Separate mechanisms for audio-tactile pitch and loudness interactions

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INTRODUCTION

We are constantly bombarded by myriad sensory signals and are tasked with sorting these for useful information about our environment. Signals conveyed by our sensory systems interact in time and space, affecting not only when and where we perceive meaningful events, but even the identity and content of these occurrences. Familiar examples of such perceptual interactions include the ventriloquism illusion (in which viewing an object biases the perceived location of a separate sound source; Thomas, 1941) and the McGurk effect (in which seeing lip movements biases the perception of simultaneously heard speech sounds; McGurk and MacDonald, 1976). A major goal in perceptual neuroscience is to understand how signals from different sensory modalities are combined to produce stable and coherent perceptual experiences.

Multisensory interactions are complex and varied, and the establishment of multisensory neural mechanisms can depend on many factors. Because we rarely perceive the world through a single modality, we develop representations that are linked across our senses; the strengths of these links reflect the history of our multisensory experiences. For instance, objects we palpate must be in close proximity to our bodies. As a result, we typically can hear sounds generated during our haptic interactions with the objects. Such correlated sensory experiences, accrued over a lifetime of co-stimulation, may pattern the neural mechanisms underlying audio-tactile interactions in simple event detection (Gescheider and Niblette, 1967; Ro et al., 2009; Tajadura-Jimenez et al., 2009; Wilson et al., 2009; Occelli et al., 2010; Wilson et al., 2010b) and event counting (Hotting and Roder, 2004; Bresciani et al., 2005; Bresciani and Ernst, 2007).

Shared neural representations can also be highly specific. Because we can simultaneously experience environmental oscillations through audition and touch (transduced by receptors in the basilar membrane and in the skin, respectively), we previously reasoned that the two sensory systems might interact in the spectral analysis of vibrations. We tested this and found that auditory tones and noise stimuli indeed systematically influence tactile frequency perception (Yau et al., 2009b), although tones do not affect tactile intensity judgments. These linked audio-tactile frequency representations may underlie our capacity to perceive textures (Lederman, 1979; Jousma and Hari, 1998; Guest et al., 2002; Yau et al., 2009a), to appreciate music (Musachia and Schroeder, 2009; Soto-Faraco and Deco, 2009), and even to comprehend speech (Gick and Derrick, 2009).

In the current study we tested the hypothesis that audio-tactile perceptual interactions are reciprocal in nature by examining how tactile distractors affect auditory tone analysis. We measured participants’ ability to discriminate the frequency or intensity of auditory pure tone stimuli in the presence or absence of simultaneous tactile vibrations. We further characterized the sensitivity of these effects to the relative timing between the auditory tones and tactile distractors. The results reported here, combined with our previous findings (Yau et al., 2009b), provide a comprehensive view of audio-tactile interactions in pitch and loudness perception.
MATERIALS AND METHODS

PARTICIPANTS
All testing procedures were performed in compliance with the policies and procedures of the Institutional Review Board for Human Use of the University of Chicago. All subjects were paid for their participation. Subjects reported normal tactile and auditory sensitivity and no history of neurological disease. Subjects did not participate in all of the experiments. Thirteen subjects (five males and eight females; mean age = 20 ± 1.5 years) participated in the auditory frequency discrimination experiment with the 200-Hz standard. Eleven subjects (four males and seven females; mean age = 20.5 ± 2.6 years) participated in the auditory frequency discrimination experiment with the 400-Hz standard. Twenty-four subjects (12 males and 12 females; mean age = 19.9 ± 2.0 years) participated in the auditory intensity discrimination experiment. Ten subjects (two males and eight females; mean age = 19.8 ± 0.4 years) participated in the frequency discrimination experiment with the timing (synchrony) manipulation. Ten subjects (five males and five females; mean age = 20.7 ± 2.5 years) participated in the intensity discrimination experiment with the timing manipulation.

