech instances as 'better' exemplars of a speech 201 category than others (e.g., Iverson & Kuhl, 1995; Utman, 1998; Utman, Blumstein, & Sullivan, 2001). Eyetracking 202 and graded electroencephalographic (EEG) responses further reveal that fine-grained acoustic details of an utterance affect its categorization (e.g., Aydelott & Bates, 2004; Mcmurray, Aslin, Tanenhaus, Spivey, & Subik, 203 2008; Mcmurray, Tanenhaus, & Aslin, 2002; Utman, 1998; Utman et al., 2001; Utman, Blumstein, & Burton, 2000) 204 and memory (e.g., Bradlow, Nygaard, & Pisoni, 1999; Goldinger, 1996; Nygaard, Sommers, & Pisoni, 1995). 205 When we move away from binary responses typical of categorical perception tasks (did you hear beach or peach?), 206 behavior suggests a rich internal structure in the representation of phonemes. Today, it is much more common 207 to conceptualize the mapping from acoustics to *perceptual phonetic categories* that are neither discrete nor symbolic 208 and instead possess rich internal structure that reflects the distributional characteristics of the experience that 209 drove category learning (Holt & Lotto, 2010; Holt, Lotto, & Kluender, 2000). 210

211 By this more contemporary perspective, there is no need to search for an invariant acoustic cue uniquely differentiating a particular phonemic contrast. Instead, phonetic categories can be considered to reside in a 212 highly multidimensional perceptual space that maps the acoustic complexity of speech across multiple 213 dimensions. Correspondingly, there is increasing appreciation that it is critical to consider *auditory* rather than 214 acoustic dimensions (like the manipulation leading to a step-wise VOT stimulus series), in appreciation of the 215 important transformations in early auditory processing that warp the perceptual space conveyed by acoustic 216 dimensions. (It is a somewhat ironic aside that some of the best evidence for nonlinearities in the mapping of 217 acoustic speech dimensions to auditory dimensions comes from the superstar of acoustic dimensions driving so 218 much research, VOT; Holt, Lotto, & Diehl, 2004). 219

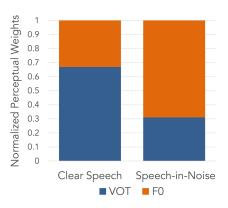
Finally, and most critically for the present review, contemporary research is rich with examples that even these auditory dimensions do not stably map to phonetic categories. Instead, the mapping is a much more dynamic process, indicating that the textbook understanding of an invariant, or even consistent, mapping from acoustics

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to speech is in need of an update, and that we should take care in concluding that neural decoding in human cortex conveys the complete mechanistic basis of human speech recognition. In the next sections, we consider some specific examples, and how they might be related to short-term plasticity and attentional modulation as well as longer-term learning about the informational environment.

### 229 1.2.1 Perceptual Weight in Speech Categorization

230 As central (and well-studied) as VOT is in signaling voicing categories in speech, there is in fact a constellation 231 of as many as 16 acoustic dimensions that co-vary with English /b/-/p/ category membership (Lisker, 1986). 232 For example, in addition to VOT, the fundamental frequency (F0, associated with voice pitch) of the following 233 vowel co-varies with  $\frac{b}{-p}$  category membership. When we utter *peach*, the following vowel tends to have a somewhat higher F0 than when we utter *beach*. Correspondingly, listeners rely upon both dimensions in 234 phonetic categorization. When VOT is acoustically ambiguous, for example, utterances with higher F0 235 frequencies are categorized as /p/ whereas those with lower F0 frequencies are categorized as /b/. Critically, 236 listeners do not rely upon these dimensions in equal measure. Rather, behavioral (Francis, Kaganovich, & 237 Driscoll-Huber, 2008; Holt & Lotto, 2006; Iverson & Kuhl, 1995), neural (Scharinger, Herrmann, Nierhaus, & 238 Obleser, 2014), and developmental (Nittrouer, Lowenstein, & Packer, 2009; Wellmann, Holzgrefe, Truckenbrodt, 239 Wartenburger, & Höhle, 2012) evidence indicates that listeners *perceptually weight* acoustic dimensions, with 240 some dimensions contributing more robustly to perception than others. 241



254 Figure 1. Listeners categorized a grid of speech 255 sounds varying across VOT and F0 as /b/ or /p/. The relationship of each acoustic dimension to 256 categorization responses was calculated using 257 regression, with normalized regression weights 258 providing a measure of perceptual weight. In Clear Speech. VOT is most diagnostic of /b/-/p/ 259 categorization. But, in white noise, F0 dominates 260 categorization of the same speech sounds.

As we will discuss in more detail below, prior research demonstrates that perceptual weights are a function of the long-term statistics of the input (Francis, Ciocca, Wong, Leung, & Chu, 2006; Holt & Lotto, 2006; Toscano & Mcmurray, 2010), they are specific to one's native language (Iverson et al., 2003; Kondaurova & Francis, 2008; 2010), and they emerge over a rather protracted developmental timeline extending at least into late childhood (Idemaru & Holt, 2013). For present purposes, the point is simply that although multiple auditory dimensions signal phonetic category identity, their contributions are not equivalent. Some dimensions carry greater perceptual weight than others. Figure 1 illustrates this for /b/ versus /p/ categorization, where perceptual weight is calculated as the normalized regression coefficient related to /b/-/p/ categorization by native-English listeners across a grid of speech syllables varying parametrically in VOT and F0. For clear speech (Figure 1, Clear Speech), both acoustic dimensions inform  $\frac{b}{-p}$ categorization, but VOT carries greater perceptual weight. It better predicts how listeners will categorize a sound than F0.

Many investigators have noted the potential for selective attention to play a role in perceptual weighting of acoustic dimensions in speech processing, in the sense that selective attention appears to be consistent with the demand to direct processing to diagnostic dimensions in the

presence of the rich acoustic information available across multiple input dimensions (Francis & Nusbaum, 2002; 262 Gordon, Eberhardt, & Rueckl, 1993; Heald & Nusbaum, 2014). This conceptualization suggests a potentially 263 dynamic process, one not rigidly wired at the culmination of development. Figure 1 provides a simple example 264 that underscores this point. When the same  $\frac{b}{-p}$  stimuli used to calculate perceptual weights across VOT 265 and F0 in clear speech (Figure 1, Clear Speech) are presented in modest levels of white noise, *categorization is* 266 more dependent on F0 and less dependent on VOT (Figure 1, Speech-in-Noise). We can speculate that the F0 267 dimension may be more robust to noise and therefore a more valuable indicator of phonetic category identity 268 under noisy conditions. Whether accomplished by processes consistent with 'attention' or through other means, 269 the shift in perceptual weights apparent in Figure 1 makes it clear that listeners rely on different acoustic 270 dimensions in speech categorization in adverse versus clear listening environments. Perceptual weights are 271 labile. VOT is the star dimension focused upon in textbook examples, but it only shines under the right 272 circumstances.

This compelling example is not a mere parlor trick of perception. It informs us that discovering a neural code for VOT, or any other acoustic dimension that informs speech perception, takes us only part of the way to understanding how the auditory system maps complex acoustics to objects for recognition. A complete account will require a deeper understanding of how acoustic dimensions are weighted in auditory recognition because

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the very dimensions that inform auditory object recognition are not fixed. Rather, listeners flexibly shift reliance
 on acoustic dimensions according to the demands of the listening environment.

