Revisiting the Stimulation-Rate-Dependent Pattern Mismatch Negativity

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Course Title

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Due Date

7 Abstract

How does the brain process and represent successive sound in close temporal proximity? By investigating mismatch negativity (MMN) components, prior research (Sussman & Gumenyuk, 2005; Sussman, Ritter & Vaughan, 1998) has suggested that temporal 10 proximity plays an important role in how sounds are represented in auditory memory. 11 Here, we investigate how predictability affects the election of mismatch negativity 12 components in auditory sequences consisting of two tones (frequent tone A = 440 Hz, 13 rare tone B = 494 Hz, fixed SOA 100 ms). In the predictable condition, tones are 14 presented in a fixed order whereas in the unpredictable condition, standards and 15 deviants are presented in a pseudo-random order. We expect to find that B tones in the 16 unpredictable condition will elicit a significant MMN while B tones in the predictable 17 conditions will not. A repeating five-tone pattern was presented at several stimulus 18 rates (200, 400, 600, and 00 ms onset-to-onset) to determine at what temporal 19 proximity the five-tone repeating unit would be represented in memory. The mismatch 20 negativity component of event-related brain potentials was used to index how the 21 sounds were organized in memory when participants had no task with the sounds. Only 22 at the 200-ms onset-to-onset pace was the five-tone sequence unitized in memory. At 23 presentation rates of 400 ms and above, the regularity (a different frequency tone 24 occurred every fifth tone) was not detected and mismatch negativity was elicited by these tones in the sequence. The results show that temporal proximity plays a role in 26 unitizing successive sounds in auditory memory. These results also suggest that global relationships between successive sounds are represented at the level of auditory cortices.

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Revisiting the Stimulation-Rate-Dependent Pattern

Mismatch Negativity

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Introduction

How does the mind organize sequences of auditory stimuli?

At every moment, a rich spectro-temporal mixture of sounds hits our eardrums and causes the cochlea to vibrate, where stereocilia convert the vibrations into electrical impulses that race thrugh the vestibulocochlear nerve to the primary auditory cortex. This part of the hearing process is well understood but

59 **ASA**

How are these mere eletrical signals processed, combines and finally formed into meanignful peceptual experiences? While similar questions in the visual domain have intrigued scientists for a very long time and most notably lead to the emegence of the 62 Gestalt psychology in the early 20th century, long before the term auditory scene analysis was coined. While the Gestalt psychologists formulated very abstract rules, which in their own view should not be limited to the visual domain but rather represent universal laws of human perception, their research was almost exclusivly carried out in the field of visual perception. Mainstream auditory perception science was largly engaged in how very basic features of sound would connect to perception. Particularly the works of A. Bregmanns gave rise to a new framework called auditory scene analysis. This proposed framwork could serve a fundamental modle of human auditory perception and, in contrast to earlyer approaches, could address questions of how humans are able to form a coherent and meangingful representation of the auditory world. Bregman suggests that the brain uses streaming and segregation to form auditory objects from spectro-temporal infromationen.

Auditory scene analysis thereby relies on two different categories of grouping,
called sequential and simultaneous integration. Simultaneous or vertical integration
refers to the grouping of concurrent properties into one or more separable auditory
objects, a process informed by temporal cues like common onset and offset, spectral and
spatial characteristics among others. Sequential integration on the other hand describes

how temporally distinct sounds are merged into one or multiple coherently perceived stream (contrary to simulatinous grouping, only one such stream can be activly perceived at any time). While vertical and horizontal grouping can come to different and therefore competing results (???), sequential grouping often takes precedence over cues for simulatoius integration (???),.