STIMULI

Auditory stimuli
Auditory stimuli were generated digitally and converted to analog signals using a digital to analog card (PCI-6229, National Instruments, Austin, TX, USA; sampling rate = 20 kHz). Stimuli were delivered binaurally via noise isolating in-ear earphones (ER6i, Etymotic Research, Elk Grove Village, IL, USA), as these allowed participants to wear noise-attenuating earmuffs (847NST, Bilsom, Winchester, VA, USA).

Tactile stimuli
The tactile distractors consisted of sinusoids that were equated in perceived intensity at different frequencies using a two-alternative forced-choice (2AFC) tracking procedure (Yau et al., 2009b). In a pilot experiment, subjects matched the intensities of stimuli whose amplitudes were initially set by the experimenter. The average intensities determined by the subjects were then used with all subjects in the main experiments. All distractors (100-, 200-, 300-, 400-, and 600-Hz) were suprathreshold and were presented to each participant’s index finger (on his or her left hand) at subjectively matched amplitudes of 6.8, 3.6, 3.2, 1.8, and 2.1 μm, respectively. The fact that the amplitude of 400-Hz stimulus was lower than that of the 300-Hz stimulus is somewhat surprising given that thresholds of Pacinian afferents (the mechanoreceptor population sensitive to vibration) are higher at 400 Hz than at 300 Hz (Muniak et al., 2007). However, the slope of the rate intensity function is steeper at 400 Hz than at 300 Hz (Muniak et al., 2007), so the overall Pacinian response may be greater at 400 Hz than at 300 Hz at suprathreshold intensities. Tactile stimuli delivered to the finger at these amplitudes are unlikely to be detected via bone-conduction (Bekesy, 1939; Dirks et al., 1976). Distractors were delivered along the axis perpendicular to the skin surface by a steel-tipped plastic stylus mounted on a Mini-shaker motor (Type 4810, Bruel & Kjer, Skodsborgvej, Nærum, Denmark). The probe had a flat, circular (8 mm diameter) contact surface. The probe tip was indented 1 mm into the skin to ensure contact throughout the stimulus presentation. The motor was equipped with an accelerometer (Type 8702B50M1, Kistler Instrument Corporation, Amherst, NY, USA) with a dynamic range of ±50 g. The accelerometer output was amplified and conditioned using a piezotron coupler (Type 5134A, Kistler Instrument Corporation, Amherst, NY, USA). This signal was then digitized (PCI-6229, National Instruments, Austin, TX, USA; sampling rate = 20 kHz) and read into a computer.

We took great care in ensuring that the tactile distractors were inaudible to guarantee that observed audio-tactile interactions truly reflected multimodal interactions. We first required subjects to wear noise isolating in-ear headphones and noise-attenuating earmuffs. Furthermore, the tactile stimulator was housed in a custom-built noise attenuation chamber to further attenuate sounds emanating from the stimulator (for a complete description, see Yau et al., 2009b).

AUDITORY FREQUENCY DISCRIMINATION

Auditory frequency discrimination with a 200-Hz standard
Participants sat facing the tactile stimulator with their left arm and hand comfortably resting in a half-cast and hand-mold. The restraints were mounted on a height-adjustable vertical stage, which allowed the stimulator to be reliably repositioned for each participant. When the participant was situated, the stimulator was gently lowered onto the distal pad of the participant’s index finger and the experiment began. Participants were tested using a 2AFC design (Figure 1A). On each trial, a pair of auditory tones, equated in perceived intensity, was presented through the headphones and the participant judged which of the two tones was higher in frequency. The tones were each presented for 1 s and were separated by a 1-s inter-stimulus interval. One interval always contained a 200-Hz tone (standard stimulus); the frequency of the tone presented during the other interval (comparison stimulus) ranged from 195 to 205 Hz. The frequency of the comparison stimulus and the stimulus interval in which it was presented were randomized across trials. On most trials, a tactile distractor was presented at the same time as the auditory comparison stimulus. Participants were instructed to ignore the tactile distractors. On the remaining trials, no distractor was presented allowing us to establish a baseline against which we could compare performance achieved in the presence of the tactile distractors. The frequency of the tactile distractor was 100-, 200-, 300-, 400-, or 600-Hz. Twenty behavioral observations were obtained for every combination of auditory comparison stimulus and tactile distractor over 10 experimental runs distributed across two to three sessions. Participants were allowed time to rest between trial blocks. No feedback was provided.

