A Gap Prepulse with a Principal Stimulus Yields a Combined Auditory Late Response

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Background and Objectives: The gap prepulse inhibition of the acoustic startle response has been used to screen tinnitus in an animal model. Here, we examined changes in the auditory late response under various conditions of gap prepulse inhibition. Subjects and Methods: We recruited 19 healthy adults (5 males, 14 females) and their auditory late responses were recorded after various stimuli with or without gap prepulsing. The N1 and P2 responses were selected for analysis. The gap prepulse inhibition was estimated to determine the optimal auditory late response in the gap prepulse paradigm. Results: We found that the gap per se generated a response that was very similar to the response elicited by sound stimuli. This critically affected the gap associated with the maximal inhibition of the stimulus response. Among the various gap-stimulus intervals (GSIs) between the gap and principal stimulus, the GSI of 150 ms maximally inhibited the response. However, after zero padding was used to minimize artifacts after a P2 response to a gap stimulus, the differences among the GSIs disappeared. Conclusions: Overall, the data suggest that both the prepulse inhibition and the gap per se should be considered when using the gap prepulse paradigm to assess tinnitus in humans.

KEY WORDS: Auditory evoked responses · Prepulse inhibition · Tinnitus.

Introduction

An auditory evoked response (AER) is an electrical potential evoked by an external auditory stimulus. The AER is thought to provide useful neurophysiological information because of its sensitive temporal resolution. Among the various AER components, the N1 and P2 peaks of the auditory late response (ALR) represent auditory perception and provide information about the psychological state of an individual [1]. Experiments using a model of neurophysiological tinnitus indicated that these components affect tinnitus [1,2]. The N1 and P2 components differ significantly between tinnitus patients and healthy subjects [2].

Prepulse inhibition (PPI) is a general neurological phenomenon wherein the magnitude of a response to a principal stimulus is reduced when a certain stimulus precedes it by 30–500 ms [3]. PPI is indicative of sensorimotor gating and can be affected by various factors such as drugs, attention, and psychological issues [4,5]. Recent studies have demonstrated that prepulsing reduced the magnitudes of the cortical auditory evoked potentials (AEPs; P50, N1, P2, and P300) in healthy humans [5].

Advances in the methods of tinnitus assessment in animals have facilitated significant developments in tinnitus research. Turner, et al. [6] used a novel method termed gap PPI (gPPI) of the acoustic startle reflex to screen for tinnitus in an animal model. This new paradigm is based on the concept that a gap in the auditory stimulus serves as a prepulse and affects the acoustic startle response. By providing a gap before the main stimulus, in the presence of background noise similar to tinnitus...
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The startle response of an animal was reduced due to PPI. However, such inhibition was not evident in animals with tinnitus. Several studies have used this model to measure behavioral changes in various animals with noise-induced tinnitus [7,8].

Previous studies have investigated the relationship between the human startle reflex and PPI in the presence of various levels of background noise; a prepulse sound inhibited the startle response in humans [9,10]. In humans, an eye-blink startle response is typically measured using electromyography of the orbicularis oculi muscle [3]. Some studies have reported an inhibited response after a small pulse sound, especially in patients with insomnia [11] and traumatic stress disorder [12]. We speculate that the prepulse paradigm can also be used for evaluating humans; the startle response is replaced by the ALR when tinnitus is present.

Ku, et al. [13] used the gPPI paradigm with humans to evaluate if it could be used for the objective assessment of tinnitus. It was found that the auditory responses of the N1P2 complex were stable. The N1P2 ratios under gap and no-gap conditions differed marginally, depending on the gap duration, gap-stimulus interval (GSI), level of background noise, and response attained at the maximum noise level with the shortest gap duration (20 ms) and the lowest level of background noise. However, as the recording of AEPs with short repetition rates is uncommon because of overlapping neural responses [14], an ALR may include a response due to a preceding gap if there is only a short interval prior to the main stimulus. This possibility should be considered if a gap serves as an “auditory stimulus” that can itself evoke an auditory response.

We explored the possibility that a “gap” can serve as a stimulus for generating an auditory response at the cortical level. We also examined the feasibility of the gPPI paradigm for ALR measurement and investigated the differences in gPPI values derived using various GSIs between a gap prepulse and the principal pulse. Additionally, we investigated the effects of gap-only stimuli in the gap prepulse paradigm by correcting for the responses evoked by the gap prepulse stimuli. Our results demonstrate how the positioning of the gap prepulse affects the gPPI.

