Variability in L2 phonemic learning originates from speech-specific capabilities: An MMN study on late bilinguals

BEGOÑA DÍAZ
Center for Brain and Cognition, Department of Technology, Universitat Pompeu Fabra, Barcelona, Spain
HOLGER MITTERER
Department of Cognitive Science, University of Malta, Msida, Malta
MIRJAM BROERSMA
Centre for Language Studies, Radboud University, Nijmegen, The Netherlands
Max Planck Institute for Psycholinguistics, Nijmegen, The Netherlands
CARLES ESCERA
Institute for Brain, Cognition, and Behavior (IR3C), University of Barcelona, Barcelona, Spain
Cognitive Neuroscience Research Group, Department of Psychiatry and Clinical Psychobiology, University of Barcelona, Barcelona, Spain
NÚRIA SEBASTIÁN-GALLÉS
Center for Brain and Cognition, Department of Technology, Universitat Pompeu Fabra, Barcelona, Spain

Address for correspondence:
Begoña Díaz
Dept. Technology
C. Roc Boronat 138
08018 Barcelona, Spain
begona.diaz@upf.edu

ABSTRACT

People differ in their ability to perceive second language (L2) sounds. In early bilinguals the variability in learning L2 phonemes stems from speech-specific capabilities (Díaz, Baus, Escera, Costa & Sebastián-Gallés, 2008). The present study addresses whether speech-specific capabilities similarly explain variability in late bilinguals. Event-related potentials were recorded (using a design similar to Díaz et al., 2008) in two groups of late Dutch–English bilinguals who were good or poor in overtly discriminating the L2 English vowels /ɛ-æ/. The mismatch negativity, an index of discrimination sensitivity, was similar between the groups in conditions involving pure tones (of different length, frequency, and presentation order) but was attenuated in poor L2 perceivers for native, unknown, and L2 phonemes. These results suggest that variability in L2 phonemic learning originates from speech-specific capabilities and imply a continuity of L2 phonemic learning mechanisms throughout the lifespan.

1. Introduction

Economic and cultural globalization has turned the learning of foreign languages into a socio-economic need. Consequently, the learning of at least one foreign language is a compulsory subject in the academic curricula of most countries in the world (around 81% of the 119 countries analysed in the UNESCO World Report on Cultural Diversity and Intercultural Dialogue in 2009[1]). One of the biggest difficulties for learners of a second language (L2) is to accurately perceive and produce the speech sounds of the new language, and only a few individuals manage to achieve high proficiency levels in these skills (Bongaerts, 1999; Sebastián-Gallés & Baus, 2005, Sebastián-Gallés & Díaz, 2012). The perception and production of speech sounds is fundamental to mastering any language: without an accurate perception of speech sounds, listeners cannot understand what others say, and without an accurate production, they cannot make themselves understood. Evidence has accumulated that in speech perception, difficulties with the perception of L2 sounds leads to inappropriate lexical competitors being active, slowing down the recognition of the intended word (Broersma, 2012; Broersma & Cutler, 2008; Broersma & Cutler, 2011; Pallier, Colomé & Sebastián-Gallés, 2001; Sebastián-Gallés, Echeverría & Bosch, 2005; Weber & Cutler, 2004;), resulting in very real problems for L2 learners in communicative situations. Whereas L2 learners can often find ways around such communicative problems by using compensatory strategies (Poulisse, Bongaerts & Kellerman, 1990), it still makes the task of communicating more difficult for them, as it makes them less efficient language users. Mastering speech sounds well is thus essential to L2 learning success.

Factors such as age of acquisition (AoA) of the L2, amount of exposure to the L2, and motivation to learn the L2 play a crucial role in ultimate L2 attainment (Birdsong, 1999), yet they do not fully account for individual variability in L2 learning.

Training studies, as well as naturalistic language studies, have shown that individuals that are similar in the aforementioned relevant factors for L2 learning do not profit from equivalent phonetic experience to the same extent (Díaz, Mitterer, Broersma & Sebastián-Gallés, 2012; Golestani, Molko, Dehaene, LeBihan & Pallier, 2007; Golestani, Paus & Zatorre, 2002; Sebastián-Gallés & Baus, 2005; Tremblay, Kraus & McGee, 1998). The present study aims to investigate the origin of the distinct outcomes in the learning of a second language. Finding the origin of individual variability in the learning of the L2 phonemes is crucial for predicting final L2 acquisition and may help to design tailored L2 learning protocols that maximize the success of L2 learning.

The present aim is addressed by using the same experimental design, with slight modifications, as a previous study with early bilinguals (Díaz, Baus, Escera, Costa & Sebastián-Gallés, 2008). This previous study suggested that there is a uniquely linguistic capability that varies across individuals and predicts L2 phonological learning (Díaz et al., 2008). In that study, early Spanish–Catalan bilinguals were selected according to their proficiency in discriminating two L2 (Catalan) vowels in several behavioural tasks (for a description of the tasks see Pallier, Bosch & Sebastián-Gallés, 1997; Sebastián-Gallés et al., 2005; Sebastián-Gallés & Soto-Faraco, 1999). The two groups were intended to represent the extreme endpoints of non-native phoneme perception and were categorized as good perceivers (GPs) or poor perceivers (PPs). Participants' sensitivity to acoustic and speech changes was assessed by means of an event-related brain potential, the mismatch negativity (MMN).

