

Oxford Brookes University – Research Archive and Digital Asset Repository (RADAR)

Pilling, M

Auditory event-related potentials (ERPs) in audiovisual speech perception.

Pilling, M (2009) Auditory event-related potentials (ERPs) in audiovisual speech perception. *Journal of speech, language, and hearing research*, 52 (4). pp. 1073-1081.

Doi: 10.1044/1092-4388(2009/07-0276)

This version is available: http://radar.brookes.ac.uk/radar/items/98c53250-30e0-afc8-f0eb-3f39c08815af/1/

Available in the RADAR: February 2012

Copyright © and Moral Rights are retained by the author(s) and/ or other copyright owners. A copy can be downloaded for personal non-commercial research or study, without prior permission or charge. This item cannot be reproduced or quoted extensively from without first obtaining permission in writing from the copyright holder(s). The content must not be changed in any way or sold commercially in any format or medium without the formal permission of the copyright holders.

This document is the postprint version of the journal article. Some differences between the published version and this version may remain and you are advised to consult the published version if you wish to cite from it.



Directorate of Learning Resources

Auditory Event Related Potentials (ERPs) in audiovisual speech perception

Michael Pilling

MRC Institute of Hearing Research, Science Road, Nottingham, NG7 2RB. United Kingdom Email: mpilling@brookes.ac.uk

Published in Journal of Speech, Language & Hearing Research, (2009), 52, 1073-1081.

Abstract

Purpose: It has recently been reported (e.g. Wassenhove et al., 2005) that audiovisual (AV) presented speech is associated with an N1/P2 auditory ERP response that is lower in peak amplitude compared to the responses associated with auditory only (AO) speech. This effect is replicated. Further comparisons are made between ERP responses to AV speech in which the visual and auditory components were in or out of synchrony, to test whether the effect is associated with the operation of integration mechanisms, as has been claimed, or occurs because of other factors such as attention.

Method: ERPs were recorded from participants presented with recordings of unimodal or AV speech syllables in a detection task.

Results: Comparisons were made between AO and AV speech, and between synchronous and asynchronous AV speech. Synchronous AV speech produced an N1/P2 with lower peak amplitudes than with AO speech, unaccounted for by linear superposition of visually-evoked responses onto auditory-evoked responses. Asynchronous AV speech produced no amplitude reduction. **Conclusion**: The dependence of N1/P2 amplitude reduction on AV synchrony is consistent with the effect resulting from the operation of integration mechanisms and validates it as an electrophysiological marker of AV integration.

Keywords: speech, ERP, audiovisual integration

Apart from by ear, speech information can also be obtained by the eye, from the visibly perceptible movements of the face that accompany speech production. When such visual speech as well as auditory speech information is present, the brain tends to *integrate* the two sources into a single unified percept (see Summerfield, 1987). The unity of visual and auditory speech is revealed in a number of behavioral markers of integration. Possibly the most notorious of these is the McGurk illusion (McGurk & MacDonald, 1975). In the McGurk illusion, an auditory CV syllable (e.g. /ba/) is presented concurrently with phonetically incongruous visual speech (e.g. a face articulating /ga/); observers typically report hearing a fusion of the two signals (e.g. /da/), illustrating the unity of auditory and visual speech perception. Other markers of the audiovisual integration of speech include the audiovisual (AV) speech-detection advantage (Bernstein et al. 2004); auditory speech-in-noise is easier to detect when presentation is AV than when auditory only (AO). AV presented speech is also usually found to be more intelligible than AO speech; such effects are particularly apparent with the auditory speech presented in noise (Erber, 1975), this advantage for AV speech intelligibility is greater than would be expected on the basis of the information from the auditory and visual modalities being processed independently (see Grant & Walden, 1996).

