

# Encoding of Illusory Continuity in Primary Auditory Cortex

Christopher I. Petkov,<sup>1,2</sup> Kevin N. O'Connor,<sup>1</sup> and Mitchell L. Sutter<sup>1,\*</sup>

<sup>1</sup>Center for Neuroscience and Section of Neurobiology, Physiology, and Behavior, University of California, Davis, Davis, CA 95616, USA

<sup>2</sup>Present address: Max Planck Institute for Biological Cybernetics, Tübingen, Germany.

\*Correspondence: [mlsutter@ucdavis.edu](mailto:mlsutter@ucdavis.edu)

DOI 10.1016/j.neuron.2007.02.031

## SUMMARY

When interfering objects occlude a scene, the visual system restores the occluded information. Similarly, when a sound of interest (a “foreground” sound) is interrupted (occluded) by loud noise, the auditory system restores the occluded information. This process, called auditory induction, can be exploited to create a continuity illusion. When a segment of a foreground sound is deleted and loud noise fills the missing portion, listeners incorrectly report hearing the foreground continuing through the noise. Here we reveal the neurophysiological underpinnings of illusory continuity in single-neuron responses from awake macaque monkeys’ primary auditory cortex (A1). A1 neurons represented the missing segment of occluded tonal foregrounds by responding to discontinuous foregrounds interrupted by intense noise as if they were responding to the complete foregrounds. By comparison, simulated peripheral responses represented only the noise and not the occluded foreground. The results reveal that many A1 single-neuron responses closely follow the illusory percept.

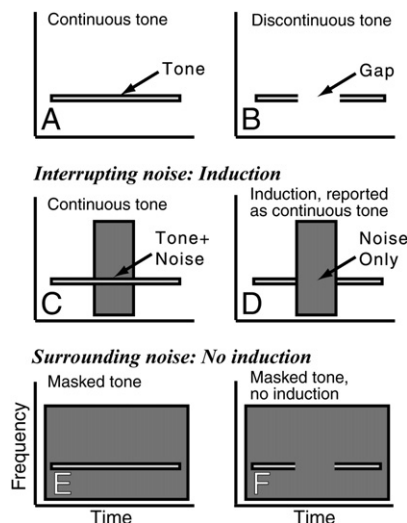
## INTRODUCTION

In natural environments, a sound of interest (a “foreground” sound) is often obscured by brief interrupting sounds produced by other objects (“background” sounds). For example, when a monkey attempts to identify another monkey’s vocalization, background bird chirps might interrupt the monkey vocalization. When interrupting background sounds are loud enough to completely obliterate a short underlying foreground segment, the auditory system fills in the occluded segment through a process called auditory induction, so-called because the foreground preceding and following the background sound induces perceptual restoration of the missing foreground segment. If there were no inducing foreground

segments preceding and following the loud noise, the foreground would be imperceptible because of masking (Figures 1E and 1F) by the background. Auditory induction is known by other names, such as amodal completion, fill-in, or phonemic restoration (Bregman, 1990; Miller et al., 2001; Petkov et al., 2003; Warren, 1970; Warren et al., 1972), and is an example of a general ability of the brain to perceptually organize sensory input to fill-in occluded information (Komatsu, 2006; Pessoa and De Weerd, 2003).

Auditory induction can be exploited to create an illusion, which was originally demonstrated with speech sounds. When segments were deleted from speech, the result was poor comprehension. However, when the removed segments were filled with loud noise, speech comprehension improved dramatically, providing compelling evidence that the brain restored the deleted information (Warren, 1970). Further studies (Bashford et al., 1988; Warren et al., 1972, 1988) demonstrated that illusory induction is not speech specific, but rather a general process that occurs with many foregrounds, including tones (Figures 1A–1D illustrate the stimulus configurations used in our study). This illusory induction has also been called the continuity illusion and is conceptually related to visual illusory contours (Day and Kasperczyk, 1983; Kanizsa, 1979), illusory motion (Assad and Maunsell, 1995), and induction (Rossi and Paradiso, 1996). An important requirement for auditory induction is that energy be present at induced frequencies. Thus, induction can be thought of as a process of perceptually organizing and assigning sound energy to various objects, selectively allocating ambiguous energy into a coherent scene, rather than the creation of an illusory percept in the absence of sensory stimulation.

Auditory induction has been studied behaviorally in humans (Kluender and Jenison, 1992; Warren, 1970; Warren et al., 1972, 1994; Wrightson and Warren, 1981), cats (Sugita, 1997), and monkeys (Miller et al., 2001; Petkov et al., 2003). However, the relationship of neuronal activity to induction remains a mystery. Psychophysical studies have provided two principles that guide the search for induction’s neural basis (Bregman, 1990; Bregman and Dannenbring, 1977; Houtgast, 1972). The first, called the “sufficiency of evidence rule,” states that during the occluding noise some neural activity should be indistinguishable from activity that would have occurred if the tone actually continued through the noise (Bregman, 1990). The

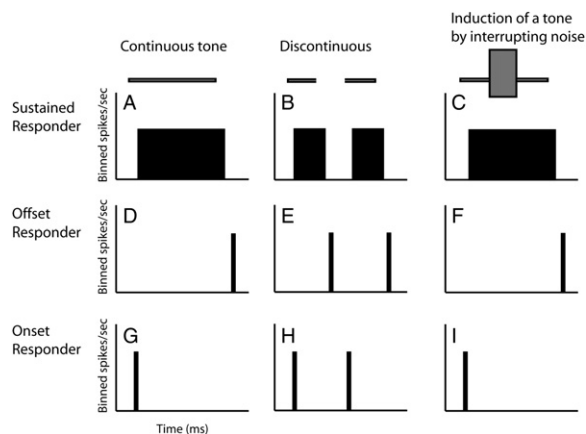


**Figure 1. Schematized Spectrograms Demonstrating Stimuli and Their Relationship to Illusory Induction and Masking**

Spectrograms of (A) continuous and (B) discontinuous tones and interrupting noise centered in (C) continuous and (D) discontinuous tones. *High-intensity* interrupting noise causes perceptual restoration (induction) of the deleted tone segment, with the tone being reported as continuous even when it is not. Here both a continuous foreground and noise are perceived. Spectrograms of surrounding noise temporally overlapping entire (E) continuous and (F) discontinuous tones. High-intensity surrounding noise masks the tone, and only noise is heard.

second, termed the “no discontinuity rule” (Bregman, 1990), states that there should be no neural evidence of transitions in the foreground (i.e., no evidence of the onset or offset of the foreground sound). This rule is based on experiments showing that induction is reduced or eliminated by placing a discontinuity or transition just prior to occluding-noise onset. For example, no induction occurs if an amplitude ramp is inserted into the foreground just prior to the interrupting noise, *even if the ramp is an amplitude increase*, which actually strengthens the foreground signal (Bregman and Dannenbring, 1977).

A heuristic model derived from these two rules helps to reveal the required single-neuron responses (Figure 2). First, to support the sufficiency of evidence rule, neurons with sustained firing to the foreground (Figures 2A–2C) should respond during induction as if the foreground were continuous (Figure 2C). Many auditory neurons have sustained responses to at least one stimulus (Wang et al., 2005). Next, to obey the no discontinuity rule, the model predicts that responses to tone transitions or discontinuities are eliminated by the occluding noise. Many auditory cortical neurons, even in awake preparations, are highly sensitive to amplitude transitions in sounds, commonly yielding phasic responses to tone onsets and offsets (Creutzfeldt et al., 1980; Erulker et al., 1956; Fishbach et al., 2001; Katsuki et al., 1959; Recanzone, 2000). Such phasic responses are well suited for detecting discontinuities in sounds and become more common as one ascends the auditory system (e.g., inferior colliculus,



**Figure 2. Heuristic Model of Single-Neuron Response Correlates of Auditory Induction**

On the top of each column are schematic spectrograms of the following stimuli: (A) a continuous tone, (B) a discontinuous tone, and (C) a discontinuous tone interrupted by intense noise. The latter (C) causes induction, and to be consistent with perception of a continuous tone during induction, responses in the third column (C, F, and I) should be like those to a continuous tone (A, D, and G). Each row shows schematized peristimulus time histograms (PSTHs) for AI neurons with sustained (A–C), offset (D–F), and onset (G–I) responses to the three stimuli.

Walton et al., 1997; thalamus, Schreiner, 1980; auditory cortex, Steinschneider et al., 1995; Eggermont, 1999). To obey the no discontinuity rule, phasic responders should fail to respond to tone transitions (Figures 2F and 2I) during induction. This model leads to a neural representation of induction as follows. When loud noise fills the gap (Figure 1D), three neural response components behave as if a continuous foreground were present, even though a part of it was deleted. Sustained responders should fire continuously (Figure 2C), as if there were no pause in the tone. Offset responders must fail to detect the offset of the initial tone segment (Figure 2F). Third, onset responders should fail to detect the reintroduction of the tone (Figure 2I).