The MMN is elicited when the auditory perceptual system detects a mismatch between frequently repeated stimuli (standard stimuli) and stimuli differing in at least one acoustic feature (deviant stimuli) (Näätänen, 2001; Näätänen, Gaillard & Mäntysalo, 1978). The MMN is a fronto-central negativity, with a reversed polarity at

temporal sites, that peaks around 100–250 ms after the detection of a change in the auditory signal and is sensitive to changes both in pure tones and speech sounds (Kujala & Näätänen, 2010; Näätänen, Lehtokoski, Lennes, Cheour, Huotilainen, Iivonen, Vainio, Alku, Ilmoniemi, Luuk, Allik, Sinkkonen, & Alho, 1997; Nenonen, Shestakova, Huotilainen & Näätänen, 2005; Winkler, Kujala, Tiitinen, Sivonen, Alku, Lehtokoski, Czigler, Csépe, Ilmoniemi & Näätänen, 1999). The MMN can be elicited during passive listening (e.g., while attending to a silent movie) and is, therefore, not influenced by engagement of cognitive processes related to task demands, strategies or motivation. Crucially, the amplitude of the MMN is directly related to the magnitude of the perceived change (Amenedo & Escera, 2000; Näätänen, 2001). There are two brain sources that contribute to the generation of the MMN: a superior temporal generator (related to the processing of the auditory sensory input against a memory trace) and a frontal generator (related to the orienting of attention towards a detected change in the auditory input) (Escera, Alho, Winkler & Näätänen, 1998; Giard, Perrin, Pernier & Bouchet, 1990; Näätänen, 1990; Yago, Escera, Alho & Giard, 2001). Note that occasionally an additional parietal MMN generator has been reported (Lavikainen, Huotilainen, Pekkonen, Ilmoniemi & Näätänen, 1994). The functional dissociation of the MMN temporal and frontal generators allows us to investigate the contributions of each MMN source to potential discrimination problems in poor L2 perceivers. Whereas ERPs cannot directly measure the activity of the MMN generators, it has been claimed that their activity can be inferred from the amplitude and latency of the MMN at frontal and mastoid electrodes. ERP source analyses indicate that temporal electrodes capture the activity only from the temporal MMN generator while the frontal electrodes receive contributions from the temporal and frontal electrodes (Giard et al., 1990). Experimental evidence showing that the frontal and temporal MMN subcomponents can be

manipulated independently support this observation. The frontal MMN, but not the temporal one, increases as a function of the probability of the standards stimuli (Sato, Hirooki, Tomiharu, Takeyuki, Naoko, Tadayoshi & Sunao, 2000), diminishes with alcohol intake (Jääskeläinen, Pekkonen, Hirvonen, Sillanaukee & Näätänen, 1996) and decreases over time (Sussman & Winkler, 2001). The temporal MMN subcomponent, but not the frontal one, shows additive MMNs for simultaneous frequency and intensity deviations (Paavilainen, Mikkonen, Kilpeläinen, Lehtinen, Saarela & Tapola , 2003) and is diminished for speech in temporal lobe epileptic patients (Hara, Ohta, Miyajima, Hara, Iino, Matsuda, Watanabe, Matsushima, Maehara & Matsuura , 2012) and stutterers (Corbera, Corral, Escera & Idiazábal, 2005) when compared to controls. The scalp distribution of MMN differences may contribute to differentiating between perceptual (related to the temporal MMN generator) and attentional (related to the frontal MMN generator) origins of differences in the MMN amplitude between groups.

The GPs and PPs tested by Díaz et al. (2008) showed equivalent MMN responses to tones of different frequencies, durations, and predictability in their presentation order. This lack of differences indicated that GPs and PPs had similar general acoustic-perceptual capacities. When the participants were tested with phonemes that belonged either to their native language (the Spanish vowels /e/ and /o/) or to an unknown language (the Estonian vowels /ö/ and/o/), GPs showed larger MMNs compared to PPs, i.e., greater discrimination sensitivity, not only to the native but also to the unknown vowel contrasts. The larger MMN in GPs for phonemes together with the lack of differences for non-linguistic stimuli between the groups was interpreted as a demonstration of a uniquely linguistic ability for (native and non-native) language learning. The difference in the amplitude of the MMN for speech between the groups was present at frontal electrodes, but absent at mastoids, which suggested that the two

groups differed in the activity of the frontal MMN generator, whose function is to reorient attention to deviations in the auditory signal. In line, a previous study (Sebastián-Gallés, Soriano-Mas, Baus, Díaz, Ressel, Pallier, Costa & Pujol, 2012) found larger white matter volume for PPs, as compared to GPs, in a right frontal brain area (i.e., the insulo/fronto-opercular region) and the volume correlated positively with the MMN amplitude (the less negative MMN, the more white matter volume).

Díaz et al. (2008)'s findings contrast with those from training studies of general auditory (i.e., non speech-specific) capabilities being at the basis of individual differences for phonological learning (Golestani et al., 2007; Lengeris & Hazan, 2010; Wong, Perrachione & Parrish, 2007; Wong, Warrier, Penhune, Roy, Sadehh, Parrish & Zatorre, 2008). For instance, Lengeris & Hazan (2010) found a positive relation between the learning of new vowels and frequency discrimination for non-speech sounds, and Wong et al. (2007) showed a positive correlation between pitch pattern identification in non-speech sounds and the learning of words of a tonal language. The contrasting results between training studies (i.e., general auditory capabilities) and Díaz et al. (2008) (i.e., uniquely linguistic ability) about the origin of individual differences in L2 learning could be due to a difference in age of acquisition of the non-native speech sounds and/or the distinct types of learning situations. Another important difference between Díaz et al. (2008) and previous training studies is the methodology employed to assess learning. Díaz et al. (2008) measured an automatic brain response during passive listening, i.e., the MMN. The use of the MMN may provide a more fine-grained measure of auditory discrimination abilities that is not influenced by differences in attention or motivation to the task.