When presented with a series of similar or repeated auditory events, rare 85 deviants (termed oddballs) result in a negative deflection of event-related responses 86 measured with EEG. These alterations are indexed by the missmatch negativity (MMN) 87 component obtained by subtracting the reponse to deviant events from the response to 88 standard events. Negativity is strongest in the fronto-temporal area of the scalp with a 89 peak latency ranging from 100 to 250 ms after stimulus onset. MMN components observed in magnetoencephalography (MEG) are called MMNm. There is a long line of 91 reserach suggesting that MMN ist pre-attentive (???). MMN has been tradionally described as an index of discrepancy between auditory input and the memory trace of 93 the preceding standard stimuli (Paavilainen, 2013). The eliction MMN is not restricted to the reptition of physically identical stimuli but can also be observed when deviant 95 events are of complex nature, e.g. when abstract auditory regularities are violated (Paavilainen, 2013). The regularities can come in the form of relationships between two 97 Saarinen et al. (1992) or multiple tones (Alain et al., 1994; Nordby et al., 1988; Schröger et al., 1996) and as with first-order MMNs could sucesdully observed in infants 99 (He et al., 2009).

Sussman et al. (1998) presented participants with a recurring five-tone pattern (A-A-A-A-A-A-A-A-A-A-A-A), "-" indicating silence between the tones). Differences in ERP following A and B tones were compared for rapid (SOA of 100 ms) and slow (SOA of 1200 ms) stimulation rates. MMN were only elicted, when stimuli were presented at a fast pace but no evidence was found at a SOA of 100 ms. In a subsquent study, Sussman & Gumenyuk (2005) used the same pattern at different SOA paces (200 ms, 400 ms, 800 ms). They also included a controll condition in which tones presentation

was pseudo-random (while keeping the porbability of B tones at 20%). Simmilarly to
their prevous study, periodic presentation at 400 ms or slower stimulus-to-stimulis pace
elicted a MMN, while at a stimulution rate of 200 ms such evidence was absent. They
attributed this to competing predicitve rules by which tones were either grouped
seuqutially or treated as distinct tones. Sussman et al. attributed this observation to
sensory memory limitations, i.e., only when auditory memory could accommodate
enough repetitions of the five-tone pattern tones could be integrated into a coherent
representation and thereby allow the extraction of the underlying relationship.

- 1. replicated the procedure by Sussman et al. in an in-class setting.
- draw on the notion
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Recently, the missmatch negativity has been described in the predictive coding framework.

A simmilar explaination can be offered in terms of predictive coding. Predictive 121 coding is a biologically plausible model proposing prediction as the key feature of 122 perception that was first described in the cortical visual system (Rao & Ballard, 1999). 123 Taking a broader view, predictive coding is part of a research tradition taking a probabilistic (or Baysian) approaches to brain function. Chharacterizing the brain as an 125 inference mashine, this line of resoning traces back to Herman von Helmholtz's work in 126 the late 19th century. In sharp contrast to traditional stimulis-driven models that 127 describe the act of perception as a bottom-up process, in probabilistic terms, perception 128 is not the direct result of sensory input, but is built by combining sensory input with 129 predictions with internal, probabilistic generative models. Using prior knowledge abou 130 the world, these models are assumed to constantly create probabilistic versions of 131 expected sensory input. Predictive coding more specifically suggests that at every processing state, predictions and actual input is constantly compared and only their 133 difference, called *prediction error*, is propagated. Perception is thus seen as the process

of improving the internal generative model by using sensory input to minimize prediciton error. Because of that, predictive coding is sometimes casually reffered to as controlled hallucination.

However, there are multiple in shortcoming in

Precision weighting

140 Hypothesis

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As layed out above, we hypothize that that two possible rules cyrry predictive value: 141 Firstly, the presentation ratio of A and B tones (9 to 1) can be used to make 142 proportion-dependent predicitons as used in classical oddball-paradigms. When tones 143 are presented in a regular fashion, as it is the cas ein the predictable condition, the 144 extracted pattern might also be used to predict the next tone. Thus, two plausible but 145 concurrent rules might guide predictions in the predictable condition, while in the 146 random codntion only the proportion-based regularity offers information to form predictions about upcoming tones. As has been shown before (???), pattern-based 148 regularities are commonly found to take precedence over proportion-based regularities. 149 If this is indeed the case, B-tones in the predictable condtiion should not be considered a 150 missmatch and thus should not elict a MMN. In contrast, since there is no way to predict B-tones in the random condition, these tones would be still considered as 152 deviant events and therefore expected to generate a MMN. Following this notion, one would also expect an expectation violation when predictable B-tones are replaced by 154 A-tones, although they would be considered "standard" events when prediction is purely guided by proportion. This would be also in line with (Sussman et al., 1998; 156 Sussman & Gumenyuk, 2005) interpretation of the original results. 157