To correctly interpret the neural evidence, it is necessary to recognize that induction comprises the percepts of *both* a continuous foreground and the interrupting noise. This is quite distinct from the percept created by loud masking noise surrounding the foreground in time (Figures 1E and 1F), which causes subjects to hear only noise (Bregman, 1990; Kluender and Jenison, 1992; Petkov et al., 2003). Therefore, when loud, interrupting (inducing) noise (Figures 1C and 1D) occludes a sound, the brain should respond as if a complete foreground and a noise were present. However, when loud surrounding (masking) noise (Figures 1E and 1F) is presented with the foreground sound, the brain should respond as if only noise were present. Here we contrast monkey primary auditory cortical responses to illusory induction and masking stimuli to evaluate whether single neurons encode the illusory induced sound features (induction) as opposed to following the physical stimulus attributes (masking). We used stimuli

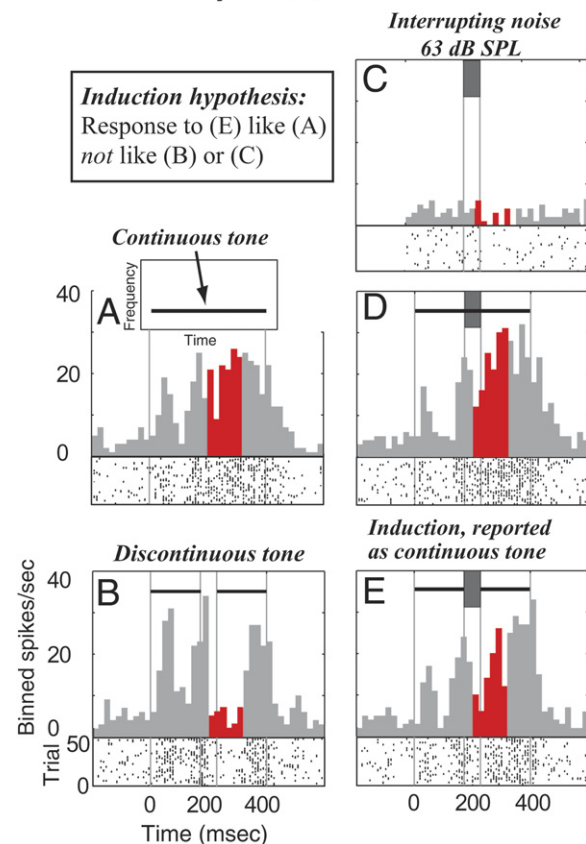
identical to those from a recent psychophysical study demonstrating auditory induction and masking in macaques (Petkov et al., 2003). Results are consistent with the model of Figure 2 and the perception of both the illusory foreground segment and interrupting noise during induction.

## RESULTS

Consistent with the hypothesized model of auditory induction, many A1 neurons represented the induced segment of occluded tones by responding to discontinuous tones occluded by intense noise (Figure 1D), as if responding to the complete tone without noise (Figure 1A). One such neuron responded to a continuous tone with an onset response followed by a pause and then sustained discharge (a sustained response, Figure 3A). When a gap was introduced (Figure 3B) during the period of sustained discharge to the continuous tone, a significant reduction in activity relative to the continuous tone response began ~35 ms after gap onset (compare Figure 3A to 3B during red/dark bins,  $p < 0.001$  bootstrap, see Experimental Procedures). However, responses to both continuous (Figure 3D) and discontinuous (Figure 3E) tones interrupted by loud inducing noise were similar to the responses to isolated complete tones (compare Figures 3D and 3E to 3A, no significant differences, bootstrap). The decrease in activity associated with gaps (Figure 3B, red/dark) was no longer observed when loud noise filled the gap (compare Figure 3B to 3E, red,  $p < 0.001$ ). In other words, the neuron responded as if the tone were complete under conditions that have been shown in monkeys and humans to cause illusory completion of the deleted segment (Kluender and Jenison, 1992; Petkov et al., 2003).

This neuron was not excited by the 63 dB SPL noise presented in isolation (Figure 3C), indicating that simple linear addition of the noise response (Figure 3C) to the discontinuous tone response (Figure 3B) cannot explain the response to the stimulus that is the linear combination of the two (Figure 3E). The stimuli in this study were carefully chosen such that the discontinuous tone with noise stimulus (Figure 3E in this example) was created by adding the discontinuous tone stimulus (Figure 3B) to the noise stimulus (Figure 3C). This allows application of a standard definition of linearity: the response to two stimuli added together is linear if it equals the sum of the response to the two stimuli presented in isolation. This definition will be used throughout the text. By this definition, while adjusting for spontaneous activity, linearity is violated because the response to the discontinuous tone with noise (Figure 3E) should be less than the response to the discontinuous tone (Figure 3B) because the noise is weakly inhibitory (Figure 3C). The response to the discontinuous tone with noise is actually larger than the response to the discontinuous tone without noise, suggesting that there is an opposite effect of the noise on the response to the discontinuous tone (net excitatory) than when the noise was presented in isolation (net inhibitory). Therefore, the dis-

## Sustained response neuron



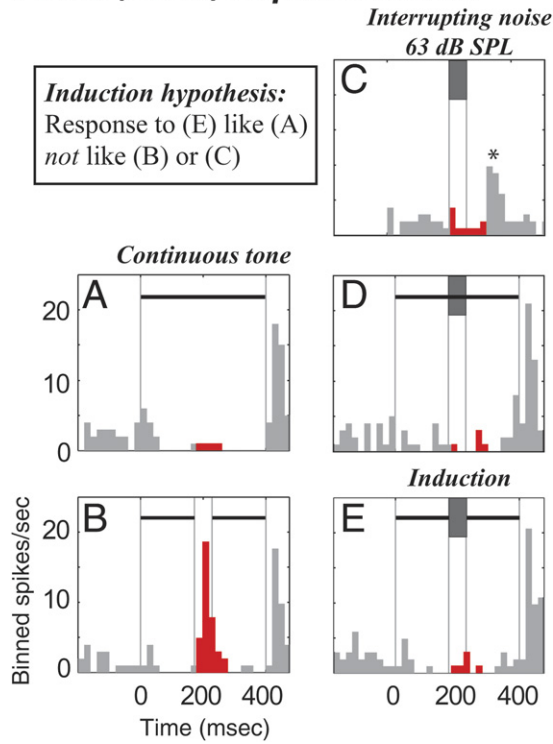
**Figure 3. Sustained Single-Neuron Response Consistent with Induction**

Above each peristimulus time histogram (PSTH) are schematized stimulus spectrograms (see [A] for spectrogram axis labels). Vertical gray lines in PSTHs align stimulus events. (A and B) PSTH and raster plots to continuous and discontinuous tones. Red (darker if printed in black and white) bins highlight times when the response is most different between continuous and discontinuous tones. (C) Response to 63 dB isolated noise. (D and E) Responses to continuous and discontinuous tones interrupted by 63 dB noise. In (C), isolated noise is time aligned to the identical noise components in the combined tone-noise stimuli, so noise onset is actually at time = 172 ms. Also in (C), the bins prior to time = -28 ms are empty because only 200 ms prestimulus spontaneous activity was collected for all stimuli. This should not be considered a lack of a response; spontaneous activity for this stimulus (C) can only be observed from time = -28 to time = 172 ms.

continuous tone plus noise response (Figure 3E) is much greater than the linear addition of the response to its components (Figures 3B and 3C).

Another neuron that behaved in a manner consistent with the perception of induction responded to tones with sustained inhibition followed by excitation to tone offset (a phasic offset response, Figure 4A). When a silent interval was introduced into the tone, the neuron responded to the first tone segment's offset with excitation (Figure 4B, red/dark). When the high-intensity interrupting (inducing) noise was added to the tones, the responses (Figures

**Phasic (offset) response neuron**



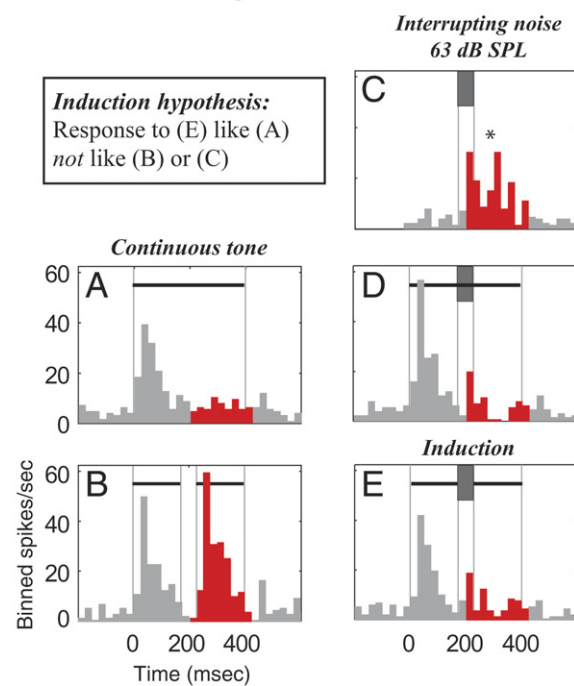
**Figure 4. A Single Neuron's Phasic (Offset) Response That Is Consistent with Induction**

Same format as Figure 3. Note that the noise-alone response, asterisk in (C), and the gap response (B) are missing during induction.