Equating auditory stimulus intensity
Two aspects of stimulus design were implemented to ensure participants did not rely on intensity information to perform the frequency discrimination task. In pilot experiments, we equated the perceived intensity of auditory stimuli at different frequencies using a 2AFC tracking procedure. The intensities of the tactile stimuli were determined using a similar procedure. On each trial, participants were presented sequentially with two 1-s stimuli separated by a 1-s inter-stimulus interval. One stimulus (the standard) was always a 200-Hz, 60.4 dB SPL (suprathreshold) tone; the other stimulus (the comparison) was a tone at one of the stimulus frequencies (195,
197, 198, 199, 201, 202, 203, and 205 Hz) tested in the frequency discrimination experiments. Participants reported which stimulus was more intense. If the participant judged the standard as more intense, the amplitude of the comparison stimulus increased on the following trial. Conversely, if the participant judged the comparison as more intense, the comparison amplitude was reduced on the following trial. The session concluded when the change in the amplitude of the comparison stimulus reversed three times. The geometric mean of the comparison stimulus amplitudes on the last 10 trials of the session was then computed. Three such measurements were recorded and averaged. The resulting mean was the stimulus amplitude at each comparison frequency that was perceived to be equally intense as a 200-Hz, 60.4 dB SPL tone. To further ensure that participants made judgments using only frequency information in the frequency discrimination experiments (and not intensive cues), the actual stimulus amplitudes used during the frequency discrimination experiments were randomly jittered on a trial-by-trial basis (the maximum jitter on any given trial was 25% of the subjectively matched amplitude).

Auditory frequency discrimination with a 400-Hz standard
The procedure was identical to that used in the auditory frequency discrimination experiment with the 200-Hz standard except that the frequency of the standard tone was 400 Hz and the frequencies of comparison stimuli were: 390, 394, 396, 398, 402, 404, 406, and 410 Hz. The amplitudes of the comparison tones were determined by equating the perceived intensity of the comparison stimuli to that of the 400-Hz, 41.72 dB SPL standard.

AUDITORY INTENSITY DISCRIMINATION
In this experiment, we wished to determine whether tactile distractors influence auditory intensity perception. In a 2AFC design, participants judged which of two sequentially presented auditory tones, equated in frequency (200 Hz), was more intense (Figure 2A). The standard amplitude was 67.2 dB SPL and the comparison amplitudes ranged from 64.8 to 69.5 dB SPL; these amplitudes were chosen because they are clearly audible but not uncomfortably loud and overlapped those used in our previous study (Yau et al., 2009b). On most trials, a tactile distractor (200- or 600-Hz) was presented at the same time as the comparison stimulus. The 200- and 600-Hz distractors were presented at four amplitude levels, each equated in perceived intensity. Amplitudes ranged from 0.9 to 3.6 and 0.5 to 2.1 μm for the 200- and 600-Hz distractors, respectively.