pattern regularity

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159 If the fixed order of the tones in the predictable state leads to a prediction of the 160 B-tones, i.e. if the pattern regularity is extracted and the proportional regularity is

irrelevant in the predictable context, we expect that the difference of 161 predictable-BAAAAAA "B" and predictable-BAAA "A" is (significantly) less negative 162 than the difference of random-BAAAAAA "B" minus random-BAAA "A". In addition, 163 we assume that the difference of predictable-BAAAA "B" and predictable-BAAA "A" is not significantly different from zero, while the difference of random-BAAAAAA "B" 165 and random-BAAA "A" is significantly less than zero. In addition, in the predictable state, the interruption of the pattern regularity with an A tone should produce a 167 significant negativity. This means that the difference between predictable-BAAAA "A" 168 and predictable-BAAA "A" is significantly more negative than the difference between 169 random-BAAAA "A" and random-BAAA "A". In addition, we expect the difference 170 between predictable-BAAA "A" and predictable-BAAA "A" to be less than zero, 171 while the difference between random-BAAA "A" and random-BAAA "A" is not significantly different from zero. 173

, this process has been characterized as stimulis-driven

In this model, these predictions are what is actually perceived while sensory information are

Predictive coding is a theoretical model based on in the fundamental idea that

• what are auditory objects?

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- what influences their formation?
- what is predicitve coding?
- how can objects be used for prediction?
 - what happends when rules conflict?
 - MMN and preidctive coding?

184 Predictive Coding

- brain as inference machine
- a. Auditory scene analysis

- b. Sussman et al.
- c. Scharf & Müller
- 189 d.

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Methods

Data Acquisition

192 Participants

Twenty-three psychology undergraduate students (2 males, average age 22.6 193 yrs., SD = 5.57, range 18 - 42 yrs.) were recruited at the Institute of Psychology at the 194 University of Leipzig. All participants reported good general health, normal hearing 195 and had normal or corrected-to-normal vision. Written informed consent was obtained 196 before the experiment. One-third (34.8%) of participants spent time enaging in musical 197 activities at time of survey, while 8.7% had no prior experience in music training. 198 Handedness was asseced using a modified version of the Edinburgh Handedness 199 Inventory (Oldfield, 1971, see appendix). A majority (00%) of parcicipants favored the 200 right hand. Participants were blinded in respect to the purpose of the experiment and 201 received course credit in compensation. 202

Study 2 Twenty healthy participants (0 males, average age 00.0 yrs., SD = 0.00,
range 00 - 00 yrs.) were recruited. Participants gave informed consent and reported
normal hearing and corrected or corrected-to-normal vision. All participants were naive
regarding the purpose of the experiment and were compensated in cource credit or
money. 00 participants (00%) had received musical training in the last 5 years before
the experiment while 00 (00%) reported no musical experiance. In addition,
participants reported if streaming occurred during the presentation of the tones.

