



Stimulus intensity modulates multisensory temporal processing

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ABSTRACT

One of the more challenging feats that multisensory systems must perform is to determine which sensory signals originate from the same external event, and thus should be integrated or “bound” into a singular perceptual object or event, and which signals should be segregated. Two important stimulus properties impacting this process are the timing and effectiveness of the paired stimuli. It has been well established that the more temporally aligned two stimuli are, the greater the degree to which they influence one another’s processing. In addition, the less effective the individual unisensory stimuli are in eliciting a response, the greater the benefit when they are combined. However, the interaction between stimulus timing and stimulus effectiveness in driving multisensory-mediated behaviors has never been explored – which was the purpose of the current study. Participants were presented with either high- or low-intensity audiovisual stimuli in which stimulus onset asynchronies (SOAs) were parametrically varied, and were asked to report on the perceived synchrony/asynchrony of the paired stimuli. Our results revealed an interaction between the temporal relationship (SOA) and intensity of the stimuli. Specifically, individuals were more tolerant of larger temporal offsets (i.e., more likely to call them synchronous) when the paired stimuli were less effective. This interaction was also seen in response time (RT) distributions. Behavioral gains in RTs were seen with synchronous relative to asynchronous presentations, but this effect was more pronounced with high-intensity stimuli. These data suggest that stimulus effectiveness plays an underappreciated role in the perception of the timing of multisensory events, and reinforces the interdependency of the principles of multisensory integration in determining behavior and shaping perception.

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1. Introduction

Our daily environment is filled with an abundance of information that our different sensory systems utilize in order to allow us to successfully navigate the world. Despite the fact that many of the objects and events in our world are specified by information carried by multiple senses, we perceive these as singular and unified. In order to create such a unified percept, the brain must be able to “bind” information that belongs together and segregate information that should be separate. The binding process by which multisensory cues are actively synthesized – a

process that represents a component of multisensory integration – has been the subject of much study. Collectively, this work has revealed dramatic changes associated with combining information across multiple senses; changes that frequently result in substantial benefits to behavior (Amlot et al., 2003; Frassinetti et al., 2002; Lovelace et al., 2003) and striking alterations in perception (McGurk and MacDonald, 1976; The Neural Bases of Multisensory Processes, 2012; Shams et al., 2002).

To solve this “binding or causal source problem,” sensory systems rely upon the statistical properties of the different sensory signals, two of the most important of which are space and time. Multisensory (e.g., visual-auditory) stimuli that are spatially and temporally concordant tend to influence one another’s processing, and may ultimately be integrated or bound, whereas those that are discordant in space and/or time tend to not influence the processing of one another (Conrey and Pisoni, 2006; Hairston et al.,

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2005; Keetels and Vroomen, 2005; Kording et al., 2007; Macaluso et al., 2004; Powers et al., 2009; Sato et al., 2007; Stevenson et al., 2012c, 2012b; van Atteveldt et al., 2007; van Wassenhove et al., 2007; Vroomen and de Gelder, 2004; Vroomen and Keetels, 2010; Wallace and Stevenson, 2014). Furthermore, it has also been shown that stimuli that are weakly effective on their own tend to give rise to the largest gains when combined (James et al., 2009; James and Stevenson, 2012b; James et al., 2012; Ross et al., 2007a; Senkowski et al., 2011; Stein et al., 2009; Stevenson et al., 2012d; Stevenson and James, 2009). Collectively, these integrative principles make a great deal of ethological sense, in that spatial and temporal proximity typically signal a common source, and in that it is highly adaptive to accentuate multisensory gain when each of the sensory signals is weak or ambiguous when presented alone.