### 283 1.2.2 Perceptual Learning Over the Long-term

284 Although speech category learning gets underway even before an infant's first birthday (Conboy & Kuhl, 2011; 285 Kuhl, 2004) there is a long developmental tail that extends into at least early adolescence in establishing the 286 perceptual weights of acoustic dimensions (Zevin, 2012). For example, the onset frequency of the third formant 287 (F3) is the acoustic dimension that best predicts English /r/-/1/ category membership in the acoustics of native 288 talkers' speech (Iverson et al., 2003), although the onset frequency of the second formant (F2) is also diagnostic 289 to a lesser degree. Among mature listeners, these distributional regularities of English speech input are reflected in /r/-/1/ perceptual categorization. Adult listeners rely more on F3 onset frequency, giving it greater 290 perceptual weight, than F2 onset frequency. But, although typically-developing native-English-learning children 291 ages 4.5, 5.5, and even 8.5 years use the dominant, F3, input dimension to accurately categorize English /r/-/l/, 292 they fail to rely upon F2 as a secondary diagnostic dimension like adults (Idemaru & Holt, 2013). This indicates 293 a much longer developmental course for phonetic category development than is typically appreciated (Zevin, 294 2012). 295

Moreover, this pattern of development underscores the fact that perceptual weighting arises, at least in part, 296 from dimensions' informativeness in signaling category identity (Holt & Lotto, 2006; McMurray & Jongman, 297 2011). The distributional regularities of speech input shape perceptual weight of input dimensions. Efficient 298 categorizers ultimately learn to perceptually weight the multiple dimensions that define speech categories in 299 relation to the dimensions' reliability, or informativeness, in signaling a category (Holt & Lotto, 2006). 300 Additionally, perceptual weight is likely to be impacted additionally by basic auditory representation (some 301 dimensions are more robustly encoded by the auditory system than others) and even task (dimensions heavily 302 weighted for phonetic categorization may be much less relied upon in identifying a talker). Either of these latter 303 factors may play a role, as well, in the perceptual weight shifts evident in Figure 1. Accordingly, some accounts 304 have emphasized learning to attend selectively to diagnostic dimensions as an important component of phonetic 305 category learning (Heald & Nusbaum, 2014; Kondaurova & Francis, 2010). (Attention-based approaches to 306 category learning and warping have long been used in vision research, e.g., Kruschke, Kappenman, & Hetrick, 307 2005; Nosofsky, 1986).

308 If efficient speech comprehension heavily relies on the process of learning and maintaining representations of 309 higher-dimensional auditory categories, then one might expect that localized patterns of neural activation 310 related to speech processing (or, indeed, seemingly *selectively* related to speech) might also be associated with 311 the emergence of new nonspeech auditory categories. As a test of this hypothesis, Leech, Holt, Devlin, and Dick 312 (2009) asked whether video game play that drives implicit nonspeech auditory categorization (Wade & Holt, 313 2005) would change responses to the trained nonspeech sounds in canonical 'speech-selective' cortex. They 314 found that subjects' ability to categorize these novel sounds after training was significantly correlated with pre-315 to-post training change in fMRI activation in a part of the left posterior superior temporal sulcus that has been implicated in speech processing and phonemic categorization (Dehaene-Lambertz et al., 2005; Desai, Liebenthal, 316 Waldron, & Binder, 2008). 317

318 Studying how adult listeners learn artificial, nonspeech auditory categories has informed thinking because it is 319 difficult to gain an experimental foothold in understanding how learning operates over long-term speech 320 category development since direct manipulation of children's speech input is infeasible. As adults learn novel, 321 artificial auditory categories they must learn to pull together auditory dimensions according to training-related 322 task demands and feedback to form new representations. Learning new auditory categories that generalize to 323 novel instances changes the partitioning of auditory representational space (Liu & Holt, 2011) in a manner that can be described as 'warping' or exaggeration of the mapping of input to emphasize categorization-relevant 324 acoustic dimensions, or alternatively as plasticity that directs selective attention to these dimensions. Indeed, 325 provided with appropriate training, listeners can learn to attend selectively to acoustic dimensions that do not 326 typically contribute to native-language speech perception (Kondaurova & Francis, 2010), and this impacts 327 electrophysiological response to speech (Ylinen et al., 2010). The observations potentially argue for construing 328 perceptual learning of auditory (including speech) categories over the long-term as involving allocation of 329 selective attention to the most diagnostic acoustic dimensions. 330

### 331 **1.2.3 Perceptual Learning Across the Short-term**

The challenge for human communication is even greater because we often encounter talkers with foreign accents, dialects, or speech idiosyncrasies. In these cases, the speech input is 'warped' relative to the pattern of

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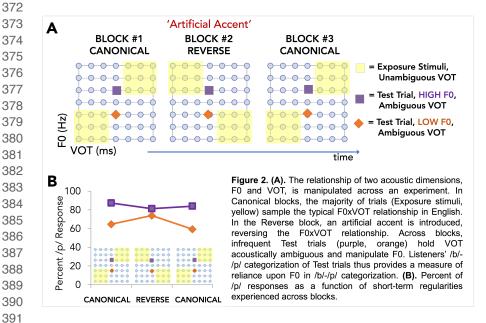
experience that established the long-term perceptual weights, with the potential for acoustic input dimensions
 to relate differently to phonetic categories.

This challenge is met by a highly flexible perceptual system capable of tracking short-term input regularities and 339 dynamically adapting reliance upon specific acoustic dimensions. Recall, from above, that both VOT and F0 340 contribute to English /b/-/p/ categorization, with VOT more diagnostic than F0 in clear speech. These 341 dimensions are also correlated in English speech productions. Stimuli with longer VOT, typical of p/, also tend 342 to have higher F0 frequencies whereas those with shorter VOT, typical of /b/, are associated with lower F0 343 frequencies. Mature listeners are sensitive to this relationship. When VOT is acoustically ambiguous and 344 insufficient to reliably signal /b/versus /p/, listeners label higher-F0 stimuli as /p/versus /p/versu345 /b/. 346

It is possible to model real-world encounters with foreign-accented speech by manipulating the short-term 347 distribution of speech experience across an experiment. For example, Idemaru and Holt (2011) had listeners 348 categorize speech sounds as beer or pier with a button press. The majority of trials were 'exposure' trials in which 349 the speech exemplars were unambiguously signaled by the dominant perceptual dimension (VOT) and the 350 secondary dimension (F0) was correlated in the canonical manner (Figure 2a). This conveyed a short-term 351 distribution of speech experience that aligned with the long-term regularities of English. Without a change in 352 task or other overt cues, Idemaru and Holt introduced a subtle 'artificial accent' by shifting the distribution 353 statistics between VOT and F0 acoustic dimensions. In the Reverse Block shown in Figure 2a, VOT continued to 354 unambiguously signal category membership across the exposure trials. But the secondary, F0, dimension was 355 now associated with the VOT dimension in manner counter to long-term English experience. In the Reverse 356 block, shorter VOTs were associated with higher F0s and longer VOTs were associated with lower F0s. This 357 produced an artificial accent that changed the short-term input regularities in a manner akin to some natural 358 foreign-language accents (Kim & Lotto, 2002).