Subjects and Methods

Subjects

We enrolled 19 adults (5 males, 14 females) with a mean age of 28.68 years (range 21—50 years). None of the participants had any history of tinnitus or neurological disease. Their hearing was normal, as shown by pure tone audiometry at 500, 1,000, 2,000, and 4,000 Hz; all hearing thresholds were below 25 dB HL (hearing level) at all tested frequencies. This study was approved by the Institutional Board of Dankook University Hospital (DKUH 2013-08-009).

ALR measurement

The system used to measure ALR is described in Fig. 1. ALR potentials were measured using the wireless electroencephalogram (EEG) headset device (Epoc, EMOTIV Inc., San Francisco, CA, USA) and the computer-based custom data acquisition (DAQ) (NI PXI-6251, National Instruments, Austin, TX, USA) system (ALR system algorithm).
(Fig. 1). The system featured the DAQ software, EEG amplification and logging software, an EEG trigger, and the auditory response signal collection software.

**Stimulus and response processing**

The stimuli used are shown in Fig. 2. All stimuli included a continuous background noise with duration of 1,000 ms. The principal pulse (white noise) was 80 dB SPL (sound pressure level) and 20 ms in duration, delivered within a 500 ms stimulus train. To measure gap-only responses, stimuli of a 50 ms gap presented at the 400 ms time point within the background noise (8 kHz, 1/3 octave band) of 60 dB SPL was used (Fig. 2). Except when gap-only stimuli were delivered, both the gap and principal stimuli within the background noise were considered a single stimulus (Fig. 2). Gaps were placed at three different time points in the stimulus train: GSIs of 50, 150, and 250 ms. The stimuli were randomly presented at inter-stimulus intervals (ISIs) of 1–3 s to avoid adaptation and temporal prediction of the acoustic stimuli, using the ER-A3 (Etymotic Research, Elk Grove Village, IL, USA) insert earphone.

**ALR measurements and response processing**

All ALRs were measured in a double-walled soundproofed room. Participants were seated in comfortable chairs, and an inspector ensured that they did not sleep or engage in habitual movements. The common mode sense electrode was placed on the ipsilateral ear (reference) and the driven light leg electrode was placed on the contralateral earlobe (ground). Headgear with four channel electrodes (AF3, AF4, F3, and F4) was placed, and measurements commenced when the impedances of all electrodes were ≤5 kΩ. Stimuli were delivered (via the earphone) to the left ear, and the contralateral ear was blocked with a sponge earplug (KE-1100, Moden zone, Seoul, Korea). For each subject, three 5 min measurements at a 128 Hz sampling rate were made. The responses were processed using a software, and the responses to no-gap and with-gap stimuli were displayed as bold and dotted lines, respectively (Fig. 3).

EEG signals transferred to the computer were processed using specialized software. Both baseline drift and high-frequency noise were removed using a 1–30 Hz band-pass filter. The responses with or without gap prepulse were averaged. With- and without-gap responses are presented together; the ranges of interest including N1 (first negative peak) and P2 (second positive peak) are highlighted using yellow bars. The amplitudes of with- and without-gap responses were automatically calculated from the N1P2 amplitude, within the region of interest.
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of interest.

Data analysis

Epoch durations of 1,000 ms (which are sufficient to include all responses to the longest GSI) were analyzed. Four peaks developing within 0.4 to 0.8 ms were measured: N1, P1, N2, and P2 peaks. Of these, the N1 and P2 peaks after both no-gap and gap stimuli were automatically identified, and the gPPI was defined using the following equation:

\[
gPPI = \frac{N1P2_{no\,gap} - N1P2_{gap}}{N1P2_{no\,gap}}
\]

Zero padding was applied to the corrected data. The responses after the zero-crossing points of P2N2 waves after gap-only stimuli were set to zero. Thus, the ALR gap-only response was subtracted from the response after both a gap and a principal pulse. Mean and standard deviation (mean ± SD) of all peak amplitudes were compared.

Statistical analysis

All data were analyzed using the Statistical Package for the Social Sciences ver. 19 (IBM, Armonk, NY, USA). Analysis of variance (ANOVA) with Bonferroni post-hoc and paired t-test were used to determine the effect of GSI on the gPPI.