Recently, a number of studies have focused on how temporal factors influence the nature of human multisensory perceptual judgments (Billock and Tsou, 2014; Conrey and Pisoni, 2006; Grant et al., 2004; Macaluso et al., 2004; van Wassenhove et al., 2007; Vroomen and de Gelder, 2004; Vroomen and Keetels, 2010). One useful construct associated with this work is the concept of a multisensory temporal binding window, defined as the epoch of time within which multisensory stimuli can influence one another's processing. The window within which multisensory influences can be demonstrated appears to be surprisingly broad, spanning several hundred milliseconds (Hillock et al., 2011; Hillock-Dunn and Wallace, 2012; Powers et al., 2012, 2009; Sarko et al., 2012; Stevenson et al., 2013b). In addition, these studies and others have revealed a number of other salient characteristics concerning multisensory temporal acuity. These include that it: 1) has a great deal of individual variability (Stevenson et al., 2012a; van Eijk et al., 2008), 2) differs depending upon stimulus type and task (Kasper et al., 2014; Megevand et al., 2013; Stevenson and Wallace, 2013; van Eijk et al., 2008, 2010; Vroomen and Stekelenburg, 2011), and 3) is malleable in response to perceptual training (Keetels and Vroomen, 2008; Powers et al., 2012, 2009; Schlesinger et al., 2014; Stevenson et al., 2012c, 2013a, 2013b; Vroomen et al., 2004) and across development (Hillock et al., 2011; Hillock-Dunn and Wallace, 2012; Joanne Jao et al., 2014; Johannsen and Roder, 2014; Lewkowicz, 2012; Polley et al., 2008; Shi and Muller, 2013) and aging (Bates and Wolbers, 2014; DeLoss et al., 2013; Diaconescu et al., 2013; Freiherr et al., 2013; Hugenschmidt et al., 2009; Mahoney et al., 2012, 2014; Mozolic et al., 2012; Stevenson et al., 2015).

Although these studies have illustrated the central importance of time in dictating human multisensory interactions, other studies have focused on the roles of space (Bertelson and Radeau, 1981; Ghose and Wallace, 2014; Kadunce et al., 2001; Krueger et al., 2009; Macaluso et al., 2004; Mahoney et al., 2015; Meredith and Stein, 1986, 1996; Radeau and Bertelson, 1974; Royal et al., 2009, 2010; Sarko et al., 2012; Vroomen et al., 2001; Wallace et al., 2004) and effectiveness (James and Stevenson, 2012a; James et al., 2012; Kim and James, 2010; Kim et al., 2012; Leone and McCourt, 2013; Liu et al., 2013; Nath and Beauchamp, 2011; Stevenson and James, 2009; Werner and Noppeney, 2010; Yalachkov et al., 2015). Collectively, we have learned a great deal from these studies about how stimulus-related factors shape the multisensory process, but most have treated time, space and effectiveness as independent contributors to the final multisensory product. In fact, these stimulus factors are complexly intertwined, with manipulations in one having effects upon the other. For example, simply changing the spatial location of an identical stimulus will impact the effectiveness of that stimulus given the differences in spatial acuity for different regions of space (Nidiffer et al. (2015) (in this issue); Stein et al., 1989). Reinforcing the importance of examining these interactions in more detail, recent neurophysiological studies in animal models have shown that manipulating one aspect of a

multisensory stimulus (e.g., spatial location) has consequent effects in both the temporal and effectiveness dimensions (Carriere et al., 2008; Ghose and Wallace, 2014; Krueger et al., 2009; Royal et al., 2009). Indeed, this work has suggested that stimulus effectiveness may play a more preeminent role than space and time in dictating multisensory interactions at the neural level. Extending this work into the domain of human performance, recent studies have shown a strong interdependency between time and space (Keetels and Vroomen, 2005; Krueger et al., 2009; Stevenson et al., 2012c). For example, Keetels and Vroomen (2005) showed that judgments concerning the order of auditory and visual stimuli were more precise when they were presented in disparate spatial locations. Stevenson et al. (2012d) showed that individuals were more likely to perceive auditory and visual stimuli as synchronous when they were presented at peripheral relative to foveal locations.

The present study seeks to expand upon these previous findings by examining for the first time the interaction between the temporal relationship of paired audiovisual stimuli and their relative effectiveness. Specifically, we tested the impact that manipulations of stimulus effectiveness (accomplished via changes in stimulus intensity and defined as rate of perceived synchrony) have on the ability of an individual to report audiovisual stimulus asynchrony. Our results illustrate that the relative effectiveness of the paired stimuli do in fact modulate how they are perceived in time. Furthermore, these studies revealed complex interactions between time and effectiveness in dictating the final behavioral outcome.

2. Methods

2.1. Participants

Participants included 51 Vanderbilt undergraduate students (21 male, mean age = 18.9, STD = 1, age range = 18–21) who were compensated with class credit. All recruitment and experimental procedures were approved by the Vanderbilt University Institutional Review Board (IRB). Data from participants who did not accurately report the perception of synchrony even when the auditory and visual presentation was objectively simultaneous (0 ms stimulus onset asynchrony; SOA) at least 50% of the time were excluded from further analysis (N = 5). Data from one additional subject was excluded for responding synchronous for all trials irrespective of SOA resulting in 45 subjects being included in all data analysis. The present study is part of a larger study investigating the interrelationship of stimulus effectiveness, and stimulus spatial and temporal factors (Nidiffer et al. (2015) (in this issue); Stevenson et al., 2012c).