DISCRIMINATION EXPERIMENTS WITH TIMING MANIPULATION
In these experiments, we wished to determine the extent to which synchronous presentation of tactile and auditory stimuli was necessary for the former to affect the perception of the latter. The sensitivity of multisensory processes to the relative timing of their component signals can reveal the underlying mechanisms of cross-modal signal integration (Bensmaia et al., 2006; Wilson et al., 2009; Yau et al., 2009b). Temporal coincidence is a primary cue used by the nervous system to determine whether sensory signals should be combined (Burr et al., 2009). Sensory events that occur simultaneously or in close temporal proximity likely emanate from a common source, while stimuli separated by longer intervals likely represent distinct events (Stein and Meredith, 1993). Multisensory interplay may result from co-stimulation in two modalities in the absence of any particular relation between the stimuli, relying on a non-specific mechanism such as rapid alerting or arousal (Driver and Noesselt, 2008). Multisensory interactions mediated by the convergence of sensory signals onto common neuronal populations may be specific to stimulus parameters like temporal frequency (Yau et al., 2009b) and (with respect to time) may only depend on the degree of temporal overlap between the signals (Stein and Meredith, 1993). In timing experiments requiring subjects to discriminate auditory frequency, three frequencies of tactile distractors were tested (100, 200, and 600 Hz). Timing experiments requiring intensity judgments employed two distractor frequencies (200 and 600 Hz). In all timing experiments, distractor amplitudes were identical to those used in the main frequency discrimination experiments. The timing experiments (Figures 3A,B) were similar to the main frequency and intensity discrimination experiments except that, on a subset of trials, the onset of the tactile distractors was 250 ms before and its offset 250 ms after the onset and offset of the auditory stimuli, respectively (the total distractor duration was 1500 ms). Critically, the overlap in the duration of auditory tones and tactile distractors was maintained (1000 ms).

DATA ANALYSIS

Psychometric functions
To quantify participants’ ability to discriminate auditory frequency we fit the following psychometric function to the data obtained from each participant:

\[ p(f_i > f_s) = \frac{1}{1 + e^{-\frac{f_i - \mu}{\sigma}}} \]

where \( p(f_i > f_s) \) is the proportion of trials a comparison tone with frequency \( f_i \) was judged to be higher in frequency than the standard stimulus \( f_s = 200 \) or 400 Hz), \( \mu \) and \( \sigma \) are free parameters corresponding to estimates of the participant’s bias and sensitivity, respectively. The bias indicates the point of subjective equality (PSE) while the sensitivity parameter denotes the change in frequency (with respect to the standard) that the participant could detect 73% of the time. The resulting sigmoidal functions range from 0 to 1. Participants’ ability to discriminate auditory intensity was similarly quantified using a psychometric function analogous to that shown in Eq. 1. These psychometric functions accurately captured discrimination performance for all conditions tested (mean correlation = 0.97, SEM = 0.002).

Statistical tests
For the frequency discrimination experiments, we first determined whether the presentation of tactile distractors significantly affected the average estimates of bias and sensitivity. We tested the effect of the distractors using a repeated-measures ANOVA (ANOVA_{RM}), with distractor condition, including the baseline condition, as the within-subjects factor. If this test was significant \( (p < 0.05) \), we then conducted an ANOVA_{RM} to test whether the effect of the distractors was significantly modulated by distractor frequency, with distractor frequency (excluding the baseline condition) as the within-subjects factor. If this ANOVA_{RM} revealed a significant main effect of distractor frequency, we performed a two-tailed paired \( t \)-test comparing
the bias (or sensitivity) estimates derived from the 100- and 600-Hz distractor conditions. For the intensity discrimination experiment, we first conducted a one-way ANOVA with distractor condition, including the baseline condition, as the within-subjects factor. If this test was significant ($p < 0.05$), we conducted a two-way ANOVA to test for effects of distractor frequency, distractor intensity, and their interaction. If this ANOVA revealed a significant main effect of distractor intensity, we performed a two-tailed paired $t$-test comparing the bias (or sensitivity) estimates derived from the lowest and highest intensity distractor conditions. For the timing manipulation experiments, we first conducted a one-way ANOVA with distractor condition, including the baseline condition, as the within-subjects factor. If this test was significant ($p < 0.05$), we tested for specific main effects and interactions using a two-way ANOVA.