359 Idemaru and Holt (2011; 2014) assessed the impact of this shift in short-term regularities across speech input 360 dimensions by observing overt categorization decisions across infrequent 'test' trials intermixed with the 361 exposure trials (orange diamond and purple square symbols, Figure 2). For these two stimuli, the dominant 362 dimension, VOT, was acoustically ambiguous and therefore provided poor information about phonetic category 363 identity. As a result, the two test stimuli differed only in the secondary, F0, dimension. As such, categorization 364 of the test stimuli provided a metric of the perceptual weight of F0 – how diagnostic the F0 dimension is in 365 signaling  $\frac{b}{-p}$  categorization. If listeners rely exclusively on VOT, categorization of the two test stimuli will not differ. But, to the extent that F0 informs category membership, categorization of the two test stimuli will 366 differ. The magnitude of this difference provides a measure of the perceptual weight of F0 in  $\frac{b}{-p}$ 367 categorization as a function of the short-term speech regularities manipulated across blocks via the exposure 368 stimuli. 369

370 It is important to point out that there was no explicit training or feedback. Listeners were not informed about 371 the shift in input from the Canonical to the Reverse block, the talker remained constant, the test trials were not



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differentiated from the exposure trials, and the task was always simply to identify the word. The range of dimension variability experienced blocks fell within across that experienced for the talker, and it went largely unnoticed by participants. Moreover, the range of values experienced across dimensions was constant across the experiment (only relationship the changed), so variability across a dimension was not a factor.

Figure 2b illustrates the impact of short-term regularities across speech input dimensions on listeners' reliance on F0 to signal /b/-/p/ categorization. When short-term input aligns with native-language

experience (Canonical Blocks), listeners relied upon the secondary, F0, dimension to make category decisions. It
 provided information across which to differentially categorize the test stimuli as /b/ (Low F0) and /p/ (High
 F0). This is simply a reflection of the fact that secondary dimensions informed categorization, albeit less robustly
 than the dominant dimensions.

However, upon introduction of the artificial accent in the Reverse block – a short-term change in input 397 regularities – reliance upon F0 to inform /b/-/p/ categorization was rapidly down-weighted (Figure 2b). When 398 the short-term input shifted such that F0 mapped to VOT in a manner inconsistent with long-term speech input 399 regularities, the F0 dimension was no longer as informative to  $\frac{b}{-p}$  categorization. Note that the down-400 weighting of F0 in informing speech categorization does not appear to reflect a wholesale shift in attention away 401 from the secondary, F0, dimension; listeners rapidly resumed reliance F0 in a final Canonical Block, indicating 402 that they continued to track F0 in the input. Rather, the data suggest a continuous, dynamic modulation of input 403 dimensions' contributions to phonetic categorization, adjusted to accommodate short-term input regularities. 404

On the face of it, this dynamic adjustment in the weighting functions with which auditory dimensions map to phonetic categories could be described as consistent with rapid adjustments in selective attention to auditory dimensions. However, our currently incomplete understanding of human auditory selective attention makes it difficult to determine definitively whether this is a viable model. The conundrum for advancing a mechanistic understanding of whether selective attention plays a role is that we do not yet have a rich body of evidence regarding the boundaries and constraints of dimension-based auditory selective attention to definitively determine whether it is playing a role. Even so, these behavioral results highlight the inherently dynamic nature of the mapping from acoustic speech input to behaviorally-relevant categories like phonemes and words.

### 412 413 **1.2.4 The Impact of Context in Speech Categorization**

414 Even quite subtle changes in distributions of sound experienced across a single input dimension can influence how an acoustic input dimension factors into phonetic categorization. To illustrate, consider categorization of 415 speech syllables that vary perceptually from /ga/ to /da/. In English speech productions, these syllables are 416 best differentiated by the third formant onset frequency (F3). Accordingly, F3 onset frequency carries a strong 417 perceptual weight in /ga/-/da/ categorization. As is typical in identification responses across a series of stop 418 consonants like /g/ and /d/, there is a rather steep slope in the transition from identifying lower F3 onsets as 419 /ga/ to identifying higher F3 onsets as /da/. This steep identification function (consistent with what is 420 traditionally interpreted as categorical perception) invites the inference that a specific range of lower-frequency 421 range of F3 onset frequencies map to /g/ and another specific, higher-frequency, range of F3 onset frequencies 422 map to /d/(e.g., Lotto & Kluender, 1998). 423

However, perception of isolated syllables only tells part of the story. Consider what happens when a simple 424 sentence precedes the /ga/-/da/ syllables. As first demonstrated long ago (Ladefoged & Broadbent, 1957), 425 preceding context can have a substantial influence on speech categorization. A contemporary example 426 demonstrates this for /ga/-/da/ (Laing, Liu, Lotto, & Holt, 2012). In this study, a precursor phrase (Please say 427 what this word is) preceded the /ga/-/da/ syllables varying in F3 onset frequency and listeners simply 428 categorized the final /ga/-/da/ syllable. In one block of trials, the precursor phrase was manipulated to subtly 429 emphasize somewhat higher frequencies in a frequency band in the range of /ga/-/da/F3 onset frequencies. 430 In another block of trials, the same precursor phrase emphasized lower frequencies in the same band. Said 431 another way, the phrases subtly differed in the long-term average spectrum of the preceding speech. On each 432 trial, listeners simply reported whether they heard ga or da in the context of one of the two precursor phrases.

The results demonstrate that phonetic perception is influenced by the long-term average spectrum of precursor sounds. In the context of a precursor sentence with exaggerated higher F3-band frequencies, the mapping of F3 onset frequency shifts to result in more /ga/ categorizations. In the context of exaggerated lower F3-band frequencies in the precursor phrase, the same speech target syllables are more often categorized as /da/. Thus, a precursor can have a substantial effect on how the F3 onset frequency input dimension maps to phonetic categories. This may provide a means by which the system accomplishes talker normalization (Assgari & Stilp, 2015; Huang & Holt, 2012; Ladefoged & Broadbent, 1957).

Perhaps more surprising, precursor contexts across which spectrally-biased long-term average spectra emerge need not be speech to impact phonetic categorization (Holt, 2005; 2006b). When a series of pure tones sampling the higher versus lower F3-band frequencies precedes /ga/-/da/ syllables, phonetic categorization is also shifted (Holt, 2005; 2006b; Laing et al., 2012). In the context of a sequence of higher-frequency tones, categorization shifts to /ga/. The same speech syllables are more often reported as /da/ when preceded by a lower-frequency sequences of tones. Here, as in the case of speech precursor sentences, the direction of the

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influence of context is *spectrally contrastive*. Higher-frequency precursors lead subsequent acoustic information 449 to be more often mapped to the category characterized by lower F3 onset frequencies, /ga/, and vice versa. This 450 pattern of spectral contrast has been observed across many speech categories (Lotto & Holt, 2006), evoked by 451 precursor sentences (Assgari & Stilp, 2015; Huang & Holt, 2012; Laing et al., 2012), single syllables (Huang & 452 Holt, 2012; Lotto & Kluender, 1998), and across nonspeech contexts varying from tones to notched noise (Holt, 453 2005; 2006a; 2006b; Lotto & Kluender, 1998). Across these studies, the findings are consistent in revealing that 454 the mapping of an input dimension to an auditory representation, here a phonetic category, is not fixed. Rather, 455 the auditory system appears to track the distribution of spectral energy evolving across the long-term average 456 spectrum of incoming speech and the mapping of subsequent acoustic information is relative to, and 457 contrastively with, the distribution of acoustic information experienced in prior context.