2.2. Stimuli

Visual and auditory stimuli were presented using E-Prime version 2.0.8.79 (Psychology Software Tools, Inc; PST). Visual stimuli were presented on a Samsung Sync Master 2233RZ 120 Hz monitor arranged so that subjects were seated at a distance of 46 cm. All visual stimuli were white circles measuring 7 mm in diameter, or approximately 1° of visual angle. Visual stimuli were presented at 0° azimuth (in front of the subject) slightly above a fixation cross. Visual stimuli were presented at two luminance levels, 7.1 cd/m² (low) and 215 cd/m² (high) on a black background of 0.28 cd/m². Luminance values were verified with a Minolta Chroma Meter CS-100. Visual stimulus durations were 10 ms, with timing confirmed using a Hameg 507 oscilloscope with a photo-voltaic cell.

Auditory stimuli were presented via a speaker mounted on the

top of the monitor at 0° azimuth angled toward the participant. Speakers were mounted 2 cm, or 2.5° above the visual presentation. Auditory stimuli consisted of a frozen white-noise burst generated at 44,100 Hz with the Matlab *rand* function with a 5 ms rise and 5 ms fall cosine gate (Fig. 1b). Auditory stimulus duration was held constant at 10 ms, with timing confirmed using a Hameg 507 oscilloscope. Auditory stimuli were presented at two intensity levels, 46 dB SPL (low) and 64 dB SPL (high), with a background noise at 41 dB SPL. All sound levels were verified with a Larson Davis sound-level meter, Model 814.

Audiovisual (AV) conditions consisted of pairs of the auditory and visual stimuli described above. Presentations were always spatially coincident at 0° azimuth and intensity levels were always matched (high-high and low-low). The temporal offset of the auditory and visual stimulus pairs were parametrically varied, including stimulus onset asynchronies (SOAs) of 0, 50, 100, and 200 ms, with timing confirmed using a Hameg 507 oscilloscope. In all asynchronous cases, visual onset preceded auditory onset.

2.3. Procedure

Participants were seated inside an unlit WhisperRoom™ (SE 2000 Series) with their forehead placed against a Headspot (University of Houston Optometry) forehead rest locked in place. A chinrest and the chair height could be adjusted to the forehead rest to ensure a comfortable seating position. Participants were asked to fixate a cross located at 0° elevation and 0° azimuth at all times and were monitored by close circuit infrared cameras throughout the experiment to ensure compliance (for experimental setup, see Fig. 1a and b).

The experiment began with detailed instructions, informing the participants that they would be presented with audiovisual stimuli, and that their task was to judge whether or not the stimuli were synchronous and to respond as quickly and accurately as possible. In the case that the participants did not perceive either the auditory or visual stimulus, they were instructed to report a unisensory perception. Participants were instructed to respond via a five-button PST serial response box were 1=synchronous, 2=asynchronous, 3=visual only, 4=auditory only, and 5=no stimulus detected. Understanding of the instructions was then confirmed, and the participants were given the opportunity to clarify any questions they had about the instructions.

Each trial began with a fixation screen of 1 s displaying the fixation cross in the center of the visual field. After that time period the fixation cross disappeared and was followed by a blank screen with a duration jittered between 500 and 1000 ms, followed by the stimulus presentation. 250 ms – 284 ms after stimulus presentation, subjects were presented with the response prompt, “Was it synchronous?” Following the participant’s response and a 500 ms delay, the fixation cross re-appeared, and the subsequent trial began. For a visual depiction of a trial, see Fig. 1c. Twenty trials of each of the stimulus conditions (4 SOAs × 2 intensity levels, unisensory V × 2 intensity levels, unisensory A × 2 intensity level) were presented for a total of 240 trials.

2.4. Analysis

For each trial in which the participant perceived an audiovisual stimulus, both the response and the response time (RT) were recorded. RTs less than 250 ms were discarded, and trials with RTs over 2000 ms were excluded to eliminate outliers. Our primary analysis focused on rates of perceived synchrony. With each condition, rates of perceived synchrony were calculated as the proportion of trials in which the participant reported a synchronous percept out of all trials in that condition, in which they perceived both stimuli, or:

$$\frac{\# \text{trials perceived synchronous}}{\# \text{trials perceived synchronous} + \# \text{trials perceived asynchronous}}.$$

It should be noted that this explicitly excludes trials in which the participant did not perceive both unisensory components of the stimuli. Rates of perceived synchrony were compared across intensity levels and SOA (for a review of methods see (Stevenson et al., 2014c)).