**RESULTS**

**EFFECT OF TACTILE DISTRACTORS ON AUDITORY FREQUENCY DISCRIMINATION**

Tactile distractors systematically influenced performance on the auditory frequency discrimination task (Figure 1). In experiments using the 200-Hz standard tone, distractors biased the perceived frequency of simultaneously heard tones [ANOVA, $F(5,71) = 6.8, p < 10^{-4}$; see Figure S1 in Supplementary Material for group psychometric functions for all experiments]. The magnitude of the bias effect depended on distractor frequency [$F(4,59) = 7.9, p < 10^{-3}$] and was larger when the frequency of the distractor was lower than that of the auditory stimuli (Figure 1B, red trace). The average PSE (bias) estimated from the discrimination performance with the 100-Hz distractor was significantly higher than the average PSE obtained with the 600-Hz distractor [$t(12) = 3.0, p = 0.01$]. We confirmed the audio-tactile pitch interaction by assessing the influence of the same tactile distractors on participants’ ability to discriminate tones in a different frequency range, centered on 400 Hz (cyan trace). Tactile distractors also biased the perceived frequency of these tones [$F(4,59) = 3.9, p < 0.01$] and the strength of the bias again depended on distractor frequency [$F(4,49) = 5.6, p < 0.01$]. The average PSE estimated with the 100-Hz distractor again differed significantly from the average value estimated with the 600-Hz distractor [$t(10) = 2.7, p = 0.02$]. In both frequency discrimination experiments, the perceived frequency of the auditory tones tended to be pulled toward the frequency of the tactile distractors. This bias effect appears to be frequency-specific, as the set of distractors that substantially biased auditory discrimination performance using the 400-Hz standard tone was expanded and shifted in frequency compared to the most effective distractors found using the 200-Hz standard. Critically, in both experiments, we observed greater bias effects when the frequency of the distractor was lower than that of the tones. This pattern was evident in individual participants’ data (Figure S2 in Supplementary Material) and across the pooled sample. Notably, this pattern mirrors the effects of auditory distractors on tactile frequency discrimination (Yau et al., 2009). Tactile distractors did not significantly affect estimates of sensitivity in the frequency discrimination experiments (Figure 1C) [$F(5,71) = 2.0, p = 0.09$ and $F(5,59) = 1.1, p = 0.40$ for the experiments using the 200- and 400-Hz standards, respectively].

**EFFECT OF TACTILE DISTRACTORS ON AUDITORY INTENSITY DISCRIMINATION**

Tactile distractors influenced performance on the auditory intensity discrimination task (Figure 2). The simultaneous presentation of tactile distractors led to an increase in perceived tone loudness.
average PSE estimated with the most intense distractor [i.e., the more intense distractor increased perceived loudness to a greater extent; $t(23) = 7.4, p < 10^{-4}$] and $t(23) = 4.8, p < 10^{-4}$ for the 200- and 600-Hz distractors, respectively]. The intensity $\times$ frequency interaction on bias estimates was not significant [$F(3,182) = 0.17, p = 0.91$]. Tactile distractors generally did not affect estimates of perceptual sensitivity in the intensity discrimination experiments (Figure 2C). While the main effect of distractor frequency on sensitivity estimates was marginally significant [$F(1,185) = 5.3, p = 0.03$], the main effect of distractor intensity and the intensity $\times$ frequency interaction failed to achieve statistical significance ($p$ values $>0.05$).

**EFFECT OF DISTRACTOR TIMING ON PITCH AND LOUDNESS INTERACTIONS**

In auditory frequency and intensity discrimination experiments, we compared the effect of distractors that were presented synchronously with the auditory tones to distractors that overlapped the tones but began earlier and terminated later. Audio-tactile pitch and loudness interactions differed in their sensitivity to this timing manipulation (Figure 3). As was the case in the main frequency and intensity discrimination experiments, the influence of auditory distractors was limited to estimates of bias (see Figure S3 in Supplementary Material for average sensitivity estimates in the timing manipulation experiments). In the frequency discrimination task (Figure 3C), tactile distractors significantly biased auditory judgments in a frequency-dependent manner [$F(2,54) = 5.9, p < 0.01$]. Critically, this effect did not depend on synchronous timing between the auditory tones and tactile distractors [$F(1,54) = 2.1, p = 0.18$], and there was no significant frequency $\times$ timing interaction [$F(2,52) = 0.05, p = 0.95$]. We previously found the auditory influence on tactile frequency perception to be similarly tolerant to stimulus onset (and offset) asynchrony (Yau et al., 2009b). In contrast, audio-tactile loudness interactions were sensitive to stimulus timing (Figure 3D). Although tactile distractors biased auditory intensity judgments when the stimuli coincided, this effect was abolished when the onset and offset timing was disrupted [$F(1,37) = 26.7, p < 10^{-4}$]. As in the main intensity discrimination experiment, the bias effect of tactile distractors did not differ across distractor frequency [$F(1,37) = 0.62, p = 0.45$] and the frequency $\times$ timing interaction was not significant [$F(1,34) = 1.5, p = 0.25$].