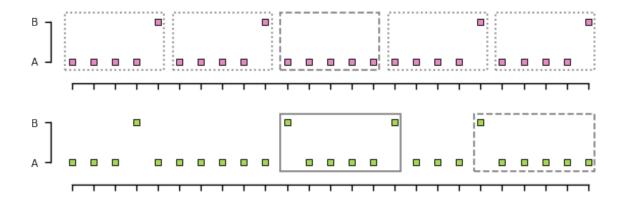


Figure 1. Tones of two different frequencies (A=440 Hz, B=449 Hz) were presented in two blocked conditions: In the "predictable" condition (top half), tones followed a simple pattern in which a single B-tone followed four A-tones. Some designated B-tones were replaced by A-tones ("pattern deviants"). In the "random" condition (lower half), tones were presented in a pseudo-random fashion ()

210 Stimuli

Stimuli consisted of pure sinusoidal tones with a duration of 50 ms (including a 10 ms 211 cosine on/off ramp), presented isochronously at a stimulation onsets asynchrony (SOA) 212 of 100 ms for study 1 and 150 ms for study 2. Participants where seated in a 213 electromagnetically shielded and sound-proofed cabin while administering a total of 40 214 blocks containing a mixture of frequent 440 Hz tones ("A" tones) and infrequent 449 Hz tones ("B" tones). In one half of the blocks, tones were presented in pseudo-random 216 order (e.g. A-A-A-B-A-B-A), "random" condition), while in the remaining block tone presentation followed a simple pattern in which a five-tone-sequence of four frequent 218 tones and one infrequent tone (i.e. A-A-A-B) was repeated cyclically ("predictable" condition). The ratio of frequent and infrequent tones was 10% for both conditions. 220 Within the predictable condition, 10% of designated (infrequent) B tones were replaced 221 by A tones, resulting in sporadic five-tone sequences consisting solely of A tones 222 (i.e. A-A-A-A), thus violating the predictability rule. To assure comparability of local 223 histories between tones in both conditions, randomly arranged tones were interspersed 224

with sequences mimicking aforementioned patterns from the predictable condition (B-A-A-A-A-B and B-A-A-A-A) in the random condition. A grand total of 2000 tones in study 1 and 4000 tones in study 2 were delivered to each participant.

228 Data Acquisition

Electrophysiological data was recorded from active silver-silver-chloride (Ag-AgCl)
electrodes using an ActiveTwo amplifier (BioSemi B.V., Amsterdam, The Netherlands).
Acquisition was monitored online to ensure optimal data quality. A total of 39 channels
were obtained using a 32-electrode-cap and 7 external electrodes. Scalp electrode
locations conformed to the international 10–20 system. Horizontal and vertical eye
movement was obtained using two bipolar configurations with electrodes placed around
the lateral canthi of the eyes and above and below the right eye. Additionally,
electrodes were placed on the tip of the nose and at the left and right mastoid sites.

Data was sampled at 512 Hz and on-line filtered at 1000 Hz.

238 Analysis Pipeline

Data prepossessing was implemented using a custom pipeline based on the MNE Python software package (Gramfort, 2013) using Python 3.7. All computations were carried out on a cluster operated by the University Computation Center of the University of Leipzig. Code used in thesis is publicly available at https://github.com/marcpabst/xmas-oddballmatch.

244 Bad Channel Detection and Interpolation

Firstly, EEG data was subject to the ZapLine procedure (de Cheveigné, 2020) to
remove line noise contamination. A fivefold detection procedure as described by
Bigdely-Shamlo et al. (2015) was then used to detect and subsequently interpolate bad
channels. This specifically included the detection of channels thain contain prolonged
segments with verry small values (i.e. flat channels), the exclusion of channels based on
robust standard deviation (deviation criterion), unusually pronounced high-frequency

noise (noisiness criterion), and the removal of channels that were poorly predicted by
nearby channels (correlation criterion and predictability criterion). Channels considered
bad by one or more of these methods were removed and interpolated using spherical
splines (Perrin et al., 1989). Electrode locations for interpolations were informed by the
BESA Spherical Head Model.

256 Independent Component Analysis

Given the $\frac{1}{f}$ power spectral density of EEG data, the estimation independent components (ICs) by independent component analysis (ICA) would be strongly influenced by high-frequency noise that is ususally considere brain-irrelevant [reference]. To mitigate this effect, a 1-Hz-high-pass filter (134th order hamming-windowed FIR) was applied prior to ICA (Winkler et al., 2015).