A follow-up, exploratory analysis was conducted on RTs. First, mean RTs were calculated across trials of each condition for each individual. Mean RTs were compared across intensity levels and SOA for perceived synchronous trials. A second exploratory RT analysis was performed using a more rigorous cumulative distribution function (CDFs) analysis. CDFs for perceived synchronous trials were calculated and compared for each condition, again averaging within each participant, and then across participants. Interactions between SOA and stimulus intensity in their CDFs were then assessed using a difference of difference calculation:

$$(CDF_{(0\text{ms high})} - CDF_{(200\text{ms high})}) - (CDF_{(0\text{ms low})} - CDF_{(200\text{ms low})}).$$

3. Results

3.1. Intensity effects on unisensory performance

To ensure that changes in stimulus intensity were actually manipulating stimulus effectiveness, a two-way, repeated-measures ANOVA (intensity × modality) confirmed that high intensity presentations were more effective than low intensity stimulus presentations ($F_{(1,39)}=8.26$, $p=0.0065$). Paired sample *t*-tests also revealed that accuracies decreased for both modalities from the high-saliency to the low-saliency condition (visual, high=97.17%, low=91.67%, $t=0.965$, $p=0.045$, $d=0.014$ and auditory, high 97.08%, low=93.75% $t=0.96$, $p=0.051$, $d=0.307$).

3.2. Effects of stimulus effectiveness and temporal factors on judgments of audiovisual synchrony

Rates of perceived synchrony were measured for high- and low-intensity audiovisual stimuli presented at temporal offsets ranging from 0 ms (synchronous) to 200 ms (visual leading asynchronies). A two-way, repeated-measures ANOVA (intensity × SOA) was performed. A significant main effect was found for SOA ($F_{(3,44)}=22.411$, $p=8.03 \times 10^{-9}$, partial- $\eta^2=0.559$). As expected, rates of perceived synchrony decreased with increasing SOA (Fig. 2a and b). The main effect for intensity was marginally significant ($F_{(1,44)}=3.171$, $p=0.082$, partial- $\eta^2=0.067$). Importantly, a significant interaction effect between SOA and intensity was also observed ($F_{(3,44)}=13.219$, $p=3.24 \times 10^{-6}$, partial- $\eta^2=0.270$).

To explore the interaction between SOA and intensity on these synchrony judgments, two sets of protected, follow-up, paired-samples *t*-tests were performed. First, *t*-tests were run comparing rates of perceived synchrony across SOAs (see Table 1 for detailed statistics). Rates of perceived synchrony were significantly reduced relative to synchronous presentations at the 100 and 200 ms SOA for the high-intensity stimuli, but only at the 200 ms SOA for the low intensity stimuli (Fig. 2a and b). Second, paired *t*-tests were run across intensity levels at each SOA. A significant difference between rates of perceived synchrony between high and low intensity stimuli was observed only for the 200 ms SOA ($t=3.5507$, $p=8.48 \times 10^{-4}$, $d=1.071$), where participants were more likely to report low intensity stimuli as synchronous relative to high intensity stimuli. No other SOAs showed significant differences between the two intensities (Fig. 2c). Thus, the interaction showed