What these above effects demonstrate is that our perception of auditory speech operates from a representation that is an integration of information derived from both the auditory and visual modalities. However there is some debate about the processing stage at which this integration process occurs (see Schwartz et al., 1998). Some models, so-called early integration models, place the integration of speech fairly early in the processing stream before phonetic categorization has occurred (e.g. Braida's 'pre-labeling model'; Braida, 1991). Other models have placed AV integration at a relatively late stage, after the information in each modality has been independently evaluated (e.g. Massaro, 1987, 1998).

Some behavioral evidence has been found to support early-integration models of AV speech perception. For instance, visual speech has been shown to influence the perception of voicing in an auditory speech stimulus (Green, 1998), consistent with visual speech interacting with auditory speech prior to phonetic categorization. Recently more direct evidence for early interactions in AV speech has been suggested by studies looking at auditory event-releted potentials (AERPs) Two studies (Besle et al., 2004; Wassenhove et al., 2005) found that the N1/P2 complex of the AERP associated with AV speech presentation was lower in amplitude compared against the responses evoked by AO speech presentation. Such a finding is important because it suggests that visual speech information may be interacting with auditory speech processing at an early (pre-categorical) processing stage. The N1/P2 wave is associated with the processing of the physical attributes of an auditory stimulus *prior* to its categorization (Näätänen & Winkler, 1999). It occurs within a time window of 100-200 ms after the onset of an auditory stimulus and is typically maximal around the scalp vertex. It is assumed to have multiple neural generators located in the dorsal surface and superior temporal plane of both temporal lobes (Scherg & Von Cramon, 1986; Näätänen & Picton, 1987).

Importantly Besle et al. (2004) and Wassenhove et al. (2005) found that lower amplitudes associated with AV presentation were not simply a result of the linear superposition of visual and auditory activity: N1/P2 amplitudes associated with AV speech were lower even when compared against the aggregate of the unimodal responses for auditory and visual speech. This demonstrated that it was the underlying AERP waveform itself that was being modulated by the AV presentation

of the speech. The effect of AV speech presentation on AERP amplitudes is consistent with visual speech producing a 'deactivation' (Wassenhove et al.) or 'depression' (Besle et al.) in the auditory cortex. The finding draws some parallels with ERP studies of binaural auditory integration. Similar amplitude reductions have been shown in the N1/P2 (along with earlier latency components) response to sounds presented binaurally compared against aggregate monaural ERP responses (e.g. McPherson & Starr, 1993), suggesting that reduced amplitudes may be a general aspect of integration processes in the auditory modality.

AV speech thus seems to be reliably associated with N1/P2 amplitude reductions compared with AO speech. However further validation is needed before the effect can be accepted as an electrophysiological correlate of an AV integration process as opposed to the result of some other factor.

An alternative explanation of the effect is that it reflects general top-down inhibition of the auditory cortex occurring in the AV condition but not in the AO condition because of the two conditions' differing task demands. Shulman et al (1997) found that auditory cortex activity was significantly reduced (as measured by Positron Emission Tomography) when participants where actively engaged with a visual stimulus compared to when passively viewing the same visual stimulus. It is possible that the presence of the talking face in the AV condition induces such top-down inhibition in the auditory cortex as a consequence of participants actively processing the visual stimulus, It is further possible that this top-down inhibition is absent with a static fixation cross or static face because, in this case, active processing does not occur. If such top-down inhibition was being induced it might well result in reduced AERP amplitudes. However, the conditions under which Shulman et al. found these modulations are quite different to those in which AV speech is presented. In Shulman et al.'s task the auditory modality was task-irrelevant: the only auditory stimulation came from ambient background noise rather than from a relevant auditory stimulus as in AV speech.