4D and 4E) were similar to the response to a complete tone in isolation (Figure 4A). This demonstrates that the neuron responded as if the tone were complete under conditions known to cause induction of the deleted segment (Figure 4E). The response to the stimulus that causes illusory continuity (Figure 4E), once again, cannot be predicted by simple linear summation of the responses to its components presented individually (Figures 4B and 4C). The first violation is the neuron's inability to detect the gap in Figure 4E (red bins) where linear summation of the responses to Figures 4B and 4C predicts a larger response during the red bins; it should be noted that the response at the same time to the nonillusory stimulus in Figure 4D is roughly linear, i.e., equal to the response in Figure 4C plus the response in Figure 4A. The second violation of linearity is the elimination of the excitatory response to noise presented in isolation (see asterisk in Figure 4C, at time ~300 ms) when the noise was presented in combination with tones (Figure 4D or 4E). While there are several possible explanations for this response, inhibition by the tone is a likely contributor.

Onset-responding neurons also behaved in a manner consistent with induction. One exemplary neuron responded to tone onset (a phasic onset response, Figure 5A). For discontinuous tones, the neuron also re-

**Phasic (onset) response neuron**



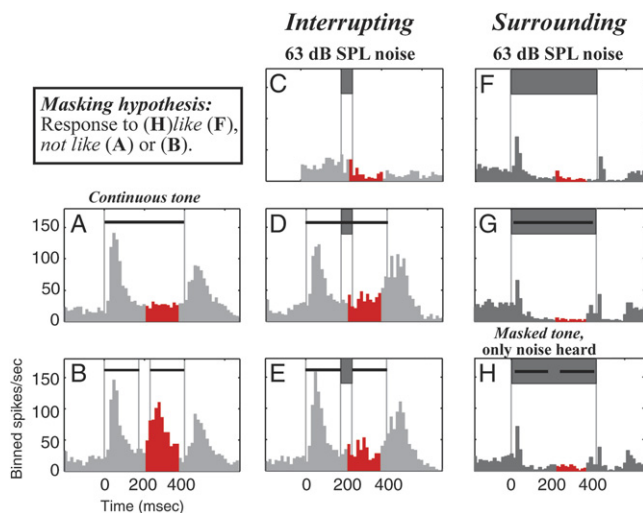
**Figure 5. Onset Response Consistent with Induction**

Same format as Figure 3.

sponded to the tone re-onset after the gap (Figure 5B) and was excited by noise (Figure 5C). This neuron also nonlinearly responded to the combined tone-noise stimuli. The gap-related response was suppressed by the presence of the noise in Figure 5E, even though the noise by itself was excitatory (Figure 5C). The result was that the stimulus known to cause illusory induction (Figure 5E) caused responses consistent with induction; that is, the response to a discontinuous tone interrupted with noise (Figure 5E) was similar to the response to a continuous tone (Figure 5A) and dissimilar to the noise (Figure 5C) and gap (Figure 5B) responses.

Another exemplary neuron responded with excitation to both tone onset and offset (Figure 6A) and with corresponding excitation to gaps in tones (middle peak of activity in Figure 6B, red/dark). Short, loud interrupting noise removed the gap-related response, making both complete and incomplete noise-interrupted tone responses (Figures 6D and 6E) similar to isolated complete tone responses (Figure 6A). Therefore, this neuron responded as if the tone were complete under conditions known to cause illusory completion of the deleted segment (Figure 6E). The four examples (Figures 3–6) demonstrate responses consistent with the induction model of Figure 2.

In contrast to the percept of induction, where occluded tone segments are heard continuing through brief interrupting noise (Figures 1C and 1D), loud noise completely surrounding tones in time (Figures 1E and 1F) creates a masking percept where only noise (and no tone) is heard



**Figure 6. A Single Neuron's Response Is Consistent with Induction for Interrupting Noise and Masking for Surrounding Noise**

Same format as Figure 3 except the right column (F–H) is for intense masking surrounding noise that causes only noise (and no tone) to be heard. Note that the differences in responses to interrupting and surrounding noise onsets likely are accounted for by differences in the onset ramps (0 ms for interrupting noise and 25 ms for surrounding noise, see [Experimental Procedures](#)).

(Bregman, 1990; Kluender and Jenison, 1992; Petkov et al., 2003). Accordingly, responses consistent with masking require only that responses to combined tone-noise stimuli resemble responses to noise presented in isolation, whereas neural responses consistent with induction require representations of both a continuous tone and the interrupting noise.

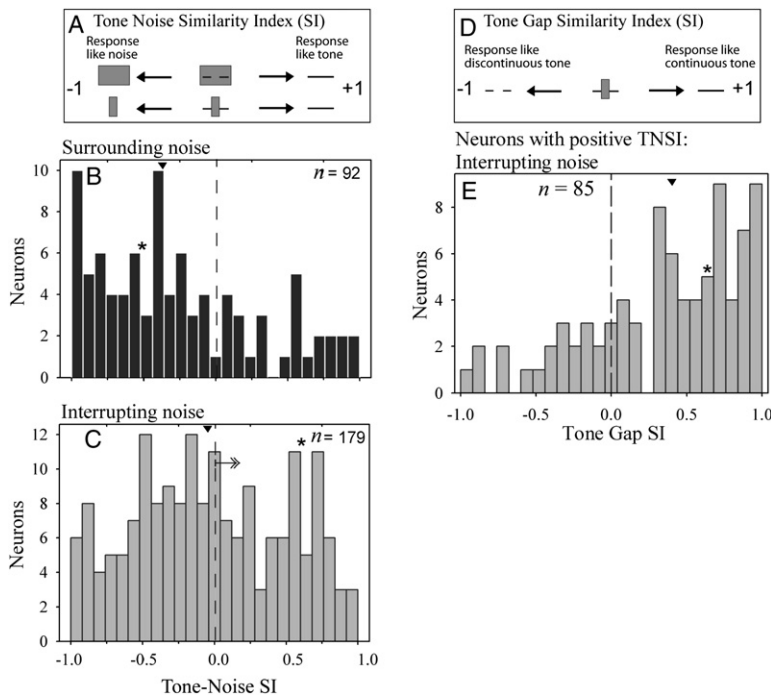
Neurons reflected the corresponding induction and masking perceptions with interrupting and surrounding noise, respectively. The neuron demonstrating induction-related responses in Figure 6E for interrupting noise responded as if being masked when presented with intense surrounding noise. For the long-duration noise, a short and weak onset response was followed by sustained inhibition (Figure 6F). This noise response differed markedly from tone responses, which had a stronger, longer excitatory component and no sustained inhibition (Figure 6A). When loud noise surrounded continuous or discontinuous tones, the neuron responded as if only noise were presented (compare Figures 6G and 6H to Figure 6F): a response consistent with masking.

Given the two distinct percepts of masking and induction, we predicted that neurons would respond to discontinuous tones with intense surrounding noise as if to isolated noise: a neuronal correlate of masking (i.e., cells only detect noise). However, for discontinuous foregrounds with intense interrupting (inducing) noise, a different result was predicted. Because during induction both the noise and the induced deleted foreground segment are perceived (Bregman, 1990; Kluender and Jenison, 1992; Warren, 1970; Warren et al., 1972), in order to be consistent with the percept of induction, the neuronal population must represent both the induced tone segment and the occluding noise. Accordingly, for interrupting noise we expected the population to represent the continuous tone (illusion) as well as the noise.

To test these predictions, differences between masking and induction were quantified using a response index, a normalized tone-noise-similarity index (TNSI, see [Exper-](#)

[imental Procedures](#), [Data Acquisition](#) and [Data Analysis](#), for details on the analysis and the neural sample). TNSI values ranged from  $-1$  to  $+1$ , signifying that responses to discontinuous tones presented with the loud noise were similar to noise ( $-1$ , masking) or tone ( $+1$ , induction) responses (Figure 7A). Responses to tones presented with intense surrounding noise were more like noise-only responses (median TNSI =  $-0.37$ ; Figure 7B), consistent with the masking percept of hearing only noise. In contrast, for loud interrupting noise, the population of neurons representing the tone and noise were more positively distributed (median TNSI =  $-0.07$ ; Figure 7C). Differences in median TNSI values for surrounding and interrupting noise were statistically significant, indicating that, for interrupting noise, more neurons responded as if the missing tone segment were present than did for surrounding noise (Kolmogorov-Smirnov [K-S] test:  $Z = 1.78$ ,  $p = 0.004$ ; Mann-Whitney:  $Z = 3.43$ ,  $p = 0.001$ ; see [Experimental Procedures](#)). This effect was observed separately for onset, sustained, and offset response components, consistent with the model in Figure 2 (sustained: interrupting noise  $n = 140$  versus surrounding noise  $n = 71$ , K-S test,  $Z = 1.60$ ,  $p = 0.012$ ; onset: interrupting noise  $n = 93$  versus surrounding noise  $n = 49$ ,  $Z = 1.52$ ,  $p = 0.02$ ; offset: interrupting noise  $n = 124$  versus surrounding noise  $n = 67$ ,  $Z = 1.40$ ,  $p = 0.04$ ).