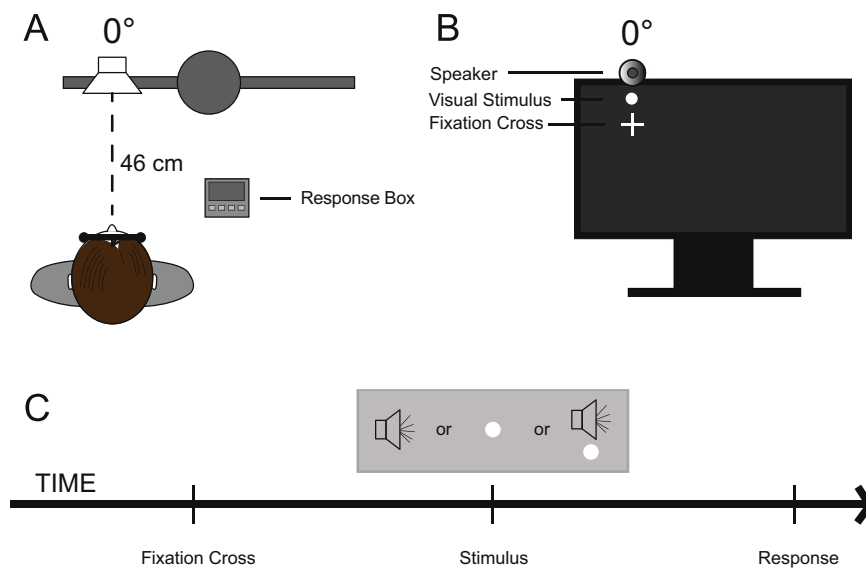


Fig. 1. A – Participant set up indicating the participant's location in relationship to the apparatus. B – Location of auditory and visual stimulus relative to fixation. C – Timeline of stimulus presentations.

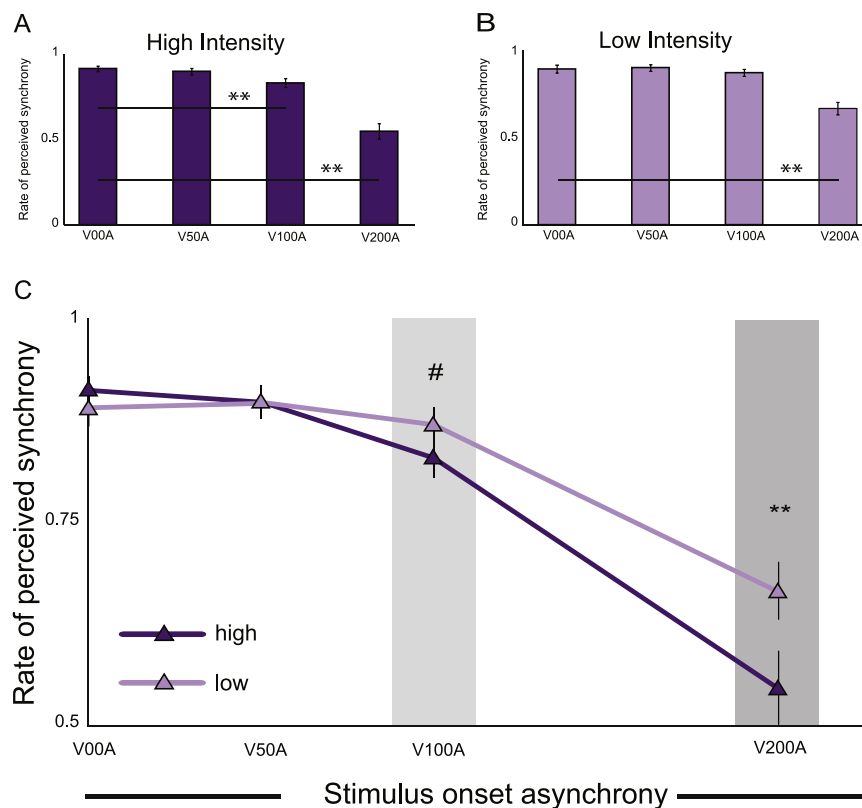


Fig. 2. A and B – Bar graphs present the changes across SOA within each level of stimulus intensity. For both saliencies, perceived synchrony decreases with increasing SOA with faster changes occurring in the high-saliency condition. C – Line graphs present the changes in perception of synchrony across stimulus intensity levels. Most notably, subjects perceived audiovisual stimuli more often as synchronous for the largest SOA at the low saliency condition. For all panels, ** indicates $p < 0.0001$, # indicates $p < 0.06$.

that individuals were more likely to report asynchronous presentations as synchronous at lower levels of stimulus intensity.

3.3. Effects of stimulus intensity and temporal factors on response times

In addition to our analysis of the synchrony judgments of participants as a follow up, a supplementary analysis was undertaken to confirm the accuracy-based findings. Mean RTs were

determined by averaging across subjects by SOA and intensity level (Table 2). Repeated measures ANOVA showed a main effect of SOA ($F_{(3,43)}=2.630$, $p=0.049$, $\text{partial-}\eta^2=0.039$). In contrast, no significant differences in mean RTs across intensities ($F_{(1,43)}=1.260$, $p=0.268$, $\text{partial-}\eta^2=0.028$) and no interaction effect between intensity and SOA ($F_{(3,43)}=0.686$, $p=0.566$, $\text{partial-}\eta^2=0.019$) were found.

In addition to an analysis of mean RTs, a cumulative distribution function (CDF) analysis was also performed. CDFs were

Table 1
Rates of perceived synchrony relative to synchronous presentations.