A more plausible alternative explanation of the amplitude reduction effect in AV speech is that it is related to shifts in attention across modalities between the AO and AV conditions. It is possible that with AV stimulus presentation attention is shifted further towards the visual modality and away from the auditory modality, leading to lower responses in the auditory cortex. Indeed, the N1/P2 response is known to be modulated by attentional shifts: reduced attention to the auditory modality has been shown to result in smaller amplitudes in this response (Hanson & Hillyard, 1980). Wassenhove et al. (2005) rejected such an explanation because with AV speech instructing participants to attend just to the visual modality did not result in any additional amplitude reduction in the auditory responses. However, this may just mean that on top of the 'automatic' shifts to the visual information in AV speech presentation, further attentional shifts towards the visual modality have no further effect on AERPs. It does not necessarily rule out attention being responsible. Besle et al. (2004) also rejected an attentional explanation of the amplitude reduction effect. They argued that in their study the attentional demands were balanced across the AO and AV conditions because in both conditions the task they gave (a speeded auditory detection task) only required attention to the auditory modality. However, this may still not fully balance attention across the AO and AV conditions. Participants may still have been actively attending to the onset of the facial movements in the AV condition which provides a salient and task-relevant alerting cue for the onset of the auditory speech signal. This would mean that attention was still different to the AO condition because in the AO condition no such visual cue was present.

Thus further evidence is needed to test if the amplitude reduction in the N1/P2 is in fact associated with integration mechanisms, rather than some other process. The experiments presented here try to provide such evidence by observing the effect of synchrony between the visual and auditory streams with AV speech on the N1/P2 amplitude reduction effect. It has been demonstrated that AV speech integration mechanisms operate only within a particular temporal window, outside of which the effects of visual speech on auditory speech perception are largely absent. This temporal window is usually found to be asymmetric: A discrepancy of ~100 ms from the recorded position of auditory speech is usually perceptible when auditory speech *leads* from its recorded position with visual speech, while a discrepancy greater than ~250 ms is needed when auditory speech *lags* from its recorded position with visual speech (Dixon & Spitz, 1980). Outside this temporal window the earlier described behavioral effects of integration tend to be either absent or greatly weakened, suggesting that the auditory and visual speech signals are no longer producing a unified percept (e.g. McGurk effect: Munhall et al., 1996; AV speech- in-noise advantage: Grant & Greenberg, 2003).

If the amplitude reduction effect is a consequence of integration mechanisms then the effect should be significantly weakened, or even abolished, for AV speech in which the auditory component is outside the temporal window of perceived synchrony with the visual signal. If, however, the amplitude reduction effect is a consequence of a process other than integration (e.g. attention) then manipulations of asynchrony should have little consequences for the AERPs: N1/P2 amplitude reduction should remain even with perceptibly asynchronous AV speech because any attention diverted towards the visual component of speech should be equal in the synchronous and asynchronous AV conditions. The experiments reported in this paper test this possibility.

A task was given in which participants detected infrequent phoneme syllables (targets) presented in a stream of standard syllables. Experiment A compared recorded ERPs to these standard syllables in unimodal auditory (AO), unimodal visual (VO) and synchronous AV conditions (AV). Experiment B recorded ERP responses to these syllables in synchronous AV (AV), asynchronous AV (AVasynch) and AO conditions.

Method

Twelve participants were used in Experiments A and twelve different participants in Experiment B, (age range 18 to 30). All had normal hearing (assessed by pure-tone audiogram),