For the neurons with positive TNSI values for interrupting noise—those hypothesized to represent the tone rather than noise during induction—we wanted to determine whether they represented the illusory tone segment because the TNSI does not rule out their representing the gap. If they responded as if a gap were present, this would suggest that the neurons neither supported masking nor induction. The tone-encoding neurons responded as if a continuous tone were presented, indicating that they represented the induced tone segment. We quantified this using a tone-gap-similarity index (TGSi). TGSi values ranged from  $-1$  to  $+1$ , signifying that responses to discontinuous tones presented with the loud noise were similar to



**Figure 7. Population of Single-Unit Responses Supports a Neural Representation of Masking for Surrounding Noise and of Induction for Interrupting Noise**

(A) Tone-noise similarity varies from +1 (response to a discontinuous tone presented with intense noise equals the isolated continuous tone response) to -1 (response to discontinuous tone presented with intense noise equals the isolated noise response). (B and C) Single-neuron TNSI distributions for surrounding and interrupting noise. (E) Neurons with positive TNSI values (see arrow in [C]) responded as if noise-interrupted discontinuous tones were continuous and not as if they were discontinuous. (D and E) Tone-gap-similarity index varies from +1 (response to discontinuous tone presented with intense interrupting noise equals the isolated continuous tone response) to -1 (response to discontinuous tone presented with intense noise equals the isolated *discontinuous* tone response). Arrowheads in (B)–(D) show the median, and asterisks (\*) show the TNSI and TGSI value for the example cell in Figure 6.

responses to a discontinuous (-1) or continuous (+1) tone presented without noise. The results with the TGSI indicated that when loud interrupting noise was used, 74% (63/85) of the neurons' responses were closer to a continuous than to a discontinuous tone response (Figure 7E). The effect was significant (one sample t test differed from 0,  $t = 6.3$ ,  $p = 0.000$ ; one sample K-S test of uniformity  $Z = 2.7$ ,  $p = 0.000$ ; median TGSI = 0.42). This indicates that most of the neurons representing the tone over the noise during the inducing stimuli are representing a continuous (rather than a discontinuous) tone, consistent with the induction percept.

The observations so far were based on recordings taken near each neuron's best-frequency response (BF, see *Experimental Procedures*). We also collected responses to 2 kHz tones—those used in the psychophysical studies of macaque induction (Petkov et al., 2003)—which were not always close to the neurons' BFs. This provided data from a larger population. We saw similar, albeit expectedly weaker, relationships in this data set (see *Supplemental Notes* and Figure S1 in the *Supplemental Data* available online). We also quantified several neuronal response characteristics that seemed to contribute toward induction (see the *Supplemental Notes* and Table S1).

### Simulated Peripheral Neuron Responses Cannot Support Induction

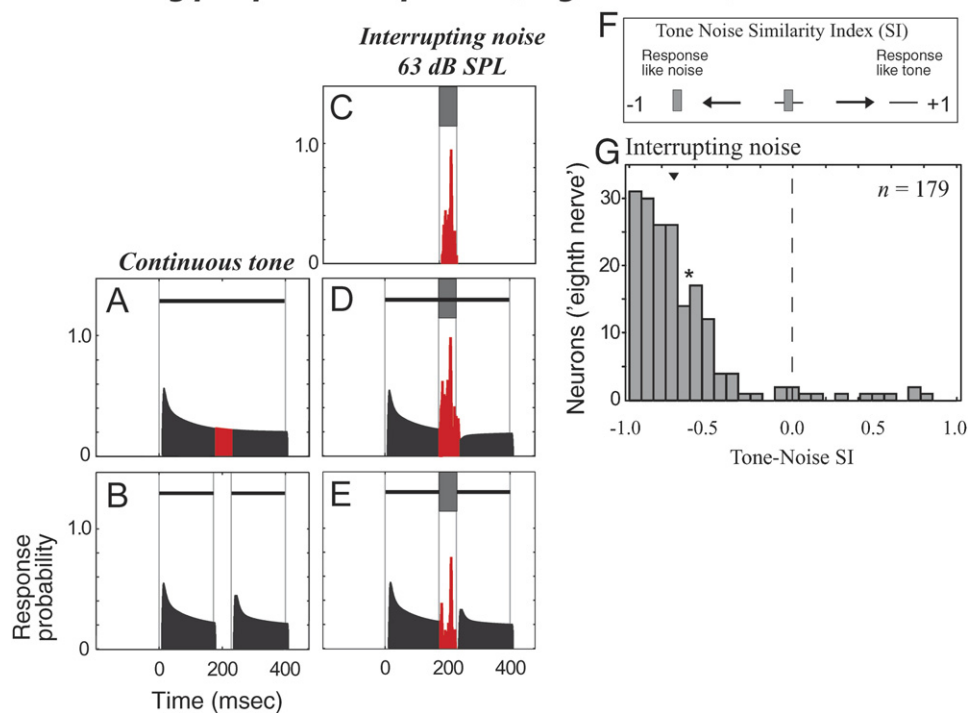
To determine whether the results we have so far observed simply reflect peripheral properties passed through the auditory system, we provided our stimuli as input to a cascaded peripheral processing model (see *Experimental Procedures*, *Simulation of Peripheral Responses*). We

then applied the same analysis for the simulated responses as we did for our A1 data.

Figure 8 shows an exemplary simulated peripheral "eighth-nerve neuron's" response to the stimuli we used to evaluate induction for A1 neurons. When the continuous tone was used as a stimulus (in this case a 2 kHz tone centered at the "BF" of this simulated neuron), the model showed a largely sustained "response," with adaptation following stimulus onset (Figure 8A). When a discontinuous tone was used there was a cessation in activity with a tone re-onset response following the gap in the tone (Figure 8B). There was little variability in tone/gap responses for these simulated peripheral neurons. Importantly, this peripheral neuron responded strongly to the interrupting noise by itself (Figure 8C), including when this noise was added to a continuous (Figure 8D) or discontinuous (Figure 8E) tone. The TNSI value for this example was negative (-0.62) showing that this example does not support induction.

Our impression from this and other examples was that the simulated peripheral neurons strongly responded to the noise, causing highly negative TNSI values (which supports masking rather than induction). This was confirmed by modeling a distribution of peripheral responses for neurons whose tone frequency/best-frequency relationships (Figure S1) matched those of the neurons sampled in A1 for Figure 7C. Thus, for stimuli containing interrupting noise, we modeled an identical distribution (in number) of "neurons" as collected for our sample of A1 neurons (see Figure S1 and *Experimental Procedures*). The distribution of interrupting noise TNSI resulting from the simulated auditory nerve neurons had a highly negative median

### Modeling peripheral responses ('eighth nerve')



**Figure 8. Simulated Peripheral Responses Support Only the Physical Properties of the Stimuli**

Shown is a modeled "eighth nerve neuron" response to the stimuli used to assess induction in our A1 neurons. Here the tonal stimuli are 2 kHz tones centered at the "BF" of the simulated neuron. For display purposes, the spontaneous "firing rate" is not shown. (A)–(E) is in the same stimulus/response format as for Figures 3A–3E. (F) is as in Figure 7A, which schematizes the range of TNSI response values that can be obtained, including how to interpret positive and negative values (positive values show that the response to the stimulus known to elicit induction are like those to a continuous tone; negative values that the response to this stimulus is as if only noise were presented). (G) shows the modeled distribution of peripheral responses. Arrowhead shows the median, and the asterisk (\*) shows the TNSI value for the example shown in (A)–(E).

( $-0.77$ , Figure 8G). The median TNSI using surrounding noise was similarly negative (median =  $-0.71$ ,  $n = 92$ ).

### DISCUSSION

Using stimuli that cause auditory induction, the population of neurons we sampled represented both the missing tone segment and the occluding noise, both of which are perceived during induction. Neurons representing the missing tone segment responded to discontinuous tones occluded by intense noise as if responding to complete tones. Consistent with the model of Figure 2, this neural code included both phasic responses that fail to detect discontinuities and sustained responses that continue through the occluded segment.