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A rather spectacular non-speech demonstration of such acoustic context effects was recently reported by 459 Chambers et al. (2017). The authors took advantage of a classic auditory stimulus, a Shepherd tone, made of 460 octave-separated pure tones distributed across all audible frequencies. If one sequentially presents two Shepherd 461 tones separated by a base frequency of 6 semitones (a musical 'tritone'), the average listener is equally likely to 462 hear a pitch shift going up or going down (although individual listeners can have quite strong and stable bias for hearing an up or down shift). However, when such a Shepherd tone pair is preceded by an acoustic context, 463 subjects' perception of the direction of this ambiguous pitch shift could be quasi-deterministically manipulated, 464 whereby the contiguity of the separate frequency elements of the context tones with the two test tones decides 465 the percept. This result shows that a basic auditory perceptual decision - the direction of a local pitch contour -466 is strongly driven by active integration with prior acoustic information. 467

468 Although speculative, these demonstrations from human behavior may be consistent with accounts of auditory 469 selective attention that emphasize optimization of auditory cortical filters for task performance and for 470 enhancing selectivity to task-relevant information via contrast enhancement (e.g., Fritz et al., 2007; Jääskeläinen et al., 2007; 2011). 471

#### 472 1.2.5 Summary 473

We began with something simple: how might the auditory system recognize a spoken word *beach* from *peach*. 474 The textbook answer to this question is straightforward and has influenced our approach to evaluating 475 neurobiological evidence for speech recognition. The traditional understanding is that the system recognizes a 476 diagnostic auditory cue, like VOT, which maps to a phonetic category. By this view, it is quite natural to conceive 477 of the mapping from input to auditory object, in the cases above phonetic categories, as examples of sensory 478 'encoding' to relatively stable features or dimensions. Thus, when we see patterns of activation in the brain that 479 correspond closely with acoustic dimensions we know to be significant in signaling a particular phonetic 480 category (Mesgarani et al., 2014) it is tempting to conclude that we have cracked the speech code. 481

The phenomena reviewed above collectively illustrate the dynamic nature of the mapping of auditory 482 dimensions to behaviorally-relevant representations and actions. They reveal the need for a less static 483 perspective on how input is mapped to behaviorally-relevant auditory representations and highlight that the 484 dividing lines between perception, attention and learning are likely to be quite blurry -- if they exist at all. The 485 very mapping of acoustics to auditory dimensions and objects is dependent upon an organism's prior history of 486 experience, the short-term experience evolving in the local input, and statistical relationships relating the present 487 sound exemplar to those experienced previously.

488 These effects are well illustrated by perception of speech, but they are not exclusive to speech. Humans and 489 other mammals are very sensitive to changes in the salience, task-relevance, and composition of the acoustic 490 dimensions of complex and ecologically important sounds (Holt & Lotto, 2006; Leech et al., 2009b; Leech, Gygi, 491 Aydelott, & Dick, 2009a; Shamma & Fritz, 2014). Indeed, listeners appear to be able to shift attention across 492 multiple simultaneously-present acoustic dimensions to home in on the ones that are diagnostic in guiding 493 behavior (Henry, Herrmann, & Obleser, 2015; Herrmann, Henry, & Obleser, 2013a; Herrmann, Henry, 494 Scharinger, & Obleser, 2013b; Herrmann, Schlichting, & Obleser, 2013c; Idemaru & Holt, 2011). As we noted 495 above, this non-spatial dimension-based auditory attention has received rather little empirical study in human auditory cognitive neuroscience. Thus, although there are suggestive connections of the phenomena reviewed 496 497 above with attention, and although selective attention has been evoked as a potential contributor to the highly dynamic mapping of input in speech perception, it remains the case that explanatory power is compromised 498 without a more solid mechanistic understanding of non-spatial dimension-based attention in auditory 499 processing. 500

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To illustrate this point, we next briefly review several illustrative nonhuman animal studies of auditory processing that provide potentially useful guideposts in making headway on this issue. Collectively, they illustrate that although 'attention' is useful as a placeholder, the phenomena to which it is directed are unlikely to arise from a single mechanism, or across a constant level of representation or timescale. These illustrative examples also offer direction in considering how to build new human paradigms that can connect better with open questions about whether auditory selective attention – and plasticity associated with it – play a substantive role in the dynamic mapping of acoustic dimensions to speech reviewed above.

## 511 512 1.3 Nonhuman Animal Neurobiological Studies

The neural mechanisms of active listening (in contrast to passive 'hearing') have been increasingly the focus of 513 research in understanding the hierarchy of cortical areas identified in the mammalian auditory system (Hackett, 514 2011; Morillon, Hackett, Kajikawa, & Schroeder, 2015). Nonhuman mammal studies have shown that behavioral 515 manipulation of attentional systems can modulate, and even alter, the topography of tonotopic maps (Bieszczad 516 & Weinberger, 2010; Weinberger, 2007), and that this modulation is important for learning. Moreover, recent 517 studies demonstrate that neuronal receptive fields in regions along the cortical hierarchy are modulated in 518 response to the behavioral relevance of auditory dimensions (Atiani et al., 2014; David, Fritz, & Shamma, 2012; 519 Shamma & Fritz, 2014; Winkowski, Bandyopadhyay, Shamma, & Kanold, 2013; Yin, Fritz, & Shamma, 2014). We 520 briefly (and selectively) review a few illustrative examples that may be useful in connecting animal 521 neurobiological frameworks with phenomena we reviewed above. 522

<u>Perceptual Weighting</u>. Multiple species exhibit sensitivity to combinations of acoustic input dimensions (e.g., 523 Atencio, Sharpee, & Schreiner, 2008), making it tempting connect these literatures with the multidimensional 524 nature of speech categories and the dynamic nature by which input dimensions are mapped to behaviorally-525 relevant categories. Indeed, a recent study demonstrates that plasticity in adult rodents impacts auditory 526 sensitivity to combinations of acoustic input dimensions (Shepard, Lin, Zhao, Chong, & Liu, 2015). Using single-527 unit recordings and electrophysiological mapping in an adult mouse model, Shepard et al. demonstrate that 528 auditory core cortical activity differentiates species-specific vocal categories. Moreover, a distinct set of core 529 auditory cortical (putative pyramidal) neurons develop increased sensitivity to specific *combinations* of auditory 530 dimensions in newly-acquired vocalization categories. At a population level, this plasticity reflects the 531 differential weighting across acoustic input dimensions associated with behaviorally-relevant vocalization 532 categories. Inasmuch as the auditory representation of behaviorally-relevant acquired categories comes to reflect 533 the combinations of acoustic dimensions signaling the categories with differential perceptual weights, this model may provide a productive framework for discovering neurobiological bases of perceptual weighting in 534 the auditory system, how these weightings emerge with experience, and how they might be dynamically re-535 weighted by short-term regularities in the input, as observed for speech (Idemaru & Holt, 2011). 536