To further reduce artifacts, Artifact Subspace Reconstruction (ASR, Mullen et al., 2015) was used to identify parts of the data with unusual characteristics (bursts) which were subsequently removed. ICA was then carried out using the *Picard* algorithm (Ablin et al., 2018, 2017) on PCA-whitened data. To avoid rank-deficiency when extracting components from data with one or more interpolated channels, PCA was also used for dimensionality reduction to obtain full-ranked data.

The EEGLAB (version 2020.0, Delorme & Makeig, 2004) software package and
the IClabel plugin (version 1.2.6, Pion-Tonachini et al., 2019) were used to
automatically classify estimated components. Only components clearly classified
(i.e. confidence above 50%) as resulting from either eye movement, muscular, or
heartbeat activity were zeroed-out in the mixing matrix before inversely transform ICs.

Filtering $\mathbf{Filtering}$

In line with recommendations from Widmann et al. (2015) and de Cheveigné & Nelken (2019), a ORDER finite impulse response (FIR) bandpass filter from 0.1 Hz to 30 Hz was applied in forward direction only (Hamming window with 0.0194 passband ripple

277 and 53 dB stopband attenuation).

278 Epoching and Averaging

Continuous data was epoched into 400 ms long segments around stimulus onsets. This included a 100 ms pre-stimulus interval which was used to perform baseline correction 280 by subtracting its mean amplitude from each epoch. The AutoReject software package 281 (Jas et al., 2017) was used to reject bad epochs. The AutoReject algorithm uses 282 cross-validations and basyan optimaziation to calculate channel-wise peak-to-peak 283 amplitude thresholds that minimizes the root mean square error (RMSE) between the 284 mean (after removing the trials marked as bad) and the median of the data (including 285 all trials). For epochs where only a small subset of channels exceeded the critical 286 threshold, bad channels were interpolated instead of removing the whole epoch. 287

288 Statistical Analysis

289 Standard Repetition Effects

290 **MMN**

The dependent variable for analysing missmatch response was calculated by averaging 291 amplitudes within a time window of ± 25 ms around the maximum negativity obtained 292 by subtracting the mean ERP timecourse following the (expected) deviant event from 293 the ERP following the (expected) standard event. To obtain mean amplitudes, ERPs to 294 4th position A tones (A-A-A-X, **boldface** indicates the tone of interest) and B tones 295 (A-A-A-B) were averaged separatly for both the random and the predictable 296 condition. For the random condition, only tones that were part of a sequence mimicking 297 the patterns from the predictable condition were included. 298

A three-way analysis of variance for repeated measures with the factors stimulus onset asynchronoy (100 ms vs. 150 ms stimulus onset asynchrony), condition (predictabe vs random presentation) and stimulus type (A tones vs. B tone).

To further analyse Following the pre-registration, a two-way analysis of variance for repeated measures to test for significant differences of mean amplitudes in the MMN window between standard and deviant tones (stimulus type) depending on the condition (predictable vs. random) was calculated seperatly for both 100 ms and 150 ms presentation. In line with Sussman & Gumenyuk (2005), FZ, F3, F4, FC1, and FC2 electrode locations were averaged. Greenhouse-Geisser correction for lack of sphericity was applied when appropriate.

For post-hock comparison, two-tailed Student's t-test were calculated for .

P-values were corrected for multiple comparisons by using the Benjamini-Hochberg

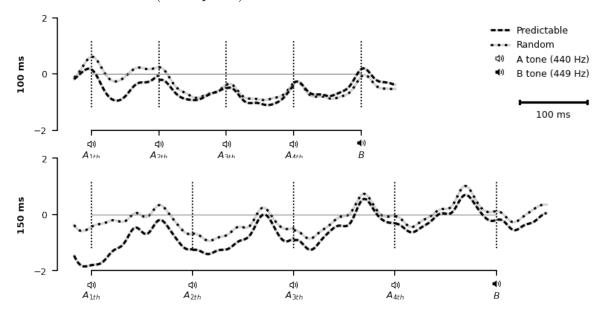
procedure.