SOA (ms)	High intensity		Low intensity	
	t-score	p-value	t-score	p-value
50	1.39	n.s.	0.56	n.s.
100	4.05	1.93e^{-4}	1.10	n.s.
200	7.60	5.61e^{-7}	5.74	5.61e^{-7}

SOA – stimulus onset asynchrony.

Table 2
Mean response times in ms.

SOA (ms)	High intensity		Low intensity	
	Mean	St. err.	Mean	St. err.
0	421.5	29.7	429.1	27.9
50	420.1	26.1	401.6	26.0
100	433.2	29.6	434.5	25.8
200	457.6	28.0	427.8	23.9

SOA – stimulus onset asynchrony.
St. err – standard error.

calculated for each subject and then averaged across participants for each condition. To explore the impact of time on these CDFs, differences between the CDFs at 0 ms SOA and 200 ms SOA for each individual were computed for both intensity levels (difference curves plotted in Fig. 3a and b) and binned into 100 ms intervals for comparison. In the high intensity condition, there was a significant effect of SOA in the RT bins spanning from 200–1000 ms, where the 0 ms SOA was associated with significantly faster RTs than the 200 ms SOA. A similar pattern was seen in the low-intensity conditions, where RTs in the bins between 200–400

and 500–600 ms showed marginally faster RTs at the 0 ms SOA. Regardless of intensity level, the greatest difference invariably occurred during the early portion of the response distribution, as seen in Fig. 3c. Here the difference in these response distributions were segmented into 100 ms bins for comparison, revealing an interaction effect where SOA showed a stronger impact on RT when stimuli were presented at high intensity levels relative to low intensity levels. Finally, to assess this interaction overall, an area-under-the-curve measure was taken for each individual's differences in response distributions across SOAs (Fig. 3d). A direct comparison of these areas under the curve confirmed the results of the binned analysis, showing that SOA had a greater impact on stimuli presented at a high intensity level ($t=28.32$, $p=1.29\text{e}^{-14}$, $d=8.539$).

4. Discussion

Previous studies have established that manipulating space, time, and stimulus intensity can greatly alter multisensory integration and the associated neural, behavioral and perceptual responses. The present study extends this work by examining how stimulus timing and intensity interact to impact perceptual and behavioral performance, showing a strong interdependency between timing and intensity in judgments of the synchrony/asynchrony of the paired stimuli (i.e., synchrony). Specifically, participants were significantly more likely to report asynchronous audiovisual stimuli as synchronous when stimulus intensity levels were lower. This novel finding illustrates that stimulus intensity plays an important and previously unreported role in the perception of the timing of multisensory events.

The results suggest that processing of multisensory stimuli that are weakly effective is less temporally precise than for more