**DISCUSSION**

In a series of psychophysical experiments, we assessed the influence of tactile distractors on participants’ ability to discriminate the frequency and intensity of auditory tones. We also determined the dependence of these perceptual interactions on the relative timing between the auditory and tactile signals. Tactile distractors systematically biased auditory frequency perception (Figure 1): Distractors at frequencies lower than that of the auditory tones induced larger bias effects than distractors at higher frequencies. Tactile distractors also biased auditory perception of intensity (Figure 2). The magnitude of this effect scaled with distractor intensity, but did not vary with distractor frequency. We also found that audio-tactile interactions in the frequency and intensity domains differ in their sensitivity to stimulus timing: Breaking the correspondence between onset and offset times of the tones and distractors disrupted loudness interactions but had no effect on pitch interactions (Figure 3).
Inseparability of auditory and tactile frequency representations, regardless of the attended modality, implies a supramodal operator for spectral analysis. Supramodal operators may also mediate perception of object shape (Amedi et al., 2001, 2002; Lacey et al., 2009), motion (Blake et al., 2004; Ricciardi et al., 2007), and microgeometric features (Zangaladze et al., 1999; Merabet et al., 2004). These studies support a metamodal view of brain organization, in which cortical areas perform particular operations regardless of input modality (Pascual-Leone and Hamilton, 2001).

Even in a shared neural representation of frequency, auditory and tactile inputs may not be conveyed with the same precision. This may explain why tactile distractors did not affect estimates of auditory sensitivity, despite robust changes in tactile sensitivity estimates with auditory distractors (Yau et al., 2009b). Changes in the sensitivity parameter (i.e., the slope of the psychometric curve) arise when a distractor unequally affects individual data points in a given psychometric function. In the current study, the range of auditory comparison frequencies was very narrow (spanning 10 and 20 Hz for the curves centered on 200 and 400 Hz, respectively). As a result, each tactile distractor affected all of the data points in individual psychometric functions equally, resulting in a uniform shift of the entire function and thus of a biasing effect with no concomitant effect on the slopes. In contrast, the range of tactile comparison frequencies we tested in our previous report was limited to 100-, 200-, and 600-Hz distractors, respectively. The black bar shows the PSE in the baseline condition. Error bars indicate SEM. The frequency bias effects were comparable across the synchronous and asynchronous timing conditions. (D) Mean bias estimates on the intensity discrimination task as a function of stimulus timing. Conventions as in (C). Intensity bias effects depended on synchronous presentation of auditory stimuli and tactile distractors.