Results

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Figure X shows EEG waveform averages (pooled FZ, F3, F4, FC1, and FC2 electrode locations) for five-tone sequences (A-A-A-B) presented in a *predictable* (top panel) and *random* contexts (lower panel).



Grand averages of the fronto-cluster (pooled FZ, F3, F4, FC1, and FC2 electrode locations) of event-related potentials for A tones (A-A-A-X) and B tones

(A-A-A-B) and their difference (B-ERP - A-ERP) are displayed in Figure X for both 320 100 ms (left panel) and 150 ms (right panel) stimulus onset asynchrony. The top half of 321 each panel shows ERPs in the predictable condition while the lower halfs depicts ERPs 322 in the random condition. For both presentation rates, clear rythms matching the 323 presentation frequency of 10 Hz (100 ms) respectively 6.667 Hz (150 ms) are seen 324 resulting from overlap of neighboring tones. Panels also show the distribution of 325 amplitude differences in the MMN latency window as defined above (110 ms to 160 ms 326 after stimulus onset) across participants and the difference of topographic maps 327 averaged over the same interval. Simmilarly, waveforms at pooled mastoid sites are 328 shown in Figure X. 329

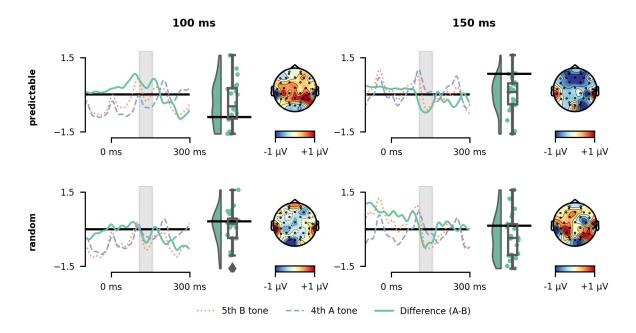


Figure 2. ERP grand averages (pooled FZ, F3, F4, FC1, and FC2 electrode locations) for an SOA of 100 ms (left) and 150 ms (right), for A tones (A-A-A-A-X, blue dashed lines) and B tones (A-A-A-A-B, orange dashed line) and their difference (B - A, green solid line). Upper panels show ERPs for tones presented in a predcitable pattern (predcitable condition) while lower panels show ERPs for tones presented in pseudo-random order (random condition). Shaded area marks MMN latency window (110 ms to 160 ms) used to calculate the distribution of amplitude differences across participants (middle of each panel) and the difference of topographic maps averaged over the same interval (right of each panel).

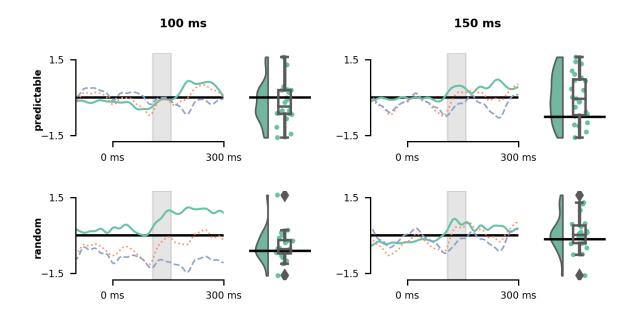


Figure 3. ERP grand averages (pooled M1, M2 electrode locations) for an SOA of 100 ms (left) and 150 ms (right), for A tones (A-A-A-A-X, blue dashed lines) and B tones (A-A-A-A-B, orange dashed line) and their difference (B - A, green solid line). Upper panels show ERPs for tones presented in a predcitable pattern (predcitable condition) while lower panels show ERPs for tones presented in pseudo-random order (random condition). Shaded area marks MMN latency window (110 ms to 160 ms) used to calculate the distribution of amplitude differences across participants.