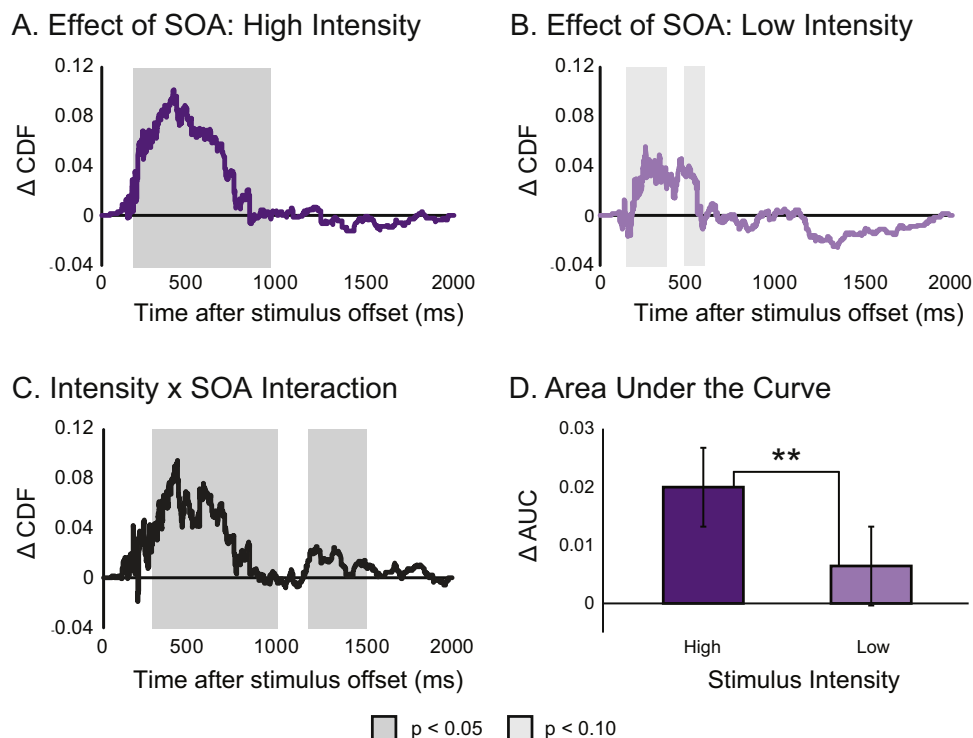


Fig. 3. A and B – Differences in cumulative distribution functions measured with synchronous (0 ms SOA) and asynchronous (200 ms SOA) presentations with both high-intensity (Panel A) and low-intensity (Panel B) presentations. C – An interaction between SOA and stimulus intensity was observed using a difference-of-difference measure. The effect of SOA was significantly more pronounced with high- than with low-intensity stimuli. D – Area-under-the-curve measures were extracted from the CDF differences displayed in Panels A and B, providing a secondary measure of interactions between SOA and stimulus intensity. **denotes a significant difference at an α -level of 0.05.

effective stimuli. Multisensory systems thus appear to compensate for increasing levels of stimulus ambiguity by increasing their tolerance for asynchronies, resulting in perceptual binding over a greater degree of temporal disparity. Adjusting the width of the temporal binding window would compensate in a manner that effects the overall magnitude of multisensory integration, a notion supported by the time-window-of-integration model (Colonius and Diederich, 2004). Ecologically, this result makes a good deal of sense, since in natural environments sensory inputs arriving from a proximal source are likely to be more intense and will arrive at their respective sensory organs in a more temporally congruent manner. In contrast, sensory information from an identical event that occurs at a greater distance from the individual will be of lower intensity and the temporal disparity at the point of the respective sensory organs will be increased. Thus, a greater tolerance for temporal offsets with low-intensity sensory signals is necessary in order to properly reflect the natural statistics of the environment. Furthermore, the nervous system may also expand its temporal filter for less effective stimuli in order to compensate for the necessity to acquire more information toward a behavioral judgment. These results parallel previous work and work in this issue showing that more peripherally presented audiovisual stimuli are more likely to be perceptually bound at wider temporal offsets (Nidiffer et al. (2015) (in this issue); Stevenson et al., 2012c).

One hallmark feature in processing information from external and internal events is the brain's capability to continuously recalibrate and update ongoing neural processes in a dynamic fashion. This occurs particularly frequently when signals are noisy or less reliable as could be the case with low-intensity stimuli. Indeed, previous research has shown that the manner in which the nervous system integrates sensory information is adaptable according to the reliability of information in each sensory modality (Andersen et al., 2004; Beauchamp et al., 2010; Beierholm et al., 2009; Besson et al., 2010; Charbonneau et al., 2013; Clemens et al., 2011; Deneve and Pouget, 2004; Ernst and Banks, 2002; Fetsch et al., 2010, 2012, 2009; Helbig et al., 2012; Ma and Pouget, 2008; Nath and Beauchamp, 2011; Noppeney et al., 2010; Rohe and Noppeney, 2015).

Furthermore, multisensory systems are characterized by a great deal of plastic capacity, a feature that has been explored extensively in the temporal realm. Thus, through recalibration (Fujisaki et al., 2004; Vroomen et al., 2004) and perceptual learning (Powers et al., 2012, 2009; Schlesinger et al., 2014; Stevenson et al., 2013b), it has been demonstrated that an individual's perception of synchrony and the window of time within which that individual will perceptually bind paired stimuli can be modified. Given the malleability of multisensory processing, it seems plausible then that these processes may become more liberal in the window of time with which they perceptually bind incoming sensory information based upon its reliability or intensity. Indeed, such real time recalibration of sensory integration has been reported previously in reference to stimulus reliability as determined by signal-to-noise ratio (Nath and Beauchamp, 2011). In this study, reweighting of stimulus inputs based on reliability impacted the magnitude of multisensory integration in STS as demonstrated through fMRI. While the current data set cannot directly assess this hypothesis, this possible explanation is intriguing and warrants direct testing in future research.