537 Dimension-based Attention to Acoustic Frequency. In both human and non-human animals, auditory attention is 538 often studied by comparing neuronal responses when the animal is engaged in a demanding behavioral task 539 (Tsunada, Liu, Gold, & Cohen, 2015) or specific readiness state (Carcea, Insanally, & Froemke, 2017), versus passive listening or less constrained activity. This makes it difficult to disambiguate effects of task, overall 540 arousal, motor activity, and cross-modal attentional allocation from the effects of attention within a given 541 dimension -- for instance, attending to a higher or lower frequency band. Recently, Schwartz and David (2017) 542 created a novel rodent experimental paradigm to direct attention to one of two frequency bands. Ferrets were 543 simultaneously presented with 2 streams of dynamically filtered narrowband noise, with each band presented 544 at a different spatial location to enable behavior. Distributed over multiple trials, one band contained embedded 545 higher-SNR 'cue' tones at the band's center frequency (serving to draw the ferret's attention to that band), with 546 both bands containing embedded 'probe' tones at lower SNRs, which served as target and foil stimuli. With 547 training, ferrets very accurately detected target and ignored foil tones. But, in contrast to what might have been 548 expected from work in vision (where attending to one part of retinotopic space increased firing for neurons 549 preferring that location) as well as in recent auditory mapping work (Da Costa, Van Der Zwaag, Miller, Clarke, 550 & Saenz, 2013), Schwartz and David (2017) found that most primary auditory cortex neurons' responses (spike 551 rate) to the narrowband noise around the attended tone frequency *decreased* compared to when the same noise 552 was ignored. By contrast, spiking to the probe tone did not change significantly depending on whether its 553 frequency band was attended or ignored. The authors suggest that this pattern may reflect very narrowly tuned adaptive suppression of *non-informative* noise around the cued frequency. This possibility will be interesting to 554 test in future work and that also harkens back to the human studies discussed above showing adaptive 555 reweighting of auditory cues based on their utility for extracting information from the speech stream. 556

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The Impact of Context. The directionality of the context-dependent speech phenomena we reviewed above, and 561 others like it in the literature, is *contrastive*. The pattern of results is such that perception is shifted *away* from the 562 acoustic input dimensions of the preceding context, consistent with neural systems that emphasize change. 563 Whether speech or nonspeech, precursors sampling a higher-frequency band shift speech categorization toward 564 categories characterized by lower-frequency spectral energy. However, the alignment of the dimension or 565 feature distinguishing the speech categories -- for example the third formant frequency band in /ga/ versus 566 /da/ - with the dimension or feature manipulated across the precursor context appears to be critical. Recall that 567 manipulating the long-term average spectrum of a precursor in the third formant (F3) frequency band shifts 568 /ga/-/da/ speech categorization. However, manipulation of the long-term average spectrum of preceding 569 speech or nonspeech in the first formant (F1) frequency band has no effect on /ga/-/da/ categorization even though manipulations to the F1 frequency band do produce contrastive context effects on vowel categorization 570 across vowels distinguished by their F1 frequencies (Huang & Holt, 2012). 571

572 Animal neurobiological studies suggest that stimulus-specific adaptation (SSA) exhibits an intriguingly similar 573 profile in both its dimension- or feature-selectivity and response characteristics (Ulanovsky, Las, & Nelken, 2003) 574 In SSA, neural responses to a particular stimulus are reduced in amplitude and delayed in latency when a 575 stimulus with similar acoustics precedes it compared to the neural response to the same stimulus presented in 576 isolation. However, SSA is not evident when the precursor stimulus is distinct enough that it fails to activate overlapping stimulus-specific neural populations (Jääskeläinen et al., 2007; May et al., 1999). In line with 577 proposals made by Ulanovsky and colleagues (2003) the depression of neural response to regularity and the 578 corresponding exaggeration of change with enhanced neural response may provide a means by which the 579 system responds to regularity present across input dimensions. (See Hermann, Henry, & Obleser, 2013a for an 580 example in human listeners). 581

582 Further, in a series of human behavioral studies of speech categorization Holt (2006) observed that the mean frequency of a distribution of tones (whether the distribution varied across 1000 Hz or included only tones 583 repeated at the mean frequency) was the best predictor of its influence of categorization of subsequent speech. 584 This resonates with findings from animal neurobiology. Ulanovsky et al. (2003) examined the response of 585 primary auditory cortex neurons to equally probable, equal-amplitude tones with 20 different frequencies. The 586 responses of the primary auditory cortex neurons to frequencies at the center of the stimulus frequency range 587 adapted the most and there was relative enhancement of responses at the eccentric frequencies furthest away 588 from the center of the frequency range. This created a U-shape in the neural tuning curves, with maximal 589 adaptation at the central frequencies and relative enhancement at the edges. This appears to arise because 590 adaptation strength is negatively associated with the frequency difference between the present stimulus and the 591 stimulus from the preceding trial (Brosch & Schreiner, 1997; Ulanovsky et al., 2003). Thus, adaptation is greatest 592 for central frequencies because central frequencies, on average, have smaller frequency differences from the 593 preceding trials compared to eccentric frequencies. Holt (2006) argued that this may relate to the observation 594 that the mean frequency of a distribution of preceding tones is the best predictor of the impact of context on 595 speech categorization. In line with proposals made by Ulanovsky and colleagues (2003; 2004), the depression of neural response to regularity and corresponding exaggeration of change with enhanced neural response may 596 provide a means by which the system responds to regularity across specific input dimensions. SSA seems to 597 have some of the right properties to support the contrastive, dimension-specific contrast effects evident in speech 598 perception (Holt, 2006). 599

600 Moving animal neurobiological studies even closer to the behavioral phenomena of speech perception, a recent 601 study of songbird forebrain demonstrates that rapid discrimination of behaviorally-relevant vocalizations 602 depends not only on specific stimulus features, but also on expectations generated from context about upcoming 603 events (Lu & Vicario, 2017). When acoustic features of a target songbird vocalization differed from the statistical distribution of a preceding context song, auditory response to the target vocalization was significantly enhanced 604 relative to when it shared the same acoustic distribution as preceding context. Thus, songbird auditory forebrain 605 is dynamically modulated by acoustic context to emphasize complex acoustic dimensions that depart from the 606 regularities build up across prior context. In mammalian species, human and nonhuman animal auditory cortex 607 also is sensitive to statistical context across extended time scales (Yaron, Hershenhoren, & Nelken, 2012). 608

In this way, the distribution of acoustic dimensions evolving in incoming input provide a means of modulating auditory processing to bias the system to down-weight the significance of dimensions well-sampled in prior input and enhance those that are novel. Although these effects are not often spoken of as selective attention, this pattern of bias toward (or away from) a particular input dimension may be another way that the auditory system directs dimension-based selective attention to behaviorally-relevant objects and events. Indeed, Jääskeläinen et al. (2011) have made the case that the tuning of auditory cortical feature-specific neural populations via SSA is

especially intriguing in light of the fact that such cortical tuning has been implicated as a mechanism of auditory
 selective attention (Fritz et al. 2007).