normal or corrected-to-normal visual acuity (assessed by Snellen chart). Stimuli consisted of high quality audiovisual recordings of a male talker articulating the CV syllables /pa/ and /ta/ from an initial neutral expression. Four examples of each syllable were used for the experiments. Recordings were normalized and calibrated to be approximately 60 dB SPL. Video recordings were edited to start from just before the initial visible articulation to the end of the articulation and converted into a sequence of still bitmap images (160×210 mm) presented on a 15" LCD screen. The screen was viewed at a distance of approx. 400 mm. The first frame of each recording was presented for 1000 ms as a still frame. After 1000 ms the first moving frame (the visible beginning of the articulation) and subsequent moving frames were presented at a rate of 25 frames-persecond. Corresponding digitized audio files were presented binaurally via Sennheiser headphones. In the AV condition the video frames of the talker were presented in synchrony with the auditory stimulus as they were naturally recorded. The duration between the onset of the first moving frame and the onset of the auditory speech was different for the four exemplars of /ta/ (range 210-296 ms) and /pa/ (range 239-401 ms) due to natural variations in the articulation of the talker. In the AVasynch condition the auditory stimulus was presented 200 ms ahead of its recorded position. This asynchrony was noted by all participants as being clearly perceptible. In this condition the duration between the first moving frame of the video and the onset of the auditory speech varied between a range of 10-96 ms for /ta/, and 39-201 for /pa/ for the different exemplars of the tokens. In the AO condition the moving frames of the talker were replaced with a static fixation cross present for the same duration. In the VO condition the moving frames of the talker were presented silently. A 34 electrode EEG cap arranged using the 10/20 system was used with the ground at AFz and Cz used for online referencing. EEG was recorded at an analogue-to-digital conversion rate of 1000 Hz (subsequently downsampled to 500 Hz to speed analysis). The target syllable (/pa/) was presented 40 times randomly in a sequence of 180 standards (/ta/) per block. Figure 1 shows a schematic diagram of a single AV standard trial. A 3500~4000 ms inter-trial interval was given in

which a blank screen was presented. Participants were instructed to listen to the speech while looking at the screen and to press a response key each time a target was presented. In Experiment A blocks were presented in the AO, VO and AV conditions. In Experiment B blocks were presented in AO, AV and AVasynch conditions. Two blocks of each condition were presented in a randomized order.

Figure 1 about here

Analysis and Results

Offline processing of EEGs was performed using Brain Vision Analyzer (V 1.05). Recordings were manually screened for artifacts and a Butterworth filter (1 - 30 Hz) applied. Data was then re-referenced using all 34 channels and channel Cz reconstructed. Ocular correction was applied using FP1 as the EOG channel using the procedure developed by Gratton et al. (1983). Response averaging was done only for the standards to avoid contamination of the AERPs from oddball and response related activity. The onset of the auditory stimulus marked the start of each epoch for averaging purposes (for the VO condition the epoch was marked at a point were the auditory stimulus would have onset had one been presented). The N1/P2 was evoked in every condition except VO and was largest around the central electrode cites (FC1, FC2, Cz, C3, C4, CP1, CP2).¹ Grand average responses on these electrode sites are shown in Figure 2 for Experiment A and Figure 3 for Experiment B.

Figures 2 and 3 about here

Peak-to-peak measures of the N1/P2 were computed for the 7 central electrodes for the averaged responses. The N1 peak was specified as the largest negative local maximum between 60-140 ms and the P2 peak as the largest positive local maximum between 130-300 ms after onset.

¹ No reliable P50 wave was observed in the grand average or in the individual waveforms. P50 responses are typically small in amplitude and more variable than later auditory components. Its absence in this study may be due to one of several factors, such as

The peak-to-peak was also calculated on the computed sum of the unimodal responses (AO+VO) to verify whether differences between the AO and AV conditions were genuine interactivity rather than linear superposition. Averaged peak-to-peak measures for the different conditions are given in Table 1 for Experiment A and Table 2 for Experiment B.

Tables 1 and 2 about here

Measures on the central electrodes were subjected to two-way ANOVAs with Condition (AO vs. AV; AO vs. AO+AV) and Electrode (seven levels) as repeated measures factors. Comparison between AO and AV showed that response amplitudes were significantly lower in the AV condition, F(1, 11) = 49.49, MSE = 1.77, p < .0001. Peak amplitudes were also significantly lower when AV speech was compared with the sum of the unimodal responses (AO+VO), F(1, 11) = 80.84, MSE = 1.39, p < .0001.² This shows that the reduction in N1/P2 peak amplitudes in the AV condition relative to the AO condition was not explained by linear superposition. This same amplitude reduction between the AV and AO conditions was replicated in Experiment B (F(1, 11) = 49.05, MSE = 0.99, p < .0001), demonstrating its reliability. In both Experiments the difference between the AO and AV conditions was found to be maximal at Cz. Critically, in the AVasynch condition amplitude reduction was not found for any electrode. In fact peak amplitudes were overall slightly larger than they were in the AO condition, though the effect was not significant (F(1, 11) = 2.88, MSE = 1.84, p > .05).