Consistent with the results of the current study, one line of previous research has provided evidence that another aspect of multisensory temporal processing, the point of subjective simultaneity (PSS), can also be modulated by stimulus intensity. The PSS, which describes the temporal offset at which the perception of synchrony is maximal, is generally found when the visual component of a multisensory stimulus slightly precedes the

auditory component. Studies investigating the role that stimulus intensity has on the PSS have generally shown that with decreases in stimulus intensity, the PSS shifts to an SOA with a greater auditory lag (Boenke et al., 2009; Neumann et al., 1992). Furthermore, stimuli presented in the periphery have also been shown to have a PSS associated with a greater visual-first SOA (Arden and Weale, 1954; Zampini et al., 2003), a result that may be derivative of changes in stimulus effectiveness. Furthermore, prior work from our lab has shown that the window within which an individual perceives temporal synchrony is in fact broader in the periphery (Stevenson et al., 2012c). This extension beyond measures of the PSS is vital, as manipulations of other stimulus factors, such as spatial disparity, have been shown to impact the window of integration without impacting the PSS (Keetels and Vroomen, 2005). The current study, however, provides the first evidence that direct manipulations of stimulus intensity alter the temporal tolerance for perceived simultaneity.

In accord with the changes in rates of perceived synchrony across levels of stimulus intensity, exploratory analysis of changes in response time CDFs across SOAs and intensity levels were also observed. Individuals were slower to respond to stimuli presented at longer SOAs, but this effect was smaller with low-intensity stimuli. This interaction effect in RTs provides converging evidence with the simultaneity judgment data, and reinforces the conclusion that at lower intensity levels (i.e., weaker effectiveness) multisensory systems are more tolerant of temporal offsets and thus capable of binding audiovisual stimuli over larger temporal intervals.

The role of effectiveness and timing is also important in the study of atypical sensory and multisensory perception. Specifically, effectiveness and timing play a role in dysfunctional multisensory integration in a number of clinical populations. For example, the ability to benefit from seeing a speaker's face while conversing in a noisy environment is dependent upon the effectiveness of the auditory signal, but individuals with schizophrenia (Ross et al., 2007b) and autism (Foxe et al., 2013) show deficits in the amount of perceptual benefit they gain at low levels of stimulus effectiveness. Furthermore, dysfunction in temporal processing of multisensory stimuli have also been shown in individuals with autism (Baum et al., 2015; Bebko et al., 2006; de Boer-Schellekens et al., 2013; Foss-Feig et al., 2010; Kwakye et al., 2011; Stevenson et al., 2014d, 2014e, 2014a, 2014b; Wallace and Stevenson, 2014; Woynarowski et al., 2013), schizophrenia (Martin et al., 2013), and dyslexia (Froyen et al., 2011; Hairston et al., 2005; Virsu et al., 2003). Since having an appropriately sized temporal binding window ensures proper multisensory integration, one avenue of future research should investigate whether or not clinical populations with enlarged temporal windows will show a corresponding widening of these windows with changing stimulus properties (i.e., stimulus intensities as described in the current study) or with increasing levels of stimulus complexity (Stevenson and Wallace, 2013; Vroomen and Stekelenburg, 2011). Furthermore, given previous demonstrations of how one can improve multisensory temporal precision through perceptual learning (Powers et al., 2012, 2009; Schlesinger et al., 2014; Stevenson et al., 2013b), future studies applying such training to clinical populations as possible therapeutic tool may prove fruitful. Emerging evidence suggests enormous developmental plasticity in multisensory temporal function (Hillock et al., 2011; Hillock-Dunn and Wallace, 2012; Stevenson et al., 2013b). Such developmental malleability could potentially be directed in clinical populations to ameliorate some of the changes in sensory function. Indeed, audiovisual training in children with reading disabilities has been used as a tool to improve reading comprehension (Kujala et al., 2001; Veuillet et al., 2007). Thus, delineating the temporal factors and constraints for multisensory binding and integration in

normative populations builds the foundation for comparisons in clinical populations, which may provide key insights into the design of effective interventional measures.

5. Conclusions

The study presented here provides novel evidence of a relationship between the perception of multisensory stimuli in relation to their stimulus effectiveness and their temporal synchrony. Specifically, these data suggest that the effectiveness of a stimulus presentation impacts how the temporal dynamics of the stimulus presentation are perceived, where tolerance for stimulus asynchronies is increased as stimulus effectiveness decreases. While the present study clearly indicates stimulus-intensity dependent changes in the window of temporal integration, it is limited to only two saliency levels, suggesting further exploration of this effect with a broader range of SOAs and salience levels that span the dynamic range of behavioral responses to provide a full picture of how the temporal principle and inverse effectiveness interact.

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