Descriptively, regardless of tone presentation rate, evoked responses for 330 fornto-central electrode locations in the MMN latency window were more negative for B 331 tones than for A tones in the random condition (100-ms-SOA: $\Delta M = -0.358 \,\mu V$, 332 150-ms-SOA: $\Delta M = -0.555 \,\mu V$) This was also true when tones were presented in a 333 predictable fashion, but only for the slower presentation rate ($\Delta M = -0.582 \,\mu V$)). In 334 contrast, when predcitable tone patterns were presented at the faster 335 100-ms-presentation rate, B tones were descriptively more positive than A tones 336 $(\Delta M = 0.383 \,\mu V).$ 337

Descriptice comparison of evoked responses from pooled left and right mostoids revealed that pseudo-randomly presented B tones were more positive in the MMN latency window than A tones (100-ms-SOA: $\Delta M = 0.746 \,\mu V$, 150-ms-SOA: $\Delta M = 0.510 \,\mu V$). A simmilar observation could be made for precitable B tones
compared to the preceding A tones at a SOA of 150 ms ($\Delta M = 0.399 \,\mu V$)) but not for
the faster presentation rate ($\Delta M = -0.132 \,\mu V$). Mean amplitudes in the MMN latency
window and their standard deviantions (SD) for all conditions are shown in Table X.

| SOA | Condition | StimulusType | Mean | SD Mean | SD |
|-----|-------------|--------------|---------|--------------|------|
| 100 | predictable | A | -0.431 | 1.23 -0.052 | 1.51 |
| | | В | -0.0477 | 1.22 -0.184 | 1.56 |
| | random | A | -0.225 | 1.82 -1.04 | 2.64 |
| | | В | -0.583 | 2.16 -0.296 | 3.23 |
| 150 | predictable | A | 0.25 | 0.967 -0.349 | 1.19 |
| | | В | -0.331 | 1.09 0.0492 | 1.33 |
| | random | A | 0.0233 | 1.75 -0.292 | 1.64 |
| | | В | -0.531 | 1.82 0.218 | 2.38 |

Statistical analysis supports these findings. For the 100 ms stimulation rate, the three-way ANOVA yielded a significant 3-way interaction effect condition x stimulus type x electrode $(F(1,19)=7.53,\,p=0.0130)$ but revelad no main effects for neither stimulus type $(F(1,19)=1.05,\,p=0.3180)$, condition $(F(1,19)=0.83,\,p=0.3730)$, nor electrode $(F(1,19)=0.04,\,p=0.8520)$. In contrast, for tones presented at a SOA of 150 ms, an effect was only found for the two-way interaction term stimulus type x electrode $(F(1,22)=20.76,\,p=0.0002)$. Mean amplitudes in the MMN latency window did not differ for factors stimulus type $(F(1,22)=0.32,\,p=0.5790)$, electrode ().

Two-way ANOVAs (Condition x Stimulus Type) were carried out seperatly for pooled fronto-central and mostoid electrode locations. For 100 ms tone presentation rate, the Condition x StimulusType interaction only revealed a significant effect for the fronto-central electrode cluster (F(1,19)=16.75, p=0.0006) but not for pooled mastoid sites (F(1,19)=2.37, p=0.1410) indicating that the 3-way interaction effect condition x stimulus type x electrode is indeed driven by the amplitude differnces in te

fronto-central electrode locations. Contrary to this, for the 150 ms presentation rate,
main effects for *stimulus type* were significant for both fronto-central and mastoid sites,
suggesting that there was both a MMN at fronto-central locations as well as a
polarity-reversal at the mastoid electrodes.

For the 150 ms stimulation rate, the 2-way ANOVA yielded a significant main effect for stimulus type $(F(1,22)=22.67,\,p=0.0001)$ but not for condition $(F(1,22)=0.95,\,p=0.3410)$ or stimulus type x condition interaction $(F(1,22)=0.03,\,p=0.8680)$. In contrast, when presenting tones with a stimulus-onset-assychmony of 100 ms, no such effects were found for the factor condition $(F(1,22)=0.95,\,p=0.3410)$, stimulus type $(F(1,22)=22.67,\,p=0.0001)$, or interaction $(F(1,22)=0.03,\,p=0.8680)$.