#### Population Codes: Perceiving a Complete Tone and Noise

We observed many neurons representing the missing tone segment in our inducing stimuli, although these were a minority of the neurons (63/179). There are two steps that led to a reduction from the 179 total neurons to 63 signaling the illusory segment. The first step was separating tone-

encoding (85/179) from noise-encoding (94/179) neurons. The second step was the observation that a minority of the tone-encoding neurons (22/85) responded as if a gap were present in the tone.

During induction, both a continuous foreground (in our case, a tone) and an interrupting noise are perceived. Thus, to be consistent with the percept, the population of neurons should represent both. A separate representation of tones and noise is consistent with the hypothesized parallel analysis of sound bandwidth for a multispectral wavelet-like analysis (Schreiner et al., 2000; Schreiner and Sutter, 1992; Sutter, 2005). Using the tone-noise-similarity index, we estimated that 94 neurons represented the noise and 85 represented the tone. It is worth considering that there are many intermediate bandwidth neurons in A1 (Recanzone et al., 2000; Schreiner et al., 1992), and many of these can respond to both the tones and noise. Such neurons could have TNSI values near 0. Thus, our approach of defining neurons with TNSI > 0 as representing the tone and those with TNSI < 0 as representing the noise likely results in some categorization errors.

Of the 85 putative tone-encoding neurons, 63 had positive tone-gap-similarity index values, indicating that they

responded more as if the deleted segment were present (induction) than absent. Why then would we find 22/85 neurons with negative TGSi values? Some might result from the classification errors in TNSi noted above causing us to inadvertently sample noise-encoding neurons. Furthermore, some neurons that represented the tone and noise might have a noise response that caused a response similar to the gap. These provide examples of how negative TGSi values could result from either statistical variation in responses (measurement noise) or reproducible responses caused by some neurons that encode both foreground and background. In either case, this suggests that there is some ambiguity in the neural code in A1 that must be resolved. This type of ambiguity in a brain area representing both foreground and background is inescapable and suggests some intriguing possibilities. First, the transformation which creates induction might not be complete at the level of A1. Second, an unbiased observer might be able to decide based on the aggregate activity of the population of foreground-encoding neurons. One decision function can result from equal weighting of all tone-encoding neurons' responses, so a positive median or mean TGSi from the distribution would lead to a decision that a continuous tone was present. Another possibility is greater weighting for neurons that more selectively represent a tone (i.e., neurons with more positive TNSi values). The aggregate of our data supports that A1 neurons represent the missing tone segment *and* the inducing noise, consistent with the entire induction percept.

### Population Codes: The Importance of Multiple Neuronal Response Types

While in this study we have found neurons that encode missing tone segments, it seems unlikely that an isolated population of "induction" neurons or a single response type could account for all induction phenomena for two reasons. First, induction has been found for every foreground sound tested. Second, the "no discontinuity" and "sufficiency of evidence" rules demand different contributions from different neuronal response types.

The general model, that a neuronal population should respond as if the induced foreground and interrupting noise were both present, is powerful because it does not depend on any single response type or physiological mechanism and therefore can be applied to any foreground sound. For the specific case of tonal foregrounds, this leads to a simplified model (Figure 2): onset and offset responses fail to detect the transitions in tones, and sustained response neurons continuously respond through the induced segment, which comply with both the no discontinuity and sufficiency of evidence rules. These conditions were both met by our results.

The psychophysical data supporting the two induction rules indicate that both sustained and phasic responses are important for induction. For an illusory inducing stimulus (Figure 1D), if neurons responding to tone *onset* fired at the reintroduction of the tone (reporting discontinuity) and

*sustained* responders continued to respond as if a continuous tone were present (reporting a continuous tone), the brain would have to resolve whether the tone actually continued through the noise. Evidence of tone onset/offset appears to weigh heavily in such resolution because psychophysical evidence of discontinuity in the tone disrupts induction (Bregman and Dannenbring, 1977). This makes sense because the noise contains energy at the tone frequency, so whether that energy comes from the tone or noise is ambiguous. If there is clear evidence of tone offset or onset at the gap, this resolves the ambiguity and suggests that the noise energy does not belong to the tone. But if only phasic neurons were present in a population, their decreases in activity might be erroneously interpreted as signaling the absence of a stimulus. However, *within a population of neurons* this reduction in activity can carry important information (Newsome et al., 1989), as long as some neurons, such as sustained responders, indicate the continuing *presence* of the sound. In this case, a decrease in activity can be just as informative as an increase because it supplies complementary information. If the brain were only to consider sustained responders, problems would also arise. A1 neurons must encode many sounds, so even an increase in activity may ambiguously represent different sounds or sound features. Thus, the joint activity from different neural response types can disambiguate the different sounds or sound mixtures that either class individually might not.

### Relationship to Masking

To what degree are processes akin to perceptual masking responsible for the interrupting noise results? Although removal of responses to transients can be thought of as a form of masking, simple peripherally mediated energetic masking (where a very loud *continuous* noise eliminates perception of a fainter foreground sound) cannot be responsible for the A1 results. Such energetic masking is commonly associated with auditory nerve responses, where large isolated *excitatory* noise responses dominate combined noise/tone responses (Rhode et al., 1978). The result is that the weak response produced by the low-intensity tone cannot be extracted from the much larger response created by the loud noise. We suspected the auditory nerve would only show a masking and not an induction correlate. The TNSi distributions obtained from our simulations of auditory nerve responses demonstrated highly negative TNSi values consistent with masking, i.e., responses more similar to the noise than the tone. In contrast, when recording in A1 using illusory inducing stimuli, more positive TNSi values were observed. This argues that factors other than known energetic masking properties of the periphery are contributing to the cortical responses. While we can rule out this simple form of masking, it is not unreasonable to predict that other more complex forms of masking, such as backward masking (Brosch et al., 1998; Pickett, 1959), contribute to induction.

### Mechanisms of Auditory Induction

We have shown that the responses of A1 neurons are consistent with induction and the simulated auditory nerve responses are not. An outstanding question is how do these cortical response properties arise and where in the brain? Our experiments were designed to address whether A1 activity *represented* the illusory sound segment, not necessarily to determine the mechanisms creating them. A1 was chosen as an initial area of study because it lies at the boundary between early and late processing of sound. From a cognitive neuroscience perspective, and because of the results of lesion studies suggesting the involvement of auditory cortex in conscious perception of sounds, e.g., [Graham et al., 1980](#); [Michel et al., 1980](#), A1 might be thought of as an early processing stage for encoding a perceptual phenomenon like induction. However, from an auditory physiology and mechanistic perspective, A1 might be thought of as a higher station in the auditory system that obtains many of its properties subcortically. By choosing A1, our results provide a crucial starting point for mechanistic studies studying in more detail how and where the response properties that we report in cortex are created and for performing studies in higher cortical areas to look for changes in these representations.

We stated in the [Introduction](#) that the continuity illusion results from trying to perceptually organize a potentially ambiguous sound signal into auditory objects. To illuminate how different brain areas might contribute to grouping sounds, we might look to another psychophysical example that involves perceptual grouping, comodulation masking release (CMR). CMR describes an increased ability to detect an unmodulated tone in the presence of a modulated noise when the noise is comodulated across bandwidth. As the envelope of the noise is confirmed across more frequency bands, it becomes easier to perceive the tone as a distinct object and to detect it. There is evidence for important contributions to CMR at the level of the cochlear nucleus ([Pressnitzer et al., 2001](#)), with progressive refinement and improvement of CMR-related properties with ascension up to A1 ([Las et al., 2005](#)). However, we cannot assume the same holds for induction. Although both induction and CMR are related to perceptual grouping, CMR and induction are very different perceptually and use very different stimuli. Therefore, while the previous CMR studies potentially provide a framework from which to view our induction results, CMR cannot directly speak to the neural origins of the cortical responses we see. The question of where the response properties that we observe are created can only be addressed by recording from many auditory stations. Still, it is an intriguing possibility that similar progressive refinement along the auditory neuroaxis might occur for induction and that auditory cortex might play a pivotal role in generating the representation of auditory objects ([Nelken et al., 2003](#)).

In addition to asking where these response properties arise, one can ask how they are created. Our findings highlight that induction requires coding multiple sound properties, through several neuronal response types,

and suggest the involvement of multiple cellular mechanisms. Explanations of the results can be made by describing how neurons respond to the time-varying stimulus spectrum and/or to the stimulus envelope. With respect to the time-varying frequency spectrum, the examples that we report lead to several intriguing possibilities, including but not limited to different nonlinear inhibitory effects ([Figures 4 and 5](#)) as well as disinhibition or facilitation ([Figures 3 and 6](#)). However, there are a plethora of possible alternative explanations, and at this point it would be premature to speculate. Additionally, we observed a much wider variety of responses than the most prominent examples shown here, typical of the heterogeneity of A1. Despite the heterogeneity of responses, the population of sampled neurons responded in a manner consistent with the induction percept. An advantage of our approach is that, regardless of the exact mechanisms shaping the induction-related responses, our analyses make it possible to evaluate the relationship between the responses of the population of A1 neurons and induction. This was achieved by comparing responses to the stimuli that cause induction to responses to individual stimulus components that were either physically present in the inducing stimulus or those that were perceived by subjects listening to the inducing stimulus. Ultimately, to reveal the mechanisms responsible for the observed correlate of induction will require recording from multiple brain regions and performing intracellular recording experiments.