the large inter-trial intervals or due to attention being partly directed towards the visual modality (even the AO condition participants were instructed to look at the fixation cross present during each trial).

 $^{^2}$ There was also a small peak latency effect: Latencies of the N1 and P2 peaks occurred earlier in the AV condition compared against the AO condition on some of the central electrodes, an effect also reported by Wassenhove et al. (2005). However the latency effect found here was statistically small where it occurred and the effect not reliably significant across Experiments A and B for the two peaks with AV speech. As a result the paper confines the discussion to the more robust amplitude reduction effect

Discussion

When AV speech was presented the resulting N1/P2 wave was significantly smaller in its peak amplitude than when auditory speech was presented (Experiment A). Furthermore the responses to AV speech were also significantly lower in amplitude when compared against the sum of unimodal responses, demonstrating that the effect associated with AV speech was not just linear superposition. This result corroborates the findings of Besle et al. (2004) and Wassenhove et al. (2005) in demonstrating a nonlinear effect on the amplitudes of AERPs. Furthermore, this amplitude reduction effect was dependent on AV synchrony (Experiment B). The responses to asynchronous AV speech exhibited no such effect. Thus, like the behavioral markers of AV integration, such as the McGurk illusion (Munhall et al. 1996), the electrophysiological amplitude reduction effect is also sensitive to synchrony between the auditory and visual modalities.

Therefore, these results are consistent with the notion that the amplitude reduction effect is associated with the operation of integration mechanisms. The results therefore provide further support for early integration models of AV speech in showing that at least some aspects of the integration of auditory and visual information occur at an early, pre-categorical stage.

The results are inconsistent with a top-down inhibition (Shulman et al., 1997) account of the amplitude reduction effect, in which the effect occurs as a consequence of active processing of the visual stimulus. The moving face was present in both the synchronous and asynchronous AV conditions so if active processing of the visual stimulus is the relevant factor then this should have occurred in both the synchronous and asynchronous AV conditions. However, the amplitude reduction effect occurred only in the synchronous condition. The results are also inconsistent with an attentional account in which attention is partly diverted away from the auditory modality by the onset of the moving face in AV conditions. Although the auditory speech was shifted from its natural recorded position, this shifted position was always later than the first moving frame of the visual stimulus, meaning that the visual stimulus still provided an effective alerting cue to the onset of the auditory speech even in the asynchronous AV condition. Nevertheless, no amplitude reduction was observed in this condition relative to the AO condition.

What do these results tell us about the neural mechanisms of integration? Besle et al. (2004), on the basis of source analysis, suggested that the modulation of AERPs occurs at least partly in the supratemporal auditory cortex. They further suggested that the superior temporal sulcus (STS), a multisensory region receiving feedforward input from both the auditory and visual corteces, and a site that several imaging studies have implicated as a likely site of AV speech integration (e.g. Sekiyama et al., 2003), could be the source of the inhibitory effects in the auditory cortex via inhibitory reentrant pathways. The results presented in this paper, together with findings from a study by Macaluso et al. (2004) are consistent with this interpretation. Macaluso et al. found that the STS exhibits a greater haemodynamic response to synchronous than perceptibly asynchronous AV speech. It is possible that this greater activity in the STS with synchronous AV speech is reflected in greater inhibition of the auditory cortical regions, resulting in a smaller N1/P2; with asynchronous AV speech the lower STS activity may mean that this structure no longer inhibits the auditory cortex. More direct support for this physiological model of the integration effect could be made if recordings from haemodynamic and ERP imaging techniques were combined in the same experiment, allowing direct comparison between activity in the STS region and N1/P2 modulation, and finer temporal synchrony-asynchrony manipulations of AV speech were made. If STS activity and ERP amplitudes were found to have similar tolerances of asynchrony it would provide further support of the involvement of this structure in producing the inhibitory effect in the auditory cortex.