In all of our frequency discrimination experiments, the influence of distractors was greatest when they were lower in frequency than the test stimuli, regardless of the modality participants attended or ignored. This response pattern is reminiscent of the finding that auditory stimuli more effectively mask (auditory) stimuli at higher frequencies than they do stimuli at lower frequencies (Moore, 2003). Accordingly, auditory filters estimated from masking studies using the notched-noise method are asymmetric at center-frequencies ranging from 100 to 800 Hz (Moore et al., 1990), a result previously ascribed to the biomechanics of the basilar membrane. That this pattern also describes the interplay between audition and touch raises another possibility: The asymmetry potentially reflects the tuning properties of auditory cortical neurons, some of which have been shown to receive both auditory and tactile inputs (Fu et al., 2003). Quite possibly, these cortical ensembles, whose involvement in acoustic frequency analysis is unchallenged, also underlie tactile frequency perception. The finding that perceptual interactions in temporal frequency are insensitive to timing disruptions is consistent with sensory-level convergence. This view is further supported by the fact that auditory and tactile stimuli exhibit frequency-dependent interactions even when one of the inputs alone fails to evoke an explicit pitch percept (Yau et al., 2009b). Indeed, our results support the hypothesis that spectral analysis of auditory and tactile inputs is mediated by a common mechanism:
(Yau et al., 2009b) was substantially broader (spanning 200 and 400 Hz for the curves centered on 200 and 400 Hz, respectively). As a result, auditory distractors affected the data points comprising individual psychometric functions to different degrees, thereby changing the slopes of the curves while also shifting the PSEs, and thus affecting both sensitivity estimates and bias estimates. Thus, the presence or absence of audio-tactile pitch interactions on the sensitivity (slope) estimates may be determined by the precision of the frequency representations. While auditory and tactile frequency representations may be integrated centrally, their precision appears to differ and may reflect differences at more peripheral stages of auditory and somatosensory processing. An obvious difference is the fact that the auditory system can exploit a place code at the receptor level (with frequency information conveyed by a receptor’s position along the basilar membrane) whereas the somatosensory system cannot.

Audio-tactile loudness interactions do not appear to be reciprocal like those in the frequency domain. Though we previously found that tactile intensity judgments were unaffected by auditory tones (Yau et al., 2009b), in the current study, tactile distractors caused auditory tones to be perceived as more intense. A number of studies have reported similar tactile enhancement of auditory loudness (Schurmann et al., 2004; Gillmeister and Eimer, 2007; Yarrow et al., 2008; Wilson et al., 2010a). The fact that audio-tactile loudness interactions do not depend on stimulus frequency hints at a nonspecific mechanism like rapid alerting or arousal. This is further supported by the sensitivity of the enhancement effect to stimulus timing. Interestingly, auditory loudness can be similarly biased by co-occurring visual stimulation (Marks et al., 2003; Odgaard et al., 2004), so intensity representations in the auditory system may be generally susceptible to non-auditory influence. Recent neurophysiological studies focusing on the role of neuronal oscillations in information processing provide a possible mechanistic explanation for both tactile and visual enhancement of auditory processing (Lakatos et al., 2007; Kayser et al., 2008). The evidence suggests that non-auditory sensory input can reset the phase of ongoing neuronal oscillations in auditory cortex. Such phase modulation results in the amplification of auditory inputs that arrive in cortex during high-excitability phases (Lakatos et al., 2007; Kayser et al., 2008). This mechanism may account for the timing dependence of the loudness interactions we observe, as well as previous neuroimaging results that reveal supra-additive integration of tactile and auditory stimulation in auditory cortex that display similar timing dependencies (Kayser et al., 2005).

The magnitude of the intensity bias did not vary with distractor frequency in our experiment, although loudness interactions have been shown to be frequency-dependent under certain conditions. For example, frequency-specific integration patterns are evident when subjects judge the combined loudness of concurrent auditory and tactile stimulation (Wilson et al., 2010a). Similarly, interactions between audition and touch can be frequency-dependent in detection paradigms (Ro et al., 2009; Tajadura-Jimenez et al., 2009; Wilson et al., 2009, 2010b). Although it is not immediately clear why we failed to see frequency-specific loudness interactions, a critical difference in these studies is that subjects deployed attention across audition and touch when making perceptual judgments, while participants directed attention to a single modality (and ignored the other) in our design. This difference potentially highlights the role of attention in giving rise to certain types of multisensory interactions (Senkowski et al., 2008; Talsma et al., 2010). Perhaps the separate neural mechanisms supporting audio-tactile frequency and intensity processing may be linked by attention signals. For instance, modulation of neuronal oscillations has been recently proposed as a mechanism for stimulus selection and binding through attention (Lakatos et al., 2009; Schroeder et al., 2010). Quite possibly, when attention is simultaneously deployed across touch and audition (rather than to one modality, as in our experiments), oscillatory activity in the neural substrates governing audio-tactile frequency and intensity processing may be synchronized and functionally linked, which would result in the frequency-specific interactions others have reported.