Crucially, training studies investigated the acquisition of new phonemes in adults while Díaz et al. (2008) studied bilinguals who learned the L2 during childhood

(i.e., at 4 years of age). It has been proposed that the neural mechanism that supports language learning changes with development. According to these views, while early language acquisition is rooted in the same brain mechanisms as native language learning, later L2 acquisition recruits distinct brain regions other than those involved in native language processing (for a debate on the topic, see Birdsong, 1999). Alternatively, one could claim that formal training and more naturalistic language learning play out differently with respect to the role of non-linguistic skills. The bilinguals studied in Díaz et al. (2008) acquired their L2 in the first years of their lives and lived in a fully bilingual society. The participants in training studies are exposed to a less naturalistic learning situation, which may involve the use of different learning strategies (as proposed by Goldinger, 2007, and McClelland, McNaughton & O'Reilly, 1995). Nevertheless, the many differences between the studies make it very difficult to accurately pinpoint a single factor (or combination of factors) that may trigger the different results. Still, these different findings make one wonder whether people who learned a second language in a natural environment, but at a later age than those studied in Díaz et al. (2008), would show a relationship between their speech-specific discrimination abilities, that is, their discrimination of any speech sound (native, L2, or unknown), and their learning of L2 phonemes.

To address the question of whether the variability in the mastery of L2 sounds is explained by speech-specific or general-acoustic capabilities in bilinguals regardless of the age at which the L2 was learned, the present study assessed and analysed the MMN with similar methods and procedures as were used in Díaz et al. (2008) but in a new bilingual population (i.e., late bilinguals) and for additional types of speech changes. In the present study, the participants were late bilinguals who learned the second language in a formal setting rather than through spontaneous social interaction. In addition, the

present study investigates, not only the discrimination sensitivity of phonemes that differed in the frequency spectral cues (as was done in Díaz et al., 2008), but also of other types of phonetic features, such as duration and nasality information.

In the present study, analogous to Díaz et al. (2008), GPs and PPs of an L2 vocalic contrast were selected from a population of Dutch–English bilinguals based on their results in different behavioural tasks (see Methods; see Díaz et al., 2012, for a detailed description of the tasks). All participants were native Dutch (L1) speakers and started learning English (L2) at the age of 10–12 in formal educational settings (Table 1). Note that 'perfect' acquisition of L2 phonology is already compromised as early as the age of 4 (Caramazza, Yeni-Komshian, Zurif & Carbone, 1973; Pallier et al., 1997); hence, in terms of phonological learning, the present sample of participants can be labeled as late bilinguals. The two groups of bilinguals differed in their ability to discriminate the English vowels /æ/ (as in cattle: /kætl/) and /ɛ/ (as in kettle: /kɛtl/). Previous research has shown that it is very difficult to learn a new phonetic contrast when the native language has a single phoneme category falling approximately in between the new contrast (Best, 1995; Best & Tyler, 2007; Flege, 1995). This is, for example, the problem that Japanese listeners must face when learning the English /r-l/ contrast (Goto, 1971). In the present case, the two English mid-front unrounded vowels /æ/ and /ɛ/ are perceptually assimilated to the only available mid-front unrounded Dutch vowel /ɛ/, which is phonetically somewhere between the two English vowels, and are therefore difficult for Dutch listeners to perceive as different vowels (Broersma, 2005; Cutler, Weber, Smits & Cooper, 2004).

After participants' behavioural measurements, GPs and PPs participated in an event-related potentials (ERP) study to measure their auditory discrimination sensitivity to both general-acoustic and speech-specific contrasts by means of the MMN brain

response. To evaluate general-acoustic capabilities, the exact same procedures from Díaz et al. (2008) were used. Participants' discrimination sensitivity to duration and frequency changes were evaluated in an oddball paradigm in which a tone was presented frequently (*standard*), while three other tones deviating in the magnitude (small, medium and large deviants) of one parameter (either frequency or duration) were presented at a lower probability (*deviants*). Participants' capacity to extract patterns from an auditory signal was evaluated by presenting a sequence of two pure alternating tones differing in frequency (*standard*). The predictable presentation of the tones was sometimes violated by repeating one of the two tones (*deviant*) (Atienza, Cantero, Grau, Gomez, Dominguez-Marin & Escera, 2003). To maximally tax the auditory perceptual system and in an effort to increase the likelihood of observing differences between participants, the tones were presented at a fast rate (one tone every 314 ms). Importantly, the lack of reliable MMN signatures for the smaller acoustic changes (i.e., small deviants) in Díaz et al. (2008) in the Duration and Frequency conditions shows that our paradigm is good at examining the limits of the participants' auditory system.

For the assessment of speech-specific capabilities, the ERP paradigm was similar to that used in Díaz et al. (2008). Native and non-native phonemes were used, but the phonetic stimuli were adapted to the languages the participants knew. The relevance of using native and non-native phonemes is that participants do not vary in their age of acquisition, motivation to learn these phonemes, or amount of exposure. The present study evaluated the discrimination sensitivity of several native phonemic cues. In our previous study, only sensitivity to changes in spectral properties was evaluated because that was the only phonemic relevant cue in the languages spoken by the participants (Spanish and Catalan). In the current study, native speech

10

discrimination abilities were measured for duration and spectral cues because these two types of information are phonemically relevant to discriminate Dutch vowels (Reinisch & Sjerps, 2013) as well as the English L2 contrast (Díaz et al., 2012). For the non-native speech contrast a new phonemic cue was included, nasality (air passes through the nose when a phoneme is pronounced, such as in the consonants /m/ or /n/ or in some French vowels), a speech feature that is not a property of Dutch vowels, which are all oral (the air passes through the mouth). Participants' discrimination sensitivity to each type of phonetic feature was evaluated by presenting in an oddball paradigm vowels that differed only in the target phonetic feature (native duration: /O:/-/O/[2]; native spectrum: /A/-/O/[2]; non-native nasalization: /O~/-/O/[2]). In a separate block, the same L2 contrast employed in the behavioural tasks for the selection of the participants, i.e., the English vowels /ɛ/ and /æ/, was presented in an oddball paradigm. A diminished MMN was expected for the group of PPs compared to GPs because the MMN amplitude should relate to the overt categorization of these vowels (Näätänen et al., 1997; Winkler et al., 1999).