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Table 1

Results of the 3-way ANOVA (condition x stimulus x electrode) for repeated measures conducted on the mean ERP-amplitudes (time window 111 - 161 ms) at electrode Fz (upper section). The significant interaction between the three factors included was further analyzed by 2-way ANOVAS (stimulus x electrode) conducted separately for the random condition (middle section) and the predictable condition (lower section).

| | Effect | DFn | DFd | ${f F}$ | p | p<.05 | ges |
|---------|--|-----|-----|---------|----------|-------|----------|
| | Condition | 1 | 19 | 0.831 | 0.373 | | 0.008 |
| | StimulusType | 1 | 19 | 1.05 | 0.318 | | 0.002 |
| α | Electrode | 1 | 19 | 0.036 | 0.852 | | 0.000331 |
| 100 ms | Condition x StimulusType | 1 | 19 | 0.051 | 0.823 | | 7.55e-05 |
| 1 | Condition x Electrode | 1 | 19 | 0.763 | 0.393 | | 0.002 |
| | StimulusType x Electrode | 1 | 19 | 0.797 | 0.383 | | 0.001 |
| | Condition x Stimulus Type x Electrode | 1 | 19 | 7.53 | 0.013 | * | 0.01 |
| | Condition | 1 | 22 | 0.08 | 0.78 | | 0.000263 |
| | StimulusType | 1 | 22 | 0.317 | 0.579 | | 0.000339 |
| α | Electrode | 1 | 22 | 0.035 | 0.854 | | 0.000301 |
| 150 ms | Condition x StimulusType | 1 | 22 | 0.16 | 0.693 | | 0.000124 |
| П | Condition x Electrode | 1 | 22 | 1.13 | 0.299 | | 0.003 |
| | StimulusType x Electrode | 1 | 22 | 20.8 | 0.000155 | * | 0.026 |
| | Condition x StimulusType x Electrode | 1 | 22 | 0.053 | 0.819 | | 4.63e-05 |

Table 2

Results of the 3-way ANOVA (condition x stimulus x electrode) for repeated measures conducted on the mean ERP-amplitudes (time window 111 - 161 ms) at electrode Fz (upper section). The significant interaction between the three factors included was further analyzed by 2-way ANOVAS (stimulus x electrode) conducted separately for the random condition (middle section) and the predictable condition (lower section).

| | | Effect | DFn | DFd | \mathbf{F} | p | p<.05 | ges |
|--------|----------|--------------------------|-----|-----|--------------|-------|-------|---------|
| | Frontal | Condition | 1 | 19 | 0.16 | .694 | | 0.003 |
| | | StimulusType | 1 | 19 | 0.006 | .938 | | 1.5e-05 |
| 100 ms | | Condition x StimulusType | 1 | 19 | 16.7 | <.001 | * | 0.013 |
| 100 | Mastoids | Condition | 1 | 19 | 1.28 | .272 | | 0.014 |
| | | StimulusType | 1 | 19 | 1.21 | .285 | | 0.004 |
| | | Condition x StimulusType | 1 | 19 | 2.37 | .141 | | 0.009 |
| | | | | | | | | |
| | Frontal | Condition | 1 | 22 | 0.947 | .341 | | 0.006 |
| | | StimulusType | 1 | 22 | 22.7 | <.001 | * | 0.038 |
| 150 ms | | Condition x StimulusType | 1 | 22 | 0.028 | .868 | | 2.2e-05 |
| 150 | Mastoids | Condition | 1 | 22 | 0.206 | .655 | | 0.001 |
| | | StimulusType | 1 | 22 | 6.56 | .018 | * | 0.018 |
| | | Condition x StimulusType | 1 | 22 | 0.122 | .730 | | 0.00028 |