Results

Effects of gap prepulsing on N1P2 amplitudes at different GSIs

All responses were averaged, yielding the grand-averaged data. The grand-averaged ALR after an 80 dB SPL (8 kHz) stimulus is shown in Fig. 3. The N1P2 amplitude was 8.66 µV, and the N1 and P2 latencies were 0.6 and 0.69 ms, respectively. The N1P2 amplitude decreased when a silent gap (50 ms) was inserted prior to the principal stimulus (Fig. 3). In this case, the N1P2 amplitude was 4.96 µV. The N1P2 amplitudes (mean ± SD) at different GSIs are listed in Table 1. A GSI of 50 ms resulted in the greatest decrease in the N1P2 amplitude.

We calculated gPPI values to explore ALR changes at various GSIs. The gPPI values at GSIs of 50, 150, and 250 ms are shown in Fig. 4. The gPPI was maximal at the GSI of 50 ms and decreased as the GSI increased. We used the paired t-test to explore how the GSI affected the gPPI. Significant differences between the gPPI values at GSIs of 50 and 150 ms (p=0.012), and at GSIs of 50 and 250 ms (p=0.001), were apparent, but not between the gPPI values at GSIs of 150 and 250 ms (p=0.435).

Effect of the gap prepulse on N1P2 amplitude (after subtracting the gap-evoked auditory response)

The decrease in the N1P2 amplitude when a gap prepulse

| GSI (msec) | 50  | 15  | 250 |
|-----------|-----|-----|-----|
| No-gap response (µV) | 9.77±3.56 | 9.71±3.16 | 10.6±4.88 |
| Gap response (µV) | 6.24±2.42 | 8.17±3.00 | 8.39±2.61 |

GSI: gap-stimulus interval

Fig. 4. Gap PPI (gPPI) values (mean±standard error of the mean) elicited by different gap-stimulus intervals (GSIs). There was a statistically significant association between the GSI of 50 ms and the highest gPPI, as compared with the GSIs of 150 ms (∗p<0.05) and 250 ms (∗∗p<0.01).

Fig. 5. Auditory late responses elicited by only a gap. The background noise was narrow-band noise of 8 kHz with a bandwidth of 1 kHz (60 dB SPL). Gaps within the background noise evoked auditory responses similar to those evoked by sound stimuli. BGN: background noise.
was applied prior to the principal stimulus is assumed to reflect the PPI of the N1P2 response. Moreover, the gap-prepulse-induced auditory response may generate interference. Previously, several studies have reported that the AEPs evoked within 300 ms of multiple stimuli changed if they were overlaid with auditory responses [15,16]. Thus, we first measured the ALR under gap-only conditions (within background noise), and acquired evoked responses that were morphologically similar to those evoked by the principal sound stimulus (Fig. 5). All peak latencies were also similar (N1=0.56, P2=0.64). Thus, the decrease in the N1P2 amplitude of the gap response may reflect the simple mathematical summation of the two waveforms (the gap-only and no-gap responses). To explore this possibility, we corrected each ALR response by subtracting the auditory response elicited by the gap-only stimulus, and then averaged the responses to derive a corrected grand-average gPPI. The N1P2 amplitudes after correction of the data obtained at various GSIs are listed in Table 2.

The gPPI values after correction are shown in Fig. 6. The mean ± SD gPPI values were 18.59 ± 27.47%, 2.26 ± 24.62%, and 17.93 ± 12.7% at GSIs of 50, 150, and 250 ms, respectively. The value was highest at the GSI of 50 ms, decreased markedly at 150 ms, and increased again at 250 ms. We used the paired t-test to examine the effect of the GSI on the corrected gPPI values. The gPPI values at GSIs of 150 and 250 ms differed significantly (*p=0.011), but those at GSIs of 50 and 150 ms (*p=0.058) and 50 and 250 ms (*p=0.922) did not.

The gPPI values after zero padding

ALRs can be classified as exogenous and endogenous responses, depending on latency [17]. An exogenous response is highly dependent on the physical properties of the stimulus, whereas an endogenous response varies with the characteristics of the stimulus and psychological status of the subject [17]. As PPI was apparent even after a very brief GSI, we considered that such inhibition was more indicative of an exogenous rather than an endogenous response. Thus, we zero padded the gap-only response to minimize any effect of an endogenous response. All responses after zero-padding correction were averaged to obtain corrected grand-averaged gPPI values. The values at GSIs of 50, 150, and 250 ms are shown in Fig. 7. The highest value occurred at the GSI of 50 ms, and the lowest at the GSI of 250 ms.