In the present study, the MMN amplitude and scalp distribution, useful for inferring the activity of the MMN generators, are compared for good and poor perceivers for acoustic and speech changes. Based on the hypothesis that differences in L2 phonetic processing in late bilinguals are caused by differences in speech-specific capabilities, as in early bilinguals (Díaz et al., 2012), we expect the MMN to be larger for GPs compared to PPs only when processing speech, regardless of the experience with the speech sounds presented (i.e., native, L2, or unknown vowels) and of the type of speech information manipulated (i.e., spectrum, duration, or nasality).

2. Methods

2.1. Participants

Selection of the experimental sample

An initial sample of 55 healthy Dutch (L1)–English (L2) late bilinguals was recruited from the Max Planck Institute for Psycholinguistics participant pool (42 females; mean age = 21.16, *SD* = 2.47). All participants had received on average 7 years (*SD* = 2.02) of English instruction during primary and secondary education, starting when they were 11 years-old (*SD* = 1.01). They had lived in the Netherlands all their life in a monolingual Dutch environment and were fluent speakers of English. No participant knew any French. Participants were college or graduate students and did not report having had language, hearing, or learning disabilities. They were paid for their participation.

All participants performed three behavioural tasks designed to evaluate their ability to perceive the English /ɛ/ - /æ/ vowel contrast that is very difficult for Dutch native listeners to discriminate (Broersma, 2005; Cutler et al., 2004). The tasks were a categorization task on a seven-step continuum that ranged from /æ/ to /ɛ/, a word identification task in which participants had to choose which word of an /æ/-/ɛ/minimal pair they heard, and a lexical decision task in which participants had to judge whether auditory strings containing /æ/ or /ɛ/ were actual English words (Broersma & Cutler, 2011). The categorization task measured acoustic-phonetic analysis, while the other two tasks evaluated phonological representations during lexical access (for a more detailed description of the task and the materials, see Díaz et al., 2012). In previous studies, a cutoff point was calculated for each task to establish a native-like performance range (the average minus 3 times the standard deviation of the native group). Bilinguals were categorized as good or poor perceivers if they performed consistently within or below natives' proficiency range respectively (Díaz et al., 2008; Díaz et al., 2012; Sebastián-

Gallés & Baus, 2005). However, this criterion could not be used in the present sample because no bilingual performed at native-like levels in all of the three tasks. Instead, we used hierarchical clustering and *k*-means procedure (IBM SPSS statistics 19) on the average accuracy for the three behavioral tasks to classify the bilinguals as good or poor perceivers (Archila-Suerte, Bunta & Hernandez, in press). A hierarchical cluster analysis using Ward's method of minimum variance with Euclidean square distance intervals revealed two large scale groups of participants. A clear demarcation point in the agglomeration schedule coefficient between two- and one-cluster solutions (74.24 and 281.24 respectively) indicated that a two-cluster solution is the most adequate to split the sample of participants (also confirmed by visual inspection of the dendrogram). The two cluster groups consisted of 28 and 27 participants each. In a second step, *k*-means clustering with fixed seeds (*k* = 2) was run. This analysis yielded two clusters of 28 and 27 participants each, the exact same groups as the previous hierarchical cluster analysis. Cluster 1 scored above cluster 2 in the categorization (log odds for cluster 1: 4.10 ± 1.12, cluster 2: 0.24 ± 0.83; $t(53) = 14.37$, $p < 0.001$) and the word identification tasks (log odds for cluster 1: 1.28 ± 0.73, cluster 2: 0.83 ± 0.51; $t(53) = 3.39$, $p < 0.05$) but were similar for the lexical decision task (A' for cluster 1: 0.70 ± 0.08, cluster 2: 0.70 ± 0.09; $t(53) > 1$).

Experimental groups

For the GP group, eight participants were randomly selected from among the 28 participants in cluster 1. They all scored within the native performance range – 3 standard deviations below the natives' mean (see Díaz et al., 2012) – in the categorization task. In addition, two of the participants scored within the native range of performance in the lexical decision task and another one in the word identification task.

For the PP group, another eight participants were randomly selected from among the 27 participants in cluster 2. They all scored below 3 standard deviations below natives' mean in all three behavioural tasks except for one (who achieved native-like levels in the lexical decision task). Table 1 displays the details of the participants from the two experimental groups and their accuracy in each behavioural task. All participants were right-handed, as assessed by the Edinburgh Handedness Inventory (Oldfield, 1971), and gave written informed consent to participate in the study. The experiment was approved by the local research ethics panel.

Table 1. *Groups' biographic details and behavioural performance in L2 phonetic tasks.* Standard errors in parenthesis.

	Good perceivers (n=8)	Poor perceivers (n=8)	*t*-test (df=14)
Biographic details			
Age	21.12 (2.74)	20.12 (1.55)	< 1
Gender	7 females	6 females	
English AoA	11.12 (1.24)	11.00 (1.06)	< 1
Years English education	7.37 (1.50)	7.50 (.92)	< 1
L2 tasks performance			
Categorization (log odds)	5.12 (.95)	0.41 (.43)	12.67**
Word identification (log odds)	1.66 (1.00)	0.68 (.33)	2.60*
Lexical decision (A')	0.73 (.07)	0.68 (.08)	1.15

*Note: *p* < .05, **p* < .001, AOA = Age of Acquisition

2.2. Stimuli

Tonal and speech stimuli were employed. The tonal stimuli were four 1000 Hz pure tones of different durations: 200 ms, 120 ms, 80 ms, 40 ms (Duration condition); four 50 ms pure tones that varied in frequency: 1000 Hz, 1030 Hz, 1060 Hz, 1090 Hz (Frequency condition); and two 50 ms pure tones with a frequency of either 500 or 1000 Hz (Pattern condition) (Table 2). All tones had 10 ms of rise/fall times.