Although this study demonstrates the importance of the temporal synchrony of the visual stimulus with auditory speech, it does not tell us what the critical features of the visual stimulus are. Two possible candidates are the pictorial content of the facial features and the dynamic

characteristics of the visual signal (Vitkovitch & Barber, 1994; 1996; Rosenblum, & Saldaña, 1996). If the dynamic characteristics of the face are the more important factor, video frame rate will be a more important factor in the modulation of AERPs than the pictorial information within the frames. Further research is needed to determine the critical aspects of the visual stimulus in AV speech presentation that lead to modulation of AERPs.

A related and unresolved question concerns the aspects of AV integration processes that are being reflected in the modulations of the AERPs? Wassenhove et al. (2005) argued that the amplitude reduction reflects an aspect of processing relating to the merging of syllabic information between the visual and auditory speech signals in the auditory cortex by a depression mechanism. They argued that when AV speech is presented information about place-of-articulation can be obtained from the visual signal, making redundant the processing of certain speech formant frequencies in the auditory signal. Similarly, Besle et al. (2004) proposed that the amplitude reduction reflects the facilitation of syllable feature analysis in the auditory cortex in the form of suppressive modulation. However, another possibility is that the modulations are reflecting the more basic structural spatiotemporal correspondence of the auditory and visual events (see Spence, 2007). Indeed consistent with this, Stekelenburg & Vroomen (2007) have recently shown that similar amplitude reductions in AERPs can be found in the perception of ecologically-valid nonspeech AV events such as hand-claps.

In conclusion the sensitivity of the reduction in N1/P2 amplitude associated with AV speech is a genuine electrophysiological marker of AV integration mechanisms. Further work will determine the precise aspects of the visual stimulus that are needed for N1/P2 amplitude reduction to occur.

References

- Bernstein, L. E., Takayanagi, S., & Auer, E. T., Jr. (2004). Auditory speech detection in noise enhanced by lipreading. Speech Communication, 44, 5-18.
- Besle, J., Fort, A., Delpuech, C., & Giard, M. H. (2004). Bimodal speech: early suppressive visual effects in the human auditory cortex. European Journal of Neuroscience, 20, 2225–2234.
- Braida, L. D. (1991). Crossmodal integration in the identification of consonant segments. <u>Quarterly</u> <u>Journal of Experimental Psychology: A, 43, 647–677.</u>
- Dixon, N. F., & Spitz, L. (1980). The detection of auditory visual desynchrony. <u>Perception, 9</u>, 719-721.
- Erber, N. P. (1975). Auditory-visual perception of speech. Journal of Speech and Hearing Disorders, 40, 481-492.
- Giarda, M. H., & Peronnet, F. (1999). Auditory-visual integration during multimodal object recognition in humans: A behavioral and electrophysiological Study. <u>Journal of Cognitive</u> <u>Neuroscience, 11</u>, 473-490.
- Grant, K. W., & Walden, P. F. (1996). Evaluating the articulation index for auditory-visual consonant recognition. Journal of the Acoustical Society of America, 100, 2415-2424.
- Grant, K. W., and Greenberg, S. (2001). Speech intelligibility derived from asynchronous processing of auditory-visual information. <u>AVSP-2001 Proceedings</u>, Scheelsminde, Denmark.
- Gratton, G., Coles, M. G. H., & Donchin, E. (1983). A new method for off-line removal of ocular artifacts. <u>Electroencephalography and Clinical Neurophysiology</u>, 55, 468-484.
- Green, K. P. (1998). The use of auditory and visual information during phonetic processing: inplications for theories of speech perception. In R. Campbell, B. Dodd, & D. Burnham,

(Eds.), <u>Hearing by Eye II: Advances in the psychology of speechreading and auditory-visual</u> <u>speech</u>. (pp. 85-108). Hove, England. Psychology Press.