All gPPI values associated with gap-only stimuli were subjected to zero-padding correction (Table 3). The mean ± SD gPPI values were 18.31 ± 22.17%, 14.49 ± 17.16%, and 13.07 ± 11.73% at GSIs of 50, 150, and 250 ms, respectively (Fig. 8). The highest gPPI value occurred at the GSI of 50 ms and then decreased as the GSI increased. We used the paired t-test to analyze the effect of the GSI; no difference was apparent among the GSIs of 50, 150, and 250 ms.

**Table 2.** Corrected N1P2 inter-peak amplitudes (mean±standard deviation) and gPPI values with- and without-gap prepulsing at different GSIs

| GSI (msec) | 50     | 150    | 250    |
|-----------|--------|--------|--------|
| No-gap response (µV) | 9.77±3.56 | 9.71±3.16 | 10.6±4.88 |
| Gap response (µV)    | 7.53±2.83 | 9.23±3.11 | 7.55±2.84 |

GSI: gap-stimulus interval

**Fig. 6.** Corrected gap PPI (gPPI) values (mean±standard error of the mean). After correction (subtraction of the gap-only stimulus), the gap-stimulus interval (GSI) of 150 ms, but not 250 ms, was associated with a near-zero gPPI (*p<0.05).

**Fig. 7.** Corrected grand-averaged gap PPI (gPPI) values after zero padding using different gap-stimulus intervals (GSIs). After zero padding, among the three GSIs evaluated, the GSI of 50 ms was associated with the highest gPPI. Notably, GSIs of 250 and 500 ms resulted in similar degrees of inhibition.

**Discussion**

We sought to identify ALR changes after gPPI in humans with normal hearing. First, we investigated gPPI under conditions very similar to those used in animal studies, and found

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Gap response (µV) 7.71 ± 2.78 8.28 ± 2.93 8.39 ± 2.61

Fig. 8. The gap PPI (gPPI) values (mean±standard error of the mean) after zero-padding correction. After correction, the gPPI did not vary according to the gap-stimulus interval (GSI).

Table 3. Corrected N1P2 inter-peak amplitudes including those obtained after zero padding (mean±standard deviation) with- and without-gap prepulsing at different GSIs

| GSI (msec) | 50    | 150   | 250   |
|-----------|-------|-------|-------|
| No-gap response (µV) | 9.77 ± 3.56 | 9.71 ± 3.16 | 10.6 ± 4.88 |
| Gap response (µV) | 7.71 ± 2.78 | 8.28 ± 2.93 | 8.39 ± 2.61 |

The ALR responses obtained using various GSIs showed that the ALR was inhibited by a gap prepulse. This was in line with the assumption that an AER can be generated using the gap prepulse paradigm [18]. Recently, various gap parameters have been shown to inhibit the effects of a gap prepulse on the ALR [19] and the eye-blink reflex in healthy adults [20]. Our results are similar to those of Ku et al. [13]. Furthermore, although direct comparisons of our data with those of animal studies would be inappropriate, we measured human responses derived using the gap prepulse paradigm; the results were similar to those of animal studies. Therefore, we support the use of the gap prepulse paradigm for tinnitus assessment in humans.

The ALR responses obtained using various GSIs showed that gap placement 50 ms prior to the principal stimulus resulted in the greatest inhibition of the response. Previous work on gPPI focused on gap duration; a longer gap prepulse was associated with greater inhibition, both for a simple gap prepulse and a gap following a prepulse when background noise was lacking between the gap and principal pulse [21]. Although a recent study on gPPI in healthy adult humans reported that the GSI did not affect the response [13], all GSIs were <100 ms. It is possible that other factors (such as gap offset or onset and background noise) affect the ALR response. Thus, the study does not confirm that the GSI between the gap and the principal stimulus is irrelevant in the context of the gap prepulse paradigm. We found that responses varied significantly with the GSI, even after the gap-only stimulus was subtracted. Our data and previous results suggest that further studies using GSIs of various durations <100 ms, with gap response subtraction, are required to confirm whether the GSI influences the gap prepulse paradigm in humans.