Table 2. *Experimental design and stimuli.* Relevant features of standard (italic type) and deviant (bold) stimuli are listed.

Condition	standard	deviant
ACOUSTIC		
Duration	*200 ms*	**120 ms, 80 ms, 40 ms**
Frequency	*1000 Hz*	**1030 Hz, 1060 Hz, 1090 Hz**
Pattern	*AB*	***AA* *B*B**
SPEECH		
Speech-1	/O/	**/O:/** (native dur.)**,** **/A/** (native freq.)**,** **/O~/** (un-known nasal)
Speech-2	/ɛ/	**/{/** (L2)

The speech stimuli were Dutch (participants' native language), English (participants' L2), and French (unknown for participants) vowels (Speech-1 condition). The Dutch and French vowels were synthesized with Klatt synthesizer (Klatt, 1980). The Dutch vowel /O/ was 120 ms long and the formant frequencies were $F1$ = 500 Hz, $F2$ = 890 Hz, $F3$ = 2600 Hz, and $F4$ = 3500 Hz (Rietveld & van Heuven, 1997). Its pitch contour linearly fell from 120 to 105 Hz over the duration of the sound, and the nasality parameter was set to 0 to create an oral vowel. The Dutch vowel /O:/ was synthesised with the exact same parameters as the vowel /O/ but with a longer duration of 180 ms (Rietveld & van Heuven, 1997). The native Dutch vowel /A/ was synthesised with the exact same parameters as the vowel /O/ except for the $F1$ and $F2$ formant frequencies (680 and 1050 Hz respectively) (Rietveld & van Heuven, 1997). The non-native French nasal /O~/ was synthesised with the exact same parameters as the vowel /O/ but differed only in the nasality parameter (which was set to 350 Hz).

In addition, participants were presented with the two endpoints of the seven-step continuum used in the categorization task for participant selection, the English vowels /ɛ/ and /æ/ (Speech-2 condition). They were synthesised with the source-filter synthesis

of the PRAAT software (Boersma, 2001). Both vowels were generated based on the average values from four English speakers (/ɛ/: $F1 = 600$ Hz, $F2 = 1800$ Hz; /æ/: $F1 = 740$ Hz, $F2 = 1630$ Hz; the other formants were identical: $F3 = 2750$ Hz, $F4 = 3400$ Hz, and $F5 = 4500$ Hz). The pitch contour for both vowels fell linearly from 108 to 90 Hz. The duration of both synthetic stimuli was 165 ms.

2.3. Procedure

The participants' central sound processing was evaluated for three different general-acoustic features (Duration, Frequency, and Pattern conditions) and for native and non-native phonetic stimuli (Speech-1 and Speech-2 conditions) (see Table 2) (Díaz et al., 2008; Corbera et al., 2005). Each condition was presented in separate blocks.

In the Duration and Frequency conditions, the tones were presented in an oddball paradigm in which the standard tone was presented with a probability of .8 (600 standard tones per block), and the probability of each deviant (small, medium and large) was .066 (50 presentations of each deviant tone per block). In the Frequency condition, a 1000 Hz tone was the standard, and a 1030 Hz (small deviant), a 1060 Hz (medium deviant), and a 1090 Hz (large deviant) tones were the deviants. In the Duration condition, the standard tone had a length of 200 ms, while the deviants had a length of 120 ms (small deviant), 80 ms (medium deviant), and 40 ms (large deviant). The stimulus onset asynchrony (SOA) was 314 ms.

In the Pattern condition, 400 trains of six tones were presented (2400 tones all together). Each train was created by presenting alternately two tones with a frequency of 1000 and 500 Hz each. Tones within and between the trains were presented at a constant SOA of 128 ms (onset-to-onset inter-train interval: 768 ms). Stimulus trains were presented in a predictable way: 'ABABAB-**B**ABABA-BABABA-**A**BABAB…' in

16

which 'A' represents the 500 Hz tone, 'B' the 1000 Hz tone, '-' indicates the beginning of the trains, and '**A**' and '**B**' denote the deviant event, i.e., repetition of the last tone presented in the preceding train.

Two blocks assessed phoneme discrimination capabilities. In the Speech-1 block, the standard stimulus was the Dutch vowel /O/ and the deviant stimulus were the Dutch vowel /O:/ (native duration deviant), the Dutch vowel /A/ (native spectrum deviant), and the French vowel /O~/ (unknown nasal deviant). The presentation probability of the standard vowel was .8 (600 standard presentations per block), and each deviant probability was .066 (50 presentations of each deviant tone per block). In the Speech-2 block, the standard stimulus was the English vowel /ɛ/ and had a presentation probability of .8 (400 standard presentations), while the deviant stimulus was the English vowel /æ/ and had a presentation probability of .2 (100 presentations). Vowels were presented with a constant SOA of 488 ms.

For all oddball conditions except for Pattern, stimuli were presented in a random order with the restriction that the first five stimulus of the blocks were always a standard and that at least one standard was presented between two deviants.

Eight presentation lists were created, including two presentations of each condition to have enough trials for each of the deviants (100 deviant trials) except for the Speech-2 condition which was presented only once because one block already contained 100 deviant trials. First, the three acoustic conditions and the Speech-1 condition appeared in random order, and after a short break, they were repeated in the reverse order. The Speech-2 block was assigned randomly to one position. Lists were counterbalanced between groups.