- Hansen, J., & Hillyard, S. J. (1980) Endogenous brain potentials associated with selective auditory attention, <u>Electroencephalography and Clinical Neurophysiology</u>, 49, 277–290.
- Macaluso, E., Dolan, R. C., Spence, D., & Driver J. (2004). Spatial and temporal factors during processing of audiovisual speech: a PET study. <u>Neuroimage, 21</u>, 725-732.
- Massaro, D. M. (1987). <u>Speech perception by ear and eye: A paradigm for psychological inquiry</u>. Hillside, NJ. Lawrence Erlbaum Associates.
- Massaro, D. M. (1998). <u>Perceiving talking faces: From speech perception to behavioral principle</u>. Cambridge, MA. MIT Press.
- McGurk, H., & MacDonald, J. (1976). Hearing lips and seeing voices. Nature, 264, 746-48.
- McPherson, D. L., & Starr, A. (1989). Binaural interaction in auditory evoked potentials: brainstem, middle- and long-latency components. <u>Hearing Research, 66</u>, 91-98.
- Munhall, K.G., Gribble, P., Sacco, L., & Ward, M. (1996). Temporal constraints on the McGurk effect. <u>Perception and Psychophysics, 58,</u> 351-362.
- Näätänen R., & Winkler, I. (1999). The concept of auditory stimulus representation in cognitive neuroscience. <u>Psychological Bulletin, 6</u>, 826–859.
- Näätänen, R., & Picton, T. (1987). The N1 wave of the human electric and magnetic response to sound: a review and an analysis of the component structure. <u>Psychophysiology</u>, 24, 375-426.
- Rosenblum, L. D., & Saldaña, H. M. (1996). An audiovisual test of kinematic primitives for visual speech perception. <u>Journal of Experimental Psychology: Human Perception and</u> <u>Performance. 22</u>, 318-331.
- Scherg, M., & Von Cramon, D. (1986) Evoked dipole source potentials of the human auditory cortex. <u>Electroencephalography and Clinical Neurology, 65</u>, 344–360.

- Schwartz, J.L., Robert-Ribes, J., and Escudier, P. (1998). Ten years after Summerfield: a taxonomy of models for audio-visual fusion in speech perception. In R. Campbell, B. Dodd, & D. Burnham, (Eds.), <u>Hearing by Eye II: Advances in the psychology of speechreading and auditory-visual speech</u>. (pp. 85-108). Hove, England. Psychology Press.
- Sekiyama, K., Kannoc, I., Miurac, S., & Sugit, Y. (2003). Auditory-visual speech perception examined by fMRI and PET. <u>Neuroscience Research</u>, 47, 277-287.
- Shulman, G. L., Corbetta, M., Buckner, R. L., Raichle, M. E., Fiezi, J. A., Miezini, F. M., & Petersen, S. E. (1997). Top-down modulation of early sensory cortex. <u>Cerebral Cortex. 7</u>, 193-206.
- Spence, C. (2007). Audiovisual multisensory integration. <u>Acoustic Science and Technology, 28</u>, 61-70.
- Stekelenburg, J. J., & Vroomen, J. (2007). Neural correlates of multisensory integration of ecologically valid audiovisual events. Journal of Cognitive Neuroscience, 19, 1964-1973.
- Summerfield, Q. (1987). Some preliminaries to a comprehensive account of audio-visual speech perception. In B. Dodd, & R. Campbell (Eds.), Hearing by Eye: <u>The Psychology of</u> <u>Lipreading</u>. (pp. 3-51). London, England. Lawrence Erlbaum Associates.
- van Wassenhove, V., Grant, K. W., & Poeppel, D. (2005) Visual speech speeds up the neural processing of auditory speech. <u>Proceedings of the National Academy of Science, 102</u>, 1181-1186.
- Vitkovitch, M., & Barber, P. J. (1994). Effect of video frame rate on subjects' ability to shadow one of two competing verbal passages. <u>Journal of Speech, Language and Hearing Research</u>, 37, 1204-1210.
- Vitkovitch, M., & Barber, P. J. (1996). Visible speech as a function of image quality: Effects of display parameters on lip reading ability. <u>Applied Cognitive Psychology</u>, 39, 121-140.