We found that the GSI of 250 ms also inhibited the response to the principal stimulus, albeit not as markedly as the GSIs of 50 and 150 ms. However, the GSI of 250 ms is clinically relevant, and temporal summation is impossible. Blumenthal [9] reported that the extent of the eye-blink inhibition by a prepulse depended on the prepulse duration and the interval between the prepulse offset and the onset of the principal pulse (the GSI in our present study). However, all GSIs applied were between 100 and 144 ms. Furthermore, whether the GSI of 50 ms was more inhibitory than GSIs of 150 or 250 ms was not investigated. Therefore, the ALR response under gap prepulse conditions is influenced not only by gap duration but also by a GSI >150 ms.

To identify the pure inhibitory effects of all GSIs, we subtracted the responses evoked by gap prepulsing; such prepulses were not inhibitory at the GSI of 150 ms. This is an interesting finding as it shows that the effect of the gap prepulse on the ALR depends on the GSI, and at some GSIs, the effect is saturated. We considered many other factors that might explain this result. As the AEP is very sensitive to measurement conditions and long measurement times, extreme caution is required during experiments for estimating such potentials. Grand-averaged data indicate gPPI trends. We sought to average all data obtained using the same GSIs; these are the most generalizable results of our study. Although statistical analysis was not possible, the most prominent inhibition was associated with the GSI of 50 ms, and the ALR response after a gap prepulse was more inhibited at shorter GSIs. Previous studies have proposed the use of the gap prepulse inhibition paradigm for tinnitus assessment in clinics [13, 22]. We also concluded that it could be a useful assessment tool for patients with tinnitus. However, according to our results, the gap prepulse itself and application of zero padding affect the result of the auditory brainstem response (ABR) after a gap prepulse stimulus, and these effects should be considered in this paradigm.

The N1P2 peak has been used to evaluate cochlear implants [23], explore the effects of drugs on auditory cognition [24], and functionally assess patients with schizophrenia [25]. Ku et al. [13] reported that the N1P2 amplitude peak was the most prominent of the various ALR peaks (N1, P2, N2, N1P2, and P2N2) in the gap prepulse paradigm. These studies suggest that our evaluation of the N1P2 peak of the gPPI ratio is appropriate when applying the paradigm. Sensory gating (inhibition of the AER by a prepulse) has also...
been evaluated [26-28]. Several works have explored the relationship between the eye-blink response and the AER (especially N1P2) in the context of the prepulse paradigm, and suggested that different mechanisms are in play [27,28]. As the eye-blink is a behavioral response that is easily influenced by a subject’s intention, assessment of an AER, especially an N1P2 response, is appropriate for tinnitus evaluation. However, the utility of the paradigm for children should be carefully considered, as ALR responses do not fully represent auditory sensitivity until late adolescence.

Multiple or paired stimuli delivered at various GSIs have been used to evaluate the auditory processing capacity [29]. However, when evaluating cortical responses, rapid re-presentation rates trigger response overlaps, making it difficult to interpret the results [14]. We found that a gap per se in the background noise evoked auditory responses. The response patterns associated with gaps in the background noise were very similar to the responses to sound stimuli. The relatively large baseline fluctuation noted in a previous study wherein gap prepulse stimuli was applied [19] may be attributed to overlaps. A recent study showed that use of the gPPI paradigm was associated with similar auditory responses in both tinnitus patients and normal subjects, perhaps because of a similar issue with overlap [22]. We used zero padding to eliminate any possibility of remnant wave fluctuation even after allowing sufficient time for a cortical response. After zero-padding correction, the gPPI did not vary according to the GSI. This challenges the conclusion of a previous study that the AEP varies with the time between the gap and principal stimulus in the gap prepulse paradigm. In conclusion, both the effect of a gap per se and that of the PPI should be considered when using the gap prepulse paradigm to evaluate humans.

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Conflicts of interest

The authors have no financial conflicts of interest.

Author Contributions

Conceptualization: Ilyong Park and Jae Yun Jung. Data curation: Jae-Hun Lee and Ilyong Park. Formal analysis: Jae-Hun Lee and Jae Yun Jung. Funding acquisition: Jae Yun Jung. Investigation: Jae Yun Jung. Methodology: Jae Yun Jung and Ilyong Park. Project administration: Jae Yun Jung and Ilyong Park. Resources: Ilyong Park. Software: Ilyong Park. Supervision: Jae Yun Jung and Ilyong Park. Validation: Jae Yun Jung. Writing—original draft: Jae-Hun Lee. Writing—review & editing: All authors.

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