ERP measurements took place in an electrically shielded, soundproof room at the Donders Centre for Cognitive Neuroimaging in Nijmegen, the Netherlands. Testing

17

took place in one session several weeks after the behavioural tests. During the EEG recording, participants sat in a comfortable chair in a dimly illuminated, sound-attenuating booth. Participants were instructed to ignore the auditory stimulation and to watch a silent movie. All the stimuli were delivered binaurally via a loudspeaker set, placed approximately 1.5 m in front of the participants, at an intensity of 80 dB. The experimental session lasted about one hour, including a ten-minute break.

2.4. Electrophysiological recording

The ERPs were recorded from the scalp using a 32-channel BrainCap with sintered Ag/AgCl electrodes and two additional electrodes located at the two mastoids (LM, RM). Eye movements were measured with electrodes attached to the infra-orbital ridge of the right eye and on the outer canthus of the right and left eyes. The common EEG/EOG reference was attached to the tip of the nose. Electrode impedances were kept below 5 kOhm. The electrophysiological signals were filtered online with a bandpass of 0.1–100 Hz and digitized at a rate of 500 Hz.

The EEG was offline filtered with a bandpass of 0.1–30 Hz and a slope of 12 dB/oct). ERPs were averaged offline for standard and deviant stimuli, separately for each participant and condition. Automatic ocular correction was performed using the method of Gratton and Coles (BrainVision Analyzer Software package v. 1.05; Brain Products GmbH, Munich, Germany). Epochs with EEG exceeding either ±100 μV after baseline subtraction at any channel, activity lower than 0.5 μV, or more than 50 μV voltage step/sampling within intervals of 200 ms were automatically rejected offline. Standard stimulus epochs occurring immediately after deviant stimulus epochs were excluded from the analysis. For each deviant at least 55 trials were accepted. Epochs included a pre-stimulus baseline of 100 ms in all cases and were 500 ms long. Baseline

was corrected, and lineal DC detrend procedure was performed on the individual segments.

2.5. Data analysis

The MMN was identified in the difference waves (obtained by subtracting ERPs elicited by the standards from those elicited by the deviants) at the Fz electrode in a 100–300 ms time window after stimulus onset for each group. For most comparisons, the MMN peaked at the same latency for both groups, except for two speech comparisons (the native frequency comparison /A/ - /O/ and the L2 comparison /æ/- /ɛ/) for which the negativity slope started at the same time point for both groups but peaked earlier for the PP group (Figure 3). For the statistical analysis, the MMN was measured for each participant group and condition as the mean amplitude in a 40 ms latency window centered in its group maximum peak (Table 3). To test whether a significant MMN was elicited in each group and condition, one sample *t*-tests were carried out to compare the amplitudes of the MMN component at the Fz electrode against the zero level (Table 3). In addition, the number of trials accepted for each participant and deviant were submitted to *t*-tests comparisons to ensure that a similar number of trials was averaged for each group of participants. The *t*-test did not reveal significant differences in the number of trial averaged for good and poor perceivers for any of the deviants (for all *t*-tests p > .1).

Table 3. *T-test of the MMN mean amplitude for the acoustic (D=Duration, F=Frequency, P=Pattern) and speech conditions at the Fz electrode.* Degrees of freedom in subscripts.

	Latency window (ms)	Poor perceivers		Good perceivers	
		Amplitude (μV)	$t_{(7)}$	Amplitude (μV)	$t_{(7)}$
D: 120 ms – 200 ms	260-300	-0.80 (0.78)	2.87*	-0.82 (1.00)	2.32[+]
D: 80 ms – 200 ms	210-250	-0.78 (0.76)	2.93*	-1.11 (1.28)	2.46*
D: 40 ms – 200 ms	155-195	-1.73 (0.83)	5.84***	-2.04 (1.31)	4.40*
F: 1,030 Hz – 1000 Hz	180-220	-0.20 (0.86)	< 1	-0.51(0.74)	1.95
F: 1,060 Hz – 1000 Hz	150-190	-0.14 (0.88)	< 1	-1.15 (1.68)	1.94
F: 1,090 Hz – 1000 Hz	150-190	-0.54 (1.36)	1.03	-0.81 (0.74)	3.08*
Pattern: AA/BB – AB/BA	110-150	-0.96 (0.53)	5.06***	-1.04 (0.64)	4.56*
Sp-1 dur.: /O:/-/O/	215-255	-0.87 (0.69)	3.58*	-1.07 (1.00)	3.01*
Sp-1 spect.: /A/-/O/	PP:90-130//GP:180-220	-0.51 (0.66)	2.21[+]	-0.97 (0.93)	2.94*
Sp-1 unknown: /O~/-/O/	180-220	-1.33 (1.15)	3.27*	-2.49 (1.29)	5.43***
Sp-2 L2: /{/-/ɛ/	PP:120-160//GP:150-190	-0.44 (1.49)	< 1	-1.71 (0.75)	6.43***

Note $^{+}p < 0.065$, $^{*}p < 0.05$, $^{***}p < 0.001$

A 5-way (2x2x2x2x3), repeated-measures ANOVA compared the MMN amplitudes for the two groups (factor 'Group') in the Tone and Speech condition types (factor 'Condition type'). To test also whether the groups differed in the scalp distribution of the MMN, the factors 'Laterality' (left electrodes: C3 and F3, and right electrodes: C4 and F4) and 'Frontality' (frontal electrodes: F3 and F4, and central electrodes: C3 and C4) were included in the ANOVA. The factor 'Deviant feature' was included nested within the factor 'Condition type'. For the Tone condition type, there were 3 deviant features: duration (the mean MMN amplitude of the three duration deviants), frequency (the mean MMN amplitude of the three frequency deviants), and pattern. For the Speech condition type, there were also 3 deviant features: duration (deviant /O:/), spectral (deviant /A/), and nasal (deviant /O~/). The L2 deviant /{/ was not included in the analysis because it was expected to elicit a larger MMN in the GP group than in the PP groups according to the behavioural tasks.