Tables

Table 1. Mean N1/P2 peak-to-peak amplitudes (in μ V) across participants for AO, AV and AO+VO in Experiment A. Standard errors are given in brackets.

	FC1	FC2	Cz	CP1	CP2	C3	C4
AO	6.36 (0.50)	6.44 (0.57)	7.90 (0.67)	5.10 (0.43)	4.63 (0.49)	4.84 (0.37)	4.83 (0.48)
AV	4.74 (0.42)	4.74 (0.44)	5.65 (0.53)	3.79 (0.47)	3.63 (0.48)	3.80 (0.34)	3.65 (0.37)
AO+VO	6.68 (0.42)	6.72 (0.46)	7.84 (0.63)	5.04 (0.43)	4.67 (0.44)	5.20 (0.47)	5.29 (0.46)

Table 2. Mean N1/P2 peak-to-peak amplitudes (in μ V) across participants for AO, AV and AVasynch in Experiment B. Standard errors are given in brackets.

	FC1	FC2	Cz	CP1	CP2	C3	C4
AO	5.06 (0.39)	4.97 (0.30)	5.83 (0.40)	3.78 (0.31)	3.40 (0.22)	3.88 (0.31)	3.43 (0.19)
AV	3.62 (0.30)	3.55 (0.27)	4.18 (0.36)	2.95 (0.27)	2.76 (0.27)	2.95 (0.26)	2.81 (0.22)
AVasynch	5.36 (0.48)	5.12 (0.41)	6.30 (0.60)	4.24 (0.45)	3.74 (0.42)	4.35 (0.38)	3.72 (0.26)

Figure headings

- Figure 1. Schematic diagram of a standard AV trial. The first moving frame occurs after a 1000 ms still frame of the talkers face in a neutral position. An example frame of the talkers face at the apex of the articulation is shown. Note that the ERP trigger is set to the onset of the auditory speech stimulus which, for the standards, occurs between a range of 210 and 296 ms after the first moving frame depending on the particular exemplar of the recorded speech being presented (on AVasynch trials the onset of the auditory speech occurred between a range of 10 and 96 ms after the first moving frame). In AO trials the video frames are replaced by a static fixation cross.
- Figure 2. Grand average ERP responses in Experiment A for the seven central electrodes (FC1, FC2, Cz, C3, C4, CP1, CP2) in AO (black line), AV (darker grey line) and VO (lighter grey line) speech conditions. Response amplitudes are corrected by a -100 ms pre-stimulus baseline. Negative deflections are downwards on the ordinate. The N1/P2 wave (present in the AO and AV conditions has a negative peak at ~100 ms and a positive peak at ~200 ms. Note that the peak amplitude of the N1/P2 wave associated with the AV condition is lower to that associated with the AO condition. This effect is most apparent at Cz.
- Figure 3. Grand average ERP responses in Experiment B for the seven central electrodes in AO (black line), AV (darker grey line) and AVasynch (lighter grey line) conditions, corrected by a -100 ms pre-stimulus baseline. Notice that, as in Figure 2, the AV condition is associated with lower peak amplitude responses than the AO condition (particularly at Cz) in the N1/P2. Note that this is not the case for the AVasynch condition, where the associated peak amplitudes are slightly higher than with the AO condition.



Figure 1