The diversity of the observed induction-related responses with tonal foregrounds suggest that the ecological pressure to maintain stable representations of interrupted sounds is important enough that it has been selected upon or acquired through multiple neuronal encoding mechanisms. As such, searching for *the* induction neuron, single brain region responsible for induction, or single cellular mechanism responsible for induction might turn out to be a futile endeavor.

### Auditory Induction and Attention

Many studies show that certain forms of auditory induction appear to be a fairly automatic process that can occur outside the focus of attention ([Bregman, 1990](#); [Micheyl et al., 2003](#)). Further, our behavioral work in monkeys ([Petkov et al., 2003](#)) supports the idea that induction cannot be entirely overridden by attention. There the animals were unable to overcome the illusion despite being rewarded for detecting a gap in the sound. These results argue that some aspects of induction might be due to processing “early” in the auditory system where the influence of “top-down” cognitive control is not as strong as in nonprimary areas of human auditory cortex ([Grady et al., 1997](#); [Petkov et al., 2004](#); [Pugh et al., 1996](#)). Such a viewpoint is also consistent with EEG induction correlates found in humans not attending sounds ([Micheyl et al., 2003](#)) and with cortical-based modeling of induction at the initial stages of auditory cortex ([Husain et al., 2005](#)).

On the other hand, there is evidence that some forms of induction can utilize feedback connections. For example, induction with speech is thought to also invoke feedback (Sivonen et al., 2006). Other forms of perceptual grouping are known to build up over time and can be influenced by the redirection of subjects' attention (Carlyon et al., 2001). Such an attention effect has yet to be demonstrated with tonal foregrounds, however, and would likely be complementary to the automatic processes already reported. Because these monkeys were passively listening, we believe the present results provide a basis for understanding the "preattentive" foundations of perceptual induction, without assuming induction arises in A1 or excluding the possibility of further top-down modulation in higher areas.

### Summary and Conclusions

In summary, our results support that, under conditions that produce induction, the illusory tone segment is represented in A1. Of the major types of neurons investigated in this study, all three responded as if the tone were present—offset responders fail to encode gap initiation, sustained responders provide activity as if the tone continued, and onset responders fail to signal the gap's termination. This result is consistent with two principles of induction (Bregman, 1990; Bregman and Dannenbring, 1977; Houtgast, 1972): (1) there should be no neural evidence of gap onset/offset, and (2) during the noise, neural activity should be indistinguishable from activity that would have occurred had the tone actually continued. We conclude that A1 neurons demonstrate the brain's ability to compensate for transient noise in the environment by "filling-in" segments of sounds occluded by noise.

### EXPERIMENTAL PROCEDURES

#### Stimuli

Stimuli, presented from speakers (O'Connor et al., 2000, 2005; Petkov et al., 2003) placed 1.5 m from the animal, were identical to those previously reported for tonal foregrounds used in psychophysical experiments (Petkov et al., 2003) except that the frequency of the foreground could also be set to the best frequency of the recorded activity (see Figure S1). The foreground was a 45 dB SPL (unfiltered calibration, Brüel & Kjær 2231 sound level meter) 400 ms tone (cosine ramped, 8 ms rise/fall times), with a sample frequency of 50 kHz, or in the cases of the tone frequency being higher than the Nyquist frequency, at 100 kHz. Transitions into and out of the silent gap—temporally centered in the tone—had 3 ms rise/fall times. The gap duration—silent portion plus transitions—was 56 ms. Noise (broadband, 25 kHz cut-off) was calibrated in RMS level (dB SPL, re 20 micro Pascals). *Interrupting noise* was unramped and temporally centered in the foreground, corresponding to noise presentation from 172 to 228 ms after initial foreground onset. When a gap was present, this corresponded to noise completely overlapping the gap (including ramps) but not the tone segments outside of the gap (Figures 1D and 3E). *Surrounding noise* (450 ms, including 25 ms onset/offset ramps reaching their plateau when the foreground began, and beginning offset transition when the foreground was completed) temporally encompassed the entire foreground. In this paper, we only report results for 63 dB SPL noise conditions. The noises within a given type and intensity were "frozen" so that the only difference between continuous and discontinuous stimuli

with noise was the presence or absence of a silent gap in the foreground. Additionally, we presented the two noise types (interrupting and surrounding, e.g., Figures 6C and 6F) in isolation and the continuous and discontinuous tones in isolation so that responses to combined tone/noise stimuli could be compared to responses to tones and noise in isolation. Interrupting noise presented in isolation (e.g., Figures 3C, 4C, and 5C) was time aligned to its occurrence in the combined stimuli (Figures 3D and 3E, 4D and 4E, and 5D and 5E) and in the figures is aligned to start at time = 172 ms (e.g., Figure 3C).

#### Data Acquisition

Standard extracellular recording techniques were used to record from the right hemispheres of two naive adult macaque monkeys, conforming to the PHS policy on animal care. Subjects were on a restricted water access protocol approved by the UC Davis animal care and use committee. Extracellular recordings occurred with macaques awake, seated, head restrained in a primate chair, designed to be "acoustically transparent," within a double-walled, sound attenuated, and foam-lined chamber (IAC: 2.9 × 3.2 × 2.0 m<sup>3</sup>, internal). For further recording and single-unit (spiking neuron) isolation details see O'Connor et al., 2005.

We recorded from 304 single units with interrupting noise, of which 153 were also recorded with surrounding noise. We recorded with the foreground frequency close to BF and at 2 kHz (the frequency used in macaque psychophysics; Petkov et al., 2003), resulting in a sample of 494 and 210 neurons recorded with interrupting and surrounding noise, respectively. Our analyses are either from neurons recorded with the tone frequency close to BF (see Results, Figure 7), from the entire sample (see Table S1 and Figure S1), or from a subdivision of the sample based on the type of response of neurons to different sounds (Figure S3).

For localizing recordings from A1, we first stereotaxically guided the electrodes to A1's relative anatomical position within the macaque auditory cortex (Paxinos et al., 2000). Then we identified A1 by its response latency, responsiveness to tones, and its direction of tonotopic gradient for best-frequency responses to tones (Merzenich and Brugge, 1973; Recanzone et al., 2000). The area extended by the tonotopic gradient in the antero-posterior direction (Hackett et al., 2001) and the medio-lateral extent of tone responsiveness (Rauschecker and Tian, 2004) supported that recordings were from field A1.

#### Data Analysis

##### Determining a Neuron's Best-Frequency Response

BF was determined using an interpolation method so as not to rely solely on the response to a single tone frequency. This incorporated the tone frequency eliciting maximal response ( $S_b$ , sum of spikes response) and the responses to the two neighboring frequencies ( $S_a$  and  $S_c$ , response to the neighboring lower and upper tone frequencies, respectively). From these three responses we determined a weighting factor as follows:

$$w = ((S_b - S_a) / ((2 * S_b) - S_a - S_c))$$

Then BF was calculated:

$$BF = f_a * 2^{(w * \text{OctRange})}$$

where  $f_a$  is the tone frequency (Hz) eliciting the  $S_a$  response and OctRange is the range in octaves between  $f_a$  and  $f_c$ . In the case of two frequencies with maximal responses, the BF was half way (in octaves) between these two frequencies (e.g.,  $w$  becomes 0.5).

We obtained a fairly even sample of BFs from our entire sample (on an octave scale) ranging from 150 to 40,025 Hz. The relationship of neuronal BF to the tone frequency used is shown in Figure S1A.

##### Tone-Noise-Similarity Index and Tone-Gap-Similarity Index

The TNSI was used to quantify how similar each single-unit's response to the discontinuous tone with intense noise (DTIN) was to the isolated tone response (T) or isolated noise response (N):

$$\text{TNSI} = (A - B) / (A + B),$$

where  $A = |N - \text{DTIN}|$  and  $B = |T - \text{DTIN}|$ . N, DTIN, and T for the analysis presented in the paper was the response in spikes counted over a time window as described below. When noise was used in isolation (e.g., Figures 3C, 4C, and 5C), the window for counting spikes was aligned with when it would occur in the combined tone/noise stimulus. The TGSi was similar to the TNSI except in the above equation the noise response (N) is replaced with the response to the tone with a gap (G), so that  $A = |G - \text{DTIN}|$ .