Further repeated-measures ANOVAs were calculated for each condition separately (Frequency, Duration, Pattern, Speech-1, and Speech-2) with the factors 'Laterality', 'Frontality', 'Deviant', and 'Group'.

When differences between the groups were found (i.e., significant main effect or interaction involving the factor 'Group'), a second ANOVA was performed with the additional factor 'MMN generator' (frontal: F3, F4 vs. mastoids: LM, RM) to investigate the contributions of the temporal and frontal MMN sources to individual differences in phoneme perception.

No correction for multiple comparisons was applied to the present planned comparisons because we had strong hypotheses based on our previous studies (Díaz et al., 2008; Sebastián-Gallés et al., 2012): we expected GPs and PPs to have similar MMNs for the tone conditions and GPs to have larger MMNs for speech sounds at frontal, but not mastoid, electrodes. All main effects are reported. However, only interactions involving the factor 'Group' are reported. Significance levels of the F-ratios were adjusted with the Greenhouse–Geisser correction for effects with more than 1 degree of freedom in the numerator, and the corrected p values are reported.

3. Results

Table 4 displays the effects that reached significance in the ANOVAs. The ANOVA comparing the GPs and PPs for all conditions together showed significant differences between the two groups ($F_{(1,14)} = 6.66$, $p < .05$). The main effect of 'Condition type' ($F_{(1,14)} = 22.58$, $p < .001$) was also significant. Crucially, the interaction 'Group' x 'Condition type' reached significance ($F_{(1,14)} = 5.91$, $p < .05$) (Figure 1). No other main effect or interactions with the factor 'Group' were significant. To investigate in which

conditions GPs and PPs differed, separate ANOVAs were performed for the acoustic and speech conditions.

Table 4. Significant effects yielded by the ANOVAs comparing the groups for all the conditions together and each condition separately.

| ANOVAs	Effect	df	F	p
Tone vs. Speech	Condition	1,14	22.58	< 0.001
	Group	1,14	6.66	< 0.05
	Group x Condition	1,14	5.91	< 0.05
Duration	---			
Frequency	Group x Frontality	1,14	4.76	< 0.05
	Group x Frontality x Deviant	1,14	3.47	= 0.05
Pattern	Group x Frontality	1,14	4.88	< 0.05
Speech-1	Group	1,14	8.13	< 0.05
	Group x Laterality x Deviant	2,14	4.81	< 0.05
	Group x MMN generator	1,14	4.36	= 0.056
Speech-2	Group	1,14	5.24	< 0.05
	Group x MMN generator	1,14	3.46	= 0.084

3.1. Acoustic conditions

MMN elicited in the Duration condition.

Both groups displayed a reliable MMN in response to the large (40 ms) and medium (80 ms) deviants compared to the standard, which had a duration of 200 ms (Table 3). The MMN elicited by the small deviant (120 ms) was significant for the PP group and marginally significant for the GP group.

When the MMN amplitudes were submitted to an ANOVA, the factor 'Group' did not reach significance ($F_{(1,14)} < 1$) and did not interact with any other factor (all $p >$ .05) (Figure 2). A main effect of 'Deviant' was observed ($F_{(2,28)} = 6.85$; $p < .05$). The large deviant elicited a more prominent MMN than the medium ($t_{(15)} = 2.94$, $p < .05$) and the small deviants ($t_{(15)} = 4.19$, $p = .001$) (amplitude means: deviant 120 ms = -.56 μV, deviant 80 ms = -.81 μV, deviant 40 ms = -1.57 μV). No differences were present between the MMNs elicited by the medium and the small deviant ($t_{(15)} < 1$).

<Insert Figure 2 about here>
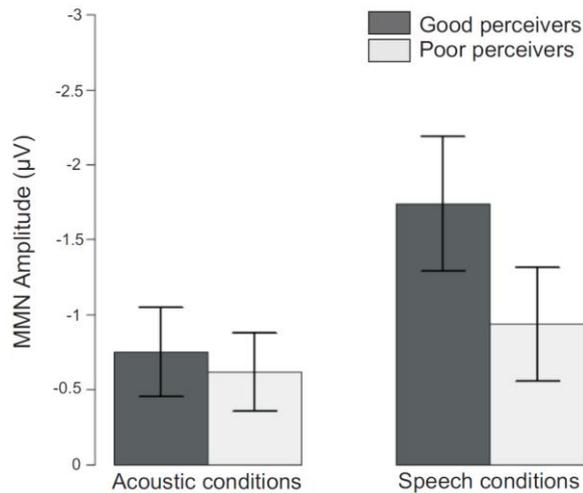
Figure 1. *Mean MMN amplitudes for the acoustic and speech conditions.* Bars represent the mean MMN amplitudes for the two types of conditions (acoustic and speech) for good and poor perceivers. Error bars represent standard deviations.

MMN elicited in the Frequency condition.

The large deviant stimuli (1090 Hz, standard: 1000 Hz) elicited a reliable MMN for the GP group. The other negativities assessed did not reach significance (Table 3). Note that despite the rather large amplitude (-1.15 µV), the negativity for the middle deviant in the GP group was not different from the zero level because of the large variability across participants (the standard deviation was ±1.68 µV and larger than the mean amplitude difference).