### 619 1.4 Building a Bridge from Animal Neurobiology to Human Phenomena

620 Like the speech perception phenomena reviewed above, these illustrative examples from nonhuman animal 621 neuroscience demonstrate that the very mapping of acoustics to auditory dimensions and objects is dependent 622 upon an organism's prior history of experience, the short-term experience evolving in the local input, and 623 statistical relationships relating the present sound exemplar to those experienced previously. Yet, despite the 624 intriguing connections reviewed above, there remains a gulf between the speech perception phenomena and the paradigms of animal neurobiological research in examining putative roles for dimension-based auditory 625 attention. It would be highly desirable to have a human behavioral paradigm that could build a bridge this gulf 626 in constructing a neurobiological model of human auditory perception, including speech perception, that 627 incorporate dimension-based auditory attention. 628

To be clear, the goal need not be to model the speech phenomenon described above directly. Rather the aim would be to develop a productive test-bed for investigating non-spatial auditory dimension-based attention in human listeners that might inform us about the auditory mechanisms available to speech perception. In this context, any such paradigm would need to include several important elements.

First, nonspeech stimuli would be desirable as the use of speech complicates direct connections with the
 informative neurobiological research with nonhuman animals. Speech also makes it challenging to isolate
 specific auditory dimensions of selective attention and assessments across speech can be 'contaminated' by
 individual differences in language ability, native-language background, and other factors. Nonspeech sounds,
 in contrast, allow for fine-grained manipulation of acoustic parameters.

638 Second, task demands should require directing attention along a specific acoustic dimension. In humans, the 639 most straightforward means of directing attention is to instruct participants to focus on a particular dimension 640 (e.g., 'pay attention to the higher sounds), or on some sub-region of that dimension while ignoring another sub-641 region (e.g., 'the cue to press the button will be a high sound, and not a low sound'). Overtly guiding participants' 642 attention to a part of the spectrum is an attractive possibility. From a practical perspective, participants' attention 643 to frequency band can be directed using relative height terms. More importantly, frequency is the primary 644 dimension of auditory representation and it has been used so productively in animal electrophysiology research 645 on dimension-based auditory attention. In addition, it relates naturally to the formant-frequency-band-specific effects so common in speech perception, as well as to visual neuroscience paradigms that overtly direct attention 646 to parts of retinotopic space. 647

648 Such explicit, symbolic (language-directed), and *endogenously driven* attention is experimentally convenient in 649 that little to no training is required for participants to understand the task. However, it does not capture more 650 exogenous attentional effects, such as those driven by the acoustic saliency or informational structure of the 651 auditory scene. These effects are vital to account for, in that decades of psychoacoustic research using variants of the 'probe-signal' paradigm (Greenberg, 1968) have shown that detection and processing of isolated or 652 embedded tones is strongly modulated by the presence and reliability of the preceding spectral context (Cusack, 653 Decks, Aikman, & Carlyon, 2004; Dai, Scharf, & Buus, 1991; Green & McKeown, 2001; Hafter, Schlauch, & Tang, 654 1993; Hübner & Hafter, 1995; Larkin & Greenberg, 1970; Mondor, 1999; Mondor & Breau, 1999; Mondor, Breau, 655 & Milliken, 1998; Reeves & Scharf, 2010; Richards & Neff, 2004; Scharf, Quigley, Aoki, Peachey, & Reeves, 1987; 656 Scharf, Reeves, & Giovanetti, 2008; Scharf, Reeves, & Suciu, 2007; Tan, Robertson, & Hammond, 2008; Woods, 657 Alain, Diaz, Rhodes, & Ogawa, 2001; Wright, 2005). Such findings are highly reminiscent of those using 658 endogenous and exogenous spatial attentional cues in vision research (reviewed in Carrasco, 2011). Nor do 659 explicitly cued attention paradigms get at the putatively attentional mechanisms underlying the dynamic 660 perceptual reweighting along multiple dimensions, as discussed above for speech phenomena. Thus, a good 661 experimental model of dimension-selective auditory attention should allow for *simultaneous* driving of more 662 sustained, endogenous, and explicitly cued attention along with moment-to-moment manipulation of acoustic 663 and informational parameters that transiently guide exogenous and endogenous attention along different 664 auditory dimensions.

Finally, it would be desirable to utilize sounds that make strong demands on integration of information within a dimension, and to be able to manipulate the difficulty of this integration to place greater or lesser demands on the system, as this is surely a factor in speech processing. At the same time, it would be advantageous to be able to manipulate the relationship of a target input dimension with competing 'distractor' dimensions across sustained sound input. This would assist in bringing studies of non-spatial dimension-based auditory attention

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of the sort directed to brief segments of speech in closer alignment with more common studies of auditory
 attention across sustained sounds, as in the classic cocktail party phenomenon. An additional benefit is that this
 approach would align well with human neuroimaging tools and the demands of listening to continuous, fluent
 speech in everyday listening.

Building a bridge between mechanisms of auditory attention in spoken language comprehension to those
revealed by non-human electrophysiology has been an active research agenda in the EEG/MEG field (e.g., Ding
& Simon, 2013; Forte et al., 2017; Kong et al., 2014; O'Sullivan et al., 2014; Skoe & Kraus, 2010; Zion Golumbic et
al., 2012). In the same spirit, here we present a novel experimental approach that meets these desiderata and we
share four insights from preliminary research.

# 1.5 Sustained Auditory Selective Attention (SASA), A Novel Approach to Investigating Non-Spatial Dimension-based Auditory Selective Attention

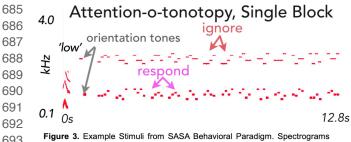


 Figure 3. Example Stimuli from SASA Behavioral Paradigm. Spectrograms (Time x Frequency) plot an example stimulus. Stimuli consisted of four-tone 'mini-sequences within a 'target' frequency band paired with a 'distractor' frequency band. A verbal cue (*high / low*) prompted listeners to monitor a specific band for mini-sequence repeats. This required listeners to maintain sustained auditory selective attention to the evolution of spectral structure across time within a specific frequency band in the context of similarlycomplex sounds in a distractor band. In recent work, we developed a novel behavioral paradigm we refer to as *SASA*, the Sustained Auditory Selective Attention paradigm. In the SASA paradigm, listeners direct attention to a series of four-tone 'mini-sequences' that fall within a specific spectra band, without any auditory spatial cues (see Figure 3). Listeners monitor for temporally-adjacent mini-sequence repeats within the attended band. This puts a high demand on encoding and integrating information across a delimited frequency range, the center frequency of which varies across trials. Adding to the challenge, target mini-sequences are accompanied by mini-sequences in a distractor frequency band that varies in its spectral distance from the target frequency band. The distractor band may also contain mini-sequence repeats. A verbal cue (*high, low*) directs

attention to a specific frequency band and brief 'orientation tones' alert listeners to the mean frequency of each
 band. Listeners report mini-sequence repeats in this target band with a key press.