#### Procedures for Comparing Responses

Many A1 responses have multiple components (inhibitory and excitatory) with high temporal precision. To prevent these responses from opposing each other, four different procedures were used to compare responses, each with their own advantages. Three involved choosing a time window over which to count spikes, and one involved comparing (correlating) entire peristimulus time histograms (PSTHs) without choice of a time window. These spike-count and correlation measures were then used to derive TNSI and TGSi values. All statistical tests with TNSI and TGSi yielded the same results (for both significant and not significant effects), regardless of which of the four procedures were used. For Figures 3–7 and associated analyses in the text, we strictly used the method based on a statistical criterion, where all neurons that contributed could be said to significantly encode gaps in tones. Elsewhere, we also report results using the other methods, including those from the entire sample, in Table S1 and the Supplemental Data.

For the methods in the paper, a statistical criterion was used to define the time window over which to count spikes. First, a difference PSTH (5 ms bins, 50 stimulus repetitions) was created by subtracting the *discontinuous* tone (without noise) PSTH from the *continuous* tone (also without noise) PSTH. Then a neuron was evaluated only if this difference PSTH, following the onset of the gap, had a maximum (of absolute value) that was significant. The statistical criterion was that two consecutive bins be two standard deviations (SD) above the prestimulus “spontaneous” activity or that one bin be four SDs above spontaneous activity. If this criterion was met, the starting and ending points of the window were determined by finding, in both directions from the maximal bin, the third consecutive bin that was below two SD; the analysis window was identified as starting and ending on these bins. The distribution of these response windows showed a narrow median width of 75 ms with a median starting position of 241 ms following tone onset (69 ms after gap onset). We also counted spikes fired to the two types of noise within these windows and saw that more spikes were elicited by the short-duration interrupting noise (median 9.9 spikes) than the longer duration surrounding noise (median 5.3 spikes), differing at the  $p < 0.05$  level (K-S test). This rules out that more negative TNSIs for surrounding noise were due to larger excitatory responses to the longer-duration noise.

This method may be preferred for evaluating neurons involved in induction because it selects a narrow statistically based window of the response to the gap, without considering the noise responses. The method selects the neurons that show that they can discriminate continuous from discontinuous tones in a statistically significant manner and therefore those most likely to represent the studied foreground differences.

#### Classifying Tone Responses

We objectively classified tone responses as a prior classification of macaque A1 responses (Recanzone, 2000). We windowed the tone stimulus response into three intervals: early (0–200 ms), late (200–400 ms), and offset (425–625 ms). Significant responses (two bins above two SD or one bin above four SD of the baseline activity) occurring within the early interval were identified as “phasic-onset” responses. “Phasic-offset” responses were defined as significant responses in the offset interval. “Sustained” (excitation) responses were identified as significant responses occurring in both the early and late periods. Sustained inhibition was difficult to detect using our standard statistical criterion (these responses were usually close

to the mean spontaneous level); thus, we assigned inhibitory responses to the “sustained” category if they were below the mean spontaneous level for more than 65 ms.

#### Statistical Analyses

To test whether interrupting versus surrounding noise distributions differed (e.g., Figure 7C versus 7B), we used the nonparametric Kolmogorov-Smirnov (K-S) test, which allows non-normally distributed data to be tested. A nonparametric test such as this was also important because the sample size for the surrounding noise distribution was smaller due to oversampling of neuronal data for the stimuli containing interrupting noise. Results were statistically the same (for significant, at  $p < 0.05$ , or not significant effects) when also using the more common nonparametric Mann-Whitney test. Although the K-S test is sensitive to differences in shape as well as central tendency, it seemed our effects were largely based on differences in central tendency, since subtracting the mean or median of the distributions between interrupting and surrounding noise (see Figures 7B and 7C) removed the significance of the observed differences. For testing whether a single distribution differed from zero, we used a one-sample t test (two-tailed prediction) and a one-sample K-S test of uniformity in the distribution.

#### Simulation of Peripheral Responses

We used Malcolm Slaney’s Auditory Toolbox (version 2), which is coded in Matlab and implements a number of models of peripheral processing (Slaney, 1998). Using the toolbox, we implemented a cascaded model to obtain simulated responses of auditory (eighth) nerve fiber/neuron responses. The first component of the model is an auditory filter bank proposed by Patterson and colleagues (Patterson et al., 1992, 1995; Slaney, 1993), which uses a  $\gamma$  tone bank of auditory filters with an equivalent rectangular bandwidth based on measurements of critical bands (ERB; Glasberg and Moore, 1990). This models basilar membrane motion, which will elicit potentials in the inner hair cells, the output of which can be considered a narrow-band auditory filter or channel. Here we used 75 channels, with upper and lower frequencies matching the range of BFs found in our A1 neuron samples. The second “Meddis” component (Meddis, 1986; Meddis et al., 1990) was added at the output stage of the ERB filterbank (Slaney, 1998). This model simulates response properties such as adaptation following stimulus onset. Standard parameters for this model were used (Meddis et al., 1990; Slaney, 1998). The output of this model is the spike probability of an auditory nerve fiber, i.e., eighth nerve neuron.

#### Supplemental Data

The Supplemental Data for this article can be found online at <http://www.neuron.org/cgi/content/full/54/1/153/DC1/>.

#### ACKNOWLEDGMENTS

We thank M. Gazzaniga, J. Johnson, E.G. Jones and M. Merzenich for comments on previous versions of this manuscript. This work was supported by grants from the National Institute on Deafness and other Communication Disorders (DC-02514), the McDonnell Foundation, the M.I.N.D. Institute, and the Alexander von Humboldt Foundation. The authors declare that they have no competing financial interests.

Received: September 28, 2006

Revised: January 30, 2007

Accepted: February 18, 2007

Published: April 4, 2007

#### REFERENCES

- Assad, J.A., and Maunsell, J.H. (1995). Neuronal correlates of inferred motion in primate posterior parietal cortex. *Nature* 373, 518–521.
- Bashford, J.A., Jr., Meyers, M.D., Brubaker, B.S., and Warren, R.M. (1988). Illusory continuity of interrupted speech: speech rate determines durational limits. *J. Acoust. Soc. Am.* 84, 1635–1638.