Multisensory studies often emphasize the specificity of sensory interactions (Stein and Meredith, 1993; Soto-Faraco and Deco, 2009; Sperdin et al., 2010). Factors such as the relative timing, location, and strength of sensory inputs can determine the degree of cross-modal interplay. Our work indicates that audio-tactile pitch interactions are highly specific to the relative frequencies of the sensory inputs, do not depend on specific stimulus onset timing, and are relatively insensitive to changes in stimulus intensity (see Yau et al., 2009b). In contrast, loudness interactions (which are unidirectional) do not appear to depend on stimulus frequency, are sensitive to stimulus timing, and can be modulated by changes in stimulus intensity. Critically, we have not investigated the spatial sensitivities of these perceptual interactions. Although some studies have argued that audio-tactile interactions are insensitive to spatial register (Murray et al., 2005; Zampini et al., 2007), others suggest that interactions can depend on the body part stimulated and its distance relative to the auditory stimulus (Tajadura-Jimenez et al., 2009). In our experiments, we always presented tactile stimuli to each participant’s left index finger and auditory stimuli binaurally through headphones. Our observed effects could potentially be strengthened or weakened in other alignments (although it is unclear how the frequency-dependent effects might be affected). Notably, Gillmeister and Eimer (2007) tested the spatial sensitivities of audio-tactile loudness interactions and found that spatial registry did not affect intensity ratings.

A large body of neuroimaging and neurophysiology studies have provided candidate regions for audio-tactile convergence (Musacchia and Schroeder, 2009; Soto-Faraco and Deco, 2009). We previously speculated that shared auditory and tactile frequency representations might reside in the caudomedial belt area (area CM), part of the auditory association cortex. We based our speculation on the response characteristics of area CM neurons (Recanzone, 2000a; Kajikawa et al., 2005) and their anatomical connections (Cappe and Barone, 2005; Hackett et al., 2007a; Smiley et al., 2007). However, neurons in area CM may be better suited for spatial localization (Rauschecker and Tian, 2000; Recanzone, 2000b) and frequency tuning in area CM may be too poor to support spectral analysis (Lakatos et al., 2005; Kayser et al., 2009). The caudolateral belt area (area CL) may be a better candidate, given its more refined tonotopic organization (Foxe, 2009). Additionally, audio-tactile interactions are thought to occur throughout primary and association auditory cortices (Foxe et al., 2000, 2002; Schroeder et al., 2001; Kayser et al., 2005; Murray et al., 2005;
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FIGURE S1 | Psychometric functions fit to data acquired in the frequency, intensity, and timing experiments. The data show the proportion of trials in which the frequency (intensity) of a given comparison stimulus is judged to be higher (louder) than that of a standard tone. Trace color indicates distractor condition. Error bars indicate SEM. (A) Frequency discrimination experiment with the 200-Hz standard. (B) Frequency discrimination experiment with the 400-Hz standard. (C) Intensity discrimination experiments. (D) Timing experiment requiring frequency judgments. (E) Timing experiment requiring intensity judgments.

FIGURE S2 | Psychometric functions fit to individual subjects in the frequency discrimination experiments. The data show the proportion of trials in which the frequency of a given comparison stimulus is judged to be higher than that of the standard tone. Trace color indicates distractor condition. Distractors at frequencies lower than that of the test stimuli induced larger bias effects in experiments using the (A) 200-Hz standard and (B) the 400-Hz standard.
Figure S3 | Sensitivity estimates for the frequency and intensity discrimination tasks with timing manipulations. Error bars indicate SEM. (A) Timing experiments requiring frequency judgments. There were no significant main or interaction effects on sensitivity ($p > 0.05$). (B) Timing experiments requiring intensity judgments. There were no significant main or interaction effects on sensitivity ($p > 0.05$).