The task meets the experimental *desiderata* outlined above in that it requires directing attention to a specific acoustic dimension, namely spectral band. (We discuss other manipulable dimensions below). The task involves nonspeech stimuli that make strong demands on integrating information (the mini-sequences) across an input dimension (the frequency band) and that can be extended across time to require sustained selective attention. Likewise, SASA requires spectrally-selective attention to a particular frequency band. In this, it aligns well with the nonhuman animal literature that has similarly capitalized on frequency as a significant acoustic input dimension across which selective attention can be directed (see Fritz et al. 2007).

In the next section, we describe four insights from utilizing this SASA paradigm among adult human listeners
 and describe how future work might exploit the approach further to make closer connections between the speech
 phenomena reviewed above and animal neurobiological models.

### 711 **1.6 Four Insights from SASA**

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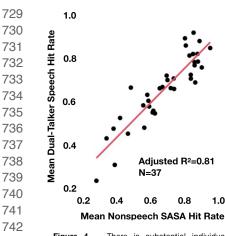


Figure 4. There is substantial individual variability in nonspeech SASA performance that is associated with speech comprehension in dual-talker conditions. Here, we plot each participant's average hit rates for nonspeech SASA versus dual-talker speech. The same relationship holds (R<sup>2</sup> = 0.81) when participants' accuracy for a standard mental rotation are included in the regression model, suggesting that the tight relationship between nonspeech SASA and dual-talker speech is not simply driven by individual differences in participants' generic ability to perform experimental tasks.

## 1.6.1 There are Substantial Individual Differences Even in Typical Young Adults

A first study examined the range of individual variation in SASA performance among healthy young-adult (N=37) university students. In this study, listeners completed a temporally-interleaved version of the nonspeech SASA task that complemented the simultaneous version shown in Figure 3. In this version, the high and low frequency bands alternated in time (every 125 ms) and listeners' task was to monitor one of the bands for mini-sequence repeats. The results are plotted in Figure 4. Even among this rather homogeneous sample of youngadult university students, there were substantial individual differences in performance on the nonspeech SASA task (apparent across the range variation on the Figure 4 x-axis). This is important in that it reveals that even healthy listeners differ in baseline ability to direct and sustain auditory selective attention to a specific acoustic dimension. Larger-scale future studies sampling a more diverse participant population have the potential to establish the range of individual variability evident among healthy listeners. This would be highly desirable as a benchmark for clinical assessment of dimension-based auditory selective attention among healthy older listeners who exhibit auditory selective attention difficulties, and among individuals with neurodevelopmental or neurodegenerative disorders that impact auditory attention (Shinn-Cunningham, 2017). It may be especially valuable that the SASA task is unlikely to be contaminated by language ability, native-language background, and other speech-specific factors.

## 1.6.2 Performance in the Nonspeech SASA Paradigm is Associated with Speech Comprehension in Dual Talker Conditions

In the same study, we also sought to examine whether the novel SASA task demanding selective attention to a specific frequency band across nonspeech stimuli relates to more common measures of auditory selective attention, specifically in the speech domain. For this reason, the same participants also completed a dual-talker speech task, similar to canonical multitalker studies of real-world listening challenges (Brungart, Simpson, Darwin, Arbogast, & Kidd, 2005). In this task, listeners attempted to detect exact repetitions in a string of 3 key words in the attended talker stream (male/female), while ignoring the other talker. As a control for overall performance, listeners also completed a version of a classic mental rotation task (Shepard & Metzler, 1971).

760 The strong relationship between SASA performance and dual-talker speech performance illustrated in Figure 4 761 indicates auditory selective attention to specific frequency bands, as measured using the novel nonspeech SASA 762 task, is strongly associated with dual-talker speech comprehension and holds even when mental rotation is 763 included as a factor in the general linear model to control for overall performance differences. This is important in that it indicates that performance in the nonspeech SASA paradigm is robustly associated with a multi-talker 764 speech comprehension challenge that demands dimension-based auditory selective attention. This is exciting 765 because it suggests that the nonspeech SASA paradigm can serve as a proxy for everyday listening challenges. 766 Whereas comprehension of speech in noise is a common model of auditory selective attention, the use of speech 767 complicates direct connections with informative neurobiological research with nonhuman animal models, 768 makes it challenging to isolate specific auditory dimensions of selective attention, and can be contaminated by 769 individual differences in language ability. The nonspeech SASA paradigm allows greater experimental control 770 over details of the target and distractor dimensions than is possible with natural speech stimuli and connects 771 directly to productive animal neurobiological models. Future studies more directly connecting this approach to 772 the speech phenomena reviewed above might, for example, take the approach of manipulating regularities 773 across which task-relevant information appears in a specific frequency band (to tap into perceptual weighting 774 and associated plasticity). Just as importantly, there is considerable opportunity to carefully manipulate 775 demands upon human spectrally-selective attention in order to address the many open questions regarding 776 basic mechanism.

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## 785 1.6.3 Listeners Can Learn to Better Deploy Dimension-based Auditory Attention

786 Especially intriguing, training can improve listeners' ability to deploy nonspatial dimension-based auditory selective attention. In a separate cohort of 787 788 listeners sampled from the same population of healthy university students, we provided two 1-hour sessions of training with feedback on the nonspeech 789 SASA task. As shown in Figure 5, most listeners improved in their ability to 790 integrate information in the target frequency band in the context of complex 791 acoustic information in a distractor frequency band. This implicates 792 behavioral training as a viable intervention that may improve dimension-793 based auditory selective attention among those with poor baseline abilities, 794 or clinical impairment of auditory selective attention. An exciting, as yet 795 unexplored, possibility is that such training might improve listeners' ability 796 to direct attention to specific frequency bands. It might be possible, for 797 example, to redirect spectral attention to higher frequencies that carry 798 significant speech information (Monson, Hunter, & Story, 2012; Monson, 799 Lotto, & Story, 2014; Vitela, Monson, & Lotto, 2015) in the context of noisy 800 surroundings that mask lower frequencies, thereby encouraging new perceptual weighting schemes beneficial to behavior. 801

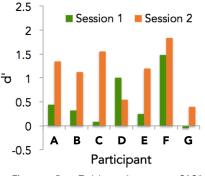


Figure 5. Training Improves SASA Performance. Brief (two-session, 90 minute) training with the SASA model paradigm improves performance in most participants.