- Bregman, A.S. (1990). *Auditory Scene Analysis* (Cambridge, MA: MIT Press).
- Bregman, A.S., and Dannenbring, G.L. (1977). Auditory continuity and amplitude edges. *Can. J. Psychol.* *31*, 151–159.
- Brosch, M., Schulz, A., and Scheich, H. (1998). Neuronal mechanisms of auditory backward recognition masking in macaque auditory cortex. *Neuroreport* *9*, 2551–2555.
- Carlyon, R.P., Cusack, R., Foxton, J.M., and Robertson, I.H. (2001). Effects of attention and unilateral neglect on auditory stream segregation. *J. Exp. Psychol. Hum. Percept. Perform.* *27*, 115–127.
- Creutzfeldt, O., Hellweg, F.C., and Schreiner, C. (1980). Thalamocortical transformation of responses to complex auditory stimuli. *Exp. Brain Res.* *39*, 87–104.
- Day, R.H., and Kasperczyk, R.T. (1983). Amodal completion as a basis for illusory contours. *Percept. Psychophys.* *33*, 355–364.
- Eggermont, J.J. (1999). Neural correlates of gap detection in three auditory cortical fields in the cat. *J. Neurophysiol.* *81*, 2570–2581.
- Erulker, S.D., Rose, J.E., and Davies, P.W. (1956). Single unit activity in the auditory cortex of the cat. *Johns Hopkins Hospital Bulletin* *39*, 55–86.
- Fishbach, A., Nelken, I., and Yeshurun, Y. (2001). Auditory edge detection: a neural model for physiological and psychoacoustical responses to amplitude transients. *J. Neurophysiol.* *85*, 2303–2323.
- Glasberg, B.R., and Moore, B.C. (1990). Derivation of auditory filter shapes from notched-noise data. *Hear. Res.* *47*, 103–138.
- Grady, C.L., Van Meter, J.W., Maisog, J.M., Pietrini, P., Krasuski, J., and Rauschecker, J.P. (1997). Attention-related modulation of activity in primary and secondary auditory cortex. *Neuroreport* *8*, 2511–2516.
- Graham, J., Greenwood, R., and Lecky, B. (1980). Cortical deafness: a case report and review of the literature. *J. Neurol. Sci.* *48*, 35–49.
- Hackett, T.A., Preuss, T.M., and Kaas, J.H. (2001). Architectonic identification of the core region in auditory cortex of macaques, chimpanzees, and humans. *J. Comp. Neurol.* *441*, 197–222.
- Houtgast, T. (1972). Psychophysical evidence for lateral inhibition in hearing. *J. Acoust. Soc. Am.* *51*, 1885–1894.
- Husain, F.T., Lozito, T.P., Ulloa, A., and Horwitz, B. (2005). Investigating the neural basis of the auditory continuity illusion. *J. Cogn. Neurosci.* *17*, 1275–1292.
- Kanizsa, G. (1979). *Organization in Vision: Essays on Gestalt Perception* (New York: Praeger).
- Katsuki, Y., Wantanabe, T., and Maruyama, N. (1959). Activity of auditory neurons in upper levels of brain of cat. *J. Neurophysiol.* *22*, 343–359.
- Kluender, K.R., and Jenison, R.L. (1992). Effects of glide slope, noise intensity, and noise duration on the extrapolation of FM glides through noise. *Percept. Psychophys.* *51*, 231–238.
- Komatsu, H. (2006). The neural mechanisms of perceptual filling-in. *Nat. Rev. Neurosci.* *7*, 220–231.
- Las, L., Stern, E.A., and Nelken, I. (2005). Representation of tone in fluctuating maskers in the ascending auditory system. *J. Neurosci.* *25*, 1503–1513.
- Meddis, R. (1986). Simulation of mechanical to neural transduction in the auditory receptor. *J. Acoust. Soc. Am.* *79*, 702–711.
- Meddis, R., Hewitt, M.J., and Shackleton, T.M. (1990). Implementation details of a computation model of the inner hair-cell/auditory-nerve synapse. *J. Acoust. Soc. Am.* *87*, 1813–1816.
- Merzenich, M., and Brugge, J. (1973). Representation of the cochlear partition of the superior temporal plane of the macaque monkey. *Brain Res.* *50*, 275–296.
- Michel, F., Peronnet, F., and Schott, B. (1980). A case of cortical deafness: clinical and electrophysiological data. *Brain Lang.* *10*, 367–377.
- Micheyl, C., Carlyon, R.P., Shtyrov, Y., Hauk, O., Dodson, T., and Pull-  
vermuller, F. (2003). The neurophysiological basis of the auditory continuity illusion: a mismatch negativity study. *J. Cogn. Neurosci.* *15*, 747–758.
- Miller, C.T., Dibble, E., and Hauser, M.D. (2001). Amodal completion of acoustic signals by a nonhuman primate. *Nat. Neurosci.* *4*, 783–784.
- Nelken, I., Fishbach, A., Las, L., Ulanovsky, N., and Farkas, D. (2003). Primary auditory cortex of cats: feature detection or something else? *Biol. Cybern.* *89*, 397–406.
- Newsome, W.T., Britten, K.H., and Movshon, J.A. (1989). Neuronal correlates of a perceptual decision. *Nature* *341*, 52–54.
- O'Connor, K.N., Barruel, P., and Sutter, M.L. (2000). Global processing of spectrally complex sounds in macaques (*Macaca mullata*) and humans. *J. Comp. Physiol. [A]* *186*, 903–912.
- O'Connor, K.N., Petkov, C.I., and Sutter, M.L. (2005). Adaptive stimulus optimization for auditory cortical neurons. *J. Neurophysiol.* *94*, 4051–4067.
- Patterson, R.D., Robinson, K., Holdsworth, J., McKeown, D., Zhang, C., and Allerhand, M.H. (1992). Complex sounds and auditory images. In *Auditory Physiology and Perception*, Y. Cazals, L. Demany, and K. Horner, eds. (Oxford: Pergamon), pp. 429–446.
- Patterson, R.D., Allerhand, M.H., and Giguere, C. (1995). Time-domain modeling of peripheral auditory processing: a modular architecture and a software platform. *J. Acoust. Soc. Am.* *98*, 1890–1894.
- Paxinos, G., Huang, X., and Toga, A.W. (2000). *The Rhesus Monkey Brain in Stereotaxic Coordinates* (San Diego: Academic Press).
- Pessoa, L., and De Weerd, P. (2003). *Filling-in: From Perceptual Completion to Cortical Reorganization* (Oxford, New York: Oxford University Press).
- Petkov, C.I., O'Connor, K.N., and Sutter, M.L. (2003). Illusory sound perception in macaque monkeys. *J. Neurosci.* *23*, 9155–9161.
- Petkov, C.I., Kang, X., Alho, K., Bertrand, O., Yund, E.W., and Woods, D.L. (2004). Attentional modulation of human auditory cortex. *Nat. Neurosci.* *7*, 658–663.
- Pickett, J. (1959). Backward masking by an intense burst of noise. *Journal of the Acoustical Society of America* *31*, 127.
- Pressnitzer, D., Meddis, R., Delahaye, R., and Winter, I.M. (2001). Physiological correlates of comodulation masking release in the mammalian ventral cochlear nucleus. *J. Neurosci.* *21*, 6377–6386.
- Pugh, K.R., offywitz, B.A., Shaywitz, S.E., Fulbright, R.K., Byrd, D., Skudlarski, P., Shankweiler, D.P., Katz, L., Constable, R.T., Fletcher, J., et al. (1996). Auditory selective attention: an fMRI investigation. *Neuroimage* *4*, 159–173.
- Rauschecker, J.P., and Tian, B. (2004). Processing of band-passed noise in the lateral auditory belt cortex of the rhesus monkey. *J. Neurophysiol.* *91*, 2578–2589.
- Recanzone, G.H. (2000). Response profiles of auditory cortical neurons to tones and noise in behaving macaque monkeys. *Hear. Res.* *150*, 104–118.
- Recanzone, G.H., Guard, D.C., and Phan, M.L. (2000). Frequency and intensity response properties of single neurons in the auditory cortex of the behaving macaque monkey. *J. Neurophysiol.* *83*, 2315–2331.
- Rhode, W.S., Geisler, C.D., and Kennedy, D.T. (1978). Auditory nerve fiber response to wide-band noise and tone combinations. *J. Neurophysiol.* *41*, 692–704.
- Rossi, A.F., and Paradiso, M.A. (1996). Temporal limits of brightness induction and mechanisms of brightness perception. *Vision Res.* *36*, 1391–1398.
- Schreiner, C. (1980). Encoding of alternating acoustical signals in the medial geniculate body of guinea pigs. *Hear. Res.* *3*, 265–278.

- Schreiner, C.E., and Sutter, M.L. (1992). Topography of excitatory bandwidth in cat primary auditory cortex: single-neuron versus multiple-neuron recordings. *J. Neurophysiol.* *68*, 1487–1502.
- Schreiner, C.E., Mendelson, J.R., and Sutter, M.L. (1992). Functional topography of cat primary auditory cortex: representation of tone intensity. *Exp. Brain Res.* *92*, 105–122.
- Schreiner, C.E., Read, H.L., and Sutter, M.L. (2000). Modular organization of frequency integration in primary auditory cortex. *Annu. Rev. Neurosci.* *23*, 501–529.
- Sivonen, P., Maess, B., and Friederici, A.D. (2006). Semantic retrieval of spoken words with an obliterated initial phoneme in a sentence context. *Neurosci. Lett.* *408*, 220–225.
- Slaney, M. (1993). An efficient implementation of the Patterson-Holdsworth auditory filter bank. Apple Computer Technical Report *35*, 1–42.
- Slaney, M. (1998). Auditory toolbox, version 2. Interval Research Corporation Technical Report *10*, 1–52.
- Steinschneider, M., Reser, D., Schroeder, C.E., and Arezzo, J.C. (1995). Tonotopic organization of responses reflecting stop consonant place of articulation in primary auditory cortex (A1) of the monkey. *Brain Res.* *674*, 147–152.
- Sugita, Y. (1997). Neuronal correlates of auditory induction in the cat cortex. *Neuroreport* *8*, 1155–1159.
- Sutter, M.L. (2005). Spectral processing in the auditory cortex. *Int. Rev. Neurobiol.* *70*, 253–298.
- Walton, J.P., Frisina, R.D., Ison, J.R., and O'Neill, W.E. (1997). Neural correlates of behavioral gap detection in the inferior colliculus of the young CBA mouse. *J. Comp. Physiol. [A]* *181*, 161–176.
- Wang, X., Lu, T., Snider, R.K., and Liang, L. (2005). Sustained firing in auditory cortex evoked by preferred stimuli. *Nature* *435*, 341–346.
- Warren, R.M. (1970). Perceptual restoration of missing speech sounds. *Science* *167*, 392–393.
- Warren, R.M., Obusek, C.J., and Ackroff, J.M. (1972). Auditory induction: perceptual synthesis of absent sounds. *Science* *176*, 1149–1151.
- Warren, R.M., Wrightson, J.M., and Puretz, J. (1988). Illusory continuity of tonal and infratonal periodic sounds. *J. Acoust. Soc. Am.* *84*, 1338–1342.
- Warren, R.M., Bashford, J.A., Jr., Healy, E.W., and Brubaker, B.S. (1994). Auditory induction: reciprocal changes in alternating sounds. *Percept. Psychophys.* *55*, 313–322.
- Wrightson, J.M., and Warren, R.M. (1981). Incomplete auditory induction of tones alternated with noise: Effects occurring below the pulsation threshold. *J. Acoust. Soc. Am.* *69*, s105–s106.