# 1.6.4 Dimension-based Auditory Attention can be Topographically Mapped in Human Primary and Non Primary Auditory Cortex

804 Acoustic frequency is a particularly attractive model for dimension-based auditory attention in that (a) 805 informative and / or disambiguating acoustic cues in ecologically-relevant environmental sounds and intentional 806 communicative signals are unevenly distributed across the spectrum and (b) frequency is topographically 807 mapped across multiple auditory areas that differentially contribute to perceptual and decision processes. 808 However, as noted by Schwartz and David (2017), it has been challenging to come up with paradigms in 809 nonhuman animals that isolate frequency-selective attention from other attentional factors -- a primary goal of 810 our human SASA paradigm. In humans, recent work on spectrally-selective attention (Da Costa et al., 2013, see 811 also Paltoglou, Sumner, & Hall, 2009) has shown that when listeners attend to either a high or low frequency 812 stream containing behavioral targets (with both streams presented simultaneously, but to different ears), voxels 813 in auditory regions with preferred frequencies near an attended frequency band show increased blood-oxygenlevel-dependent (BOLD) activation, whereas voxels with preferred frequencies far from the attended frequency 814 band show decreased BOLD activity. Using an innovative melody-monitoring paradigm in a three-frequency-815 band stimulus, Riecke et al. (2016) showed that the topography of spectral attention significantly echoed 816

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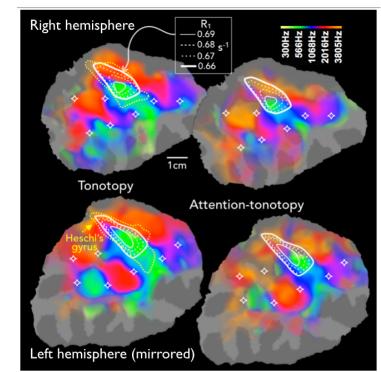
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tonotopic maps in early auditory areas; in putative secondary areas, attended frequency could be decoded using 841 multivoxel-pattern classification approaches, but did not seem to follow tonotopic progressions. 842



865 Figure 6. Group average Tonotopy and Attention-Tonotopy 866 maps on flattened superior temporal lobe patches, with R1 contours showing putative auditory core, from Dick et al. 867 (2017). Isocontour lines show quantitative R1 values for the 868 group-averaged putative auditory core, and color maps 869 showing group-averaged best frequency in both conditions. 870 The stars are fiduciary points to assist in visual comparisons of 871 maps across conditions; the outline of Heschl's gyrus is in 872 yellow dashed lines. The average tonotopic map is characterized by two pairs of three interlacing best-frequency 873 'fingers,' with the high-frequency fingers (red/orange colormap) 874 showing greatest frequency preference medially and extending 875 laterally, where they meet interdigitated lower-frequency 876 fingers (green/yellow colormap) extending lateral to medial, 877 with the longest 'middle' lower-frequency finger extending 878 about halfway into auditory core. This pattern is evident in 879 Fourier-analysis-derived maps of the Attention-tonotopy condition but not in the 'randomized control' for which the 880 attentional response was phase-cancelled (not shown here) 881

Using our non-speech SASA paradigm, we have spectral-based recently examined auditory selective attention in human cortex, combining functional MRI with high-resolution quantitative MRI in order to identify putative auditory core (Dick et al. 2017). Here, we observed that human primary and much of non-primary auditory cortical activation is strongly modulated by spectrally-directed auditory selective attention to five different frequency bands, in a manner that recapitulates its tonotopic sensory organization. The detailed, graded activation profiles elicited by single frequency bands (without distractors) were strongly associated with attentionally-driven activation when these frequency bands were accompanied by distractors (acoustic stimuli as in Figure 3, Figure 6 shows group average maps for tonotopic and 'attention-o-tonotopic' conditions from Dick et al., 2017). Moreover, systematic spatial maps of 'dis-preferred frequency' (the frequency that drove the smallest response at each voxel) could also be recapitulated by frequencydirected attention to those same frequencies. Finally, the graded frequency preferences observed in small patches across auditory cortex were closely aligned to those evoked by attention to those frequencies in the presence of distractor frequency bands.

1.6.5 SASA Overview and Future Directions These initial studies using the SASA paradigm demonstrate that we can non-invasively observe dimension-based auditory selective attention in the human brain by embedding task-relevant information in different regions of the frequency spectrum - here the dimension along which attention is directed. A major advantage of this approach is that brings human auditory cortical paradigms into closer alignment with informative electrophysiological animal research. Additionally, since behavioral research using the same paradigm indicates the close association of performance in this nonspeech SASA task with

882 comprehension of multi-talker speech it builds a bridge across which to connect traditional approaches in human 883 listeners like perception of speech in noise with these productive animal paradigms. Since training in the 884 nonspeech SASA paradigm leads to improvements in the ability to direct attention to specific frequency bands, 885 the pairing of training with these neuroimaging approaches can present new opportunities for understanding 886 how dimension-based auditory selective attention relates to short- and long-term plasticity. 887

Notably, these first studies using the SASA paradigm did not manipulate listeners' attention to auditory 888 dimensions other than spectral band, nor did they explore any other means of directing attention than through 889 specific verbal instruction. The SASA paradigm can accommodate explicit attention to other dimensions 890 through varying the acoustic character of the individual sequence elements, which are not limited to pure tones 891 but can be complex tones or synthetic sound objects. For instance, attention can be directed to durational or 892 timbral characteristics that define the task-relevant mini-sequence stream - similar to the way that listeners at a 893 concert will attend to spectrally and temporally overlapping flute or oboe lines in an orchestral piece. A SASA 894

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variant more analogous to the dimension-based dynamic reweighting effects discussed above might provide 897 multiple probabilistic acoustic cues that would predict the occurrence of a mini-sequence repeat. As an example, 898 explicit attention could be directed a given spectral band (as in the original SASA), but the acoustic 899 characteristics of the constituent tone elements would vary constantly in two dimensions (duration and 900 envelope) in both attended and unattended bands. Target mini-sequences would be more likely to occur when 901 preceded by tones of shorter duration, or a combination cue of shorter duration and sharper onset envelope. 902 Such a configuration would allow for listeners to discover and selectively direct attention along the acoustic 903 dimension(s) that are task-informative, as in the speech examples above. The dynamics of this (putative) 904 functionally-driven attentional reweighting could be directly compared to parallel manipulations in speech or 905 speech-like domains.

### 906 907 **1.7 Summary and Conclusions**

Was that a *beach* or a *peach*? This rather simple example, the auditory dimensions of which evolve across just 10s 908 of milliseconds, proves to involve more complex processing that has traditionally been described. In contrast to 909 early accounts of speech processing that emphasized rather static mapping of input dimensions to discrete 910 phonemic representations, contemporary research highlights that speech perception involves selective 911 weighting of acoustic input dimensions as a function of context and both short- and long-term input regularities. 912 We have attempted to make a case that selective attention to specific, non-spatial auditory dimensions may be 913 an important contributor in this dynamic mapping of speech input to behaviorally-relevant representations and 914 actions. Yet, the state of our understanding is such that there remain many open questions regarding this 915 putative link. We do not yet have a deep understanding of human auditory selective attention, especially as it 916 relates to directing attention to specific, non-spatial dimensions evolving within a sound object of the sort 917 potentially demanded by speech phenomena reviewed above. Nevertheless, there are important parallels 918 emerging in animal neurobiological research. This work suggests that the phenomena we refer to as involving selective attention are likely to draw from multiple neurobiological mechanisms. The hope is that paradigms that 919 put human and nonhuman animal research into closer alignment, as in the case of the SASA paradigm we 920 reviewed above, can facilitate progress in discovering the basic mechanisms of auditory selective attention 921 available to support higher-level processing like that demanded by speech to move us beyond selective attention 922 as a cognitive placeholder. 923

### 924 Author Note

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