The ANOVA did not reveal group differences ($F_{(1,14)}$ = 2.44, p = .140). The factor 'Group' interacted significantly with 'Frontality' ($F_{(1,14)}$ = 4.76, $p < .05$) and the triple interaction 'Group' x 'Frontality' x 'Deviant type' was also significant ($F_{(1,14)}$ = 3.47, p = .05). To explore whether group differences caused the triple interaction, t-test comparisons between the two groups were performed separately for each deviant type at each frontal and central electrodes. None of the t-test yielded significant results (all $p > 0.1$).
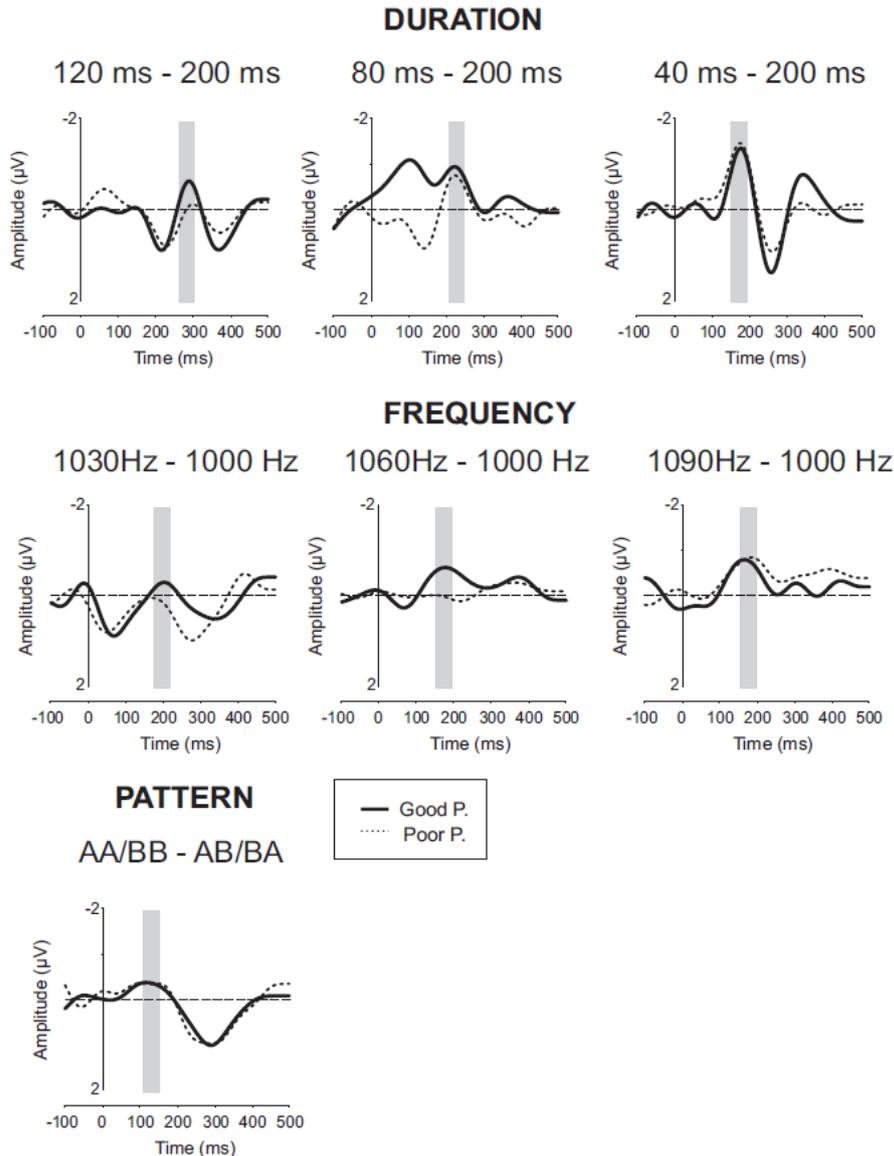
MMN elicited in the Pattern condition.

Both groups showed a reliable MMN response (Table 3). The statistical comparisons revealed that the negativity was not different between the two groups ($F_{(1,14)} < 1$) (Figure 2). The factor 'Frontality' was significant ($F_{(1,14)} = 11.65$, $p < .05$) due to the enhancement of the negativity at frontal electrodes (frontal = -.84 µV, central = -.40 µV). The factor 'Frontality' interacted significantly with the factor 'Group' ($F_{(1,14)} = 4.88$, $p < .05$). The subsequent *t*-test comparisons of the MMN amplitude at each electrode did not reveal any significant difference between the two groups (all p >.05).

3.2. Speech conditions

MMN elicited in the Speech-1 condition

The native duration deviant phoneme /O:/ (different from the standard phoneme only in duration) elicited a reliable MMN in both groups of participants; the native spectrum deviant phoneme /A/ (different from the standard phoneme only in the spectrum) elicited a reliable MMN in the GP group and a marginally significant MMN in the PP group. The non-native deviant, the unknown French nasal vowel /O~/ (different from the standard phoneme only in nasalization) triggered a reliable MMN in both groups.

The analysis revealed a significant main effect of 'Group': GPs showed larger MMNs for phonetic stimuli than PPs ($F_{(1,14)} = 8.13$, $p < .05$; GP = -1.73 µV, PP = -.93 µV) (Figure 3). This effect was not qualified by a 'Group' x 'Deviant' interaction ($F_{(1,14)} < 1$), meaning that the PP group showed smaller mismatch detection for all kinds of deviants (spectrum, duration, and nasality). In addition, there was a main effect of the

24

DURATION

120 ms - 200 ms

80 ms - 200 ms

40 ms - 200 ms

FREQUENCY

1030Hz - 1000 Hz

1060Hz - 1000 Hz

1090Hz - 1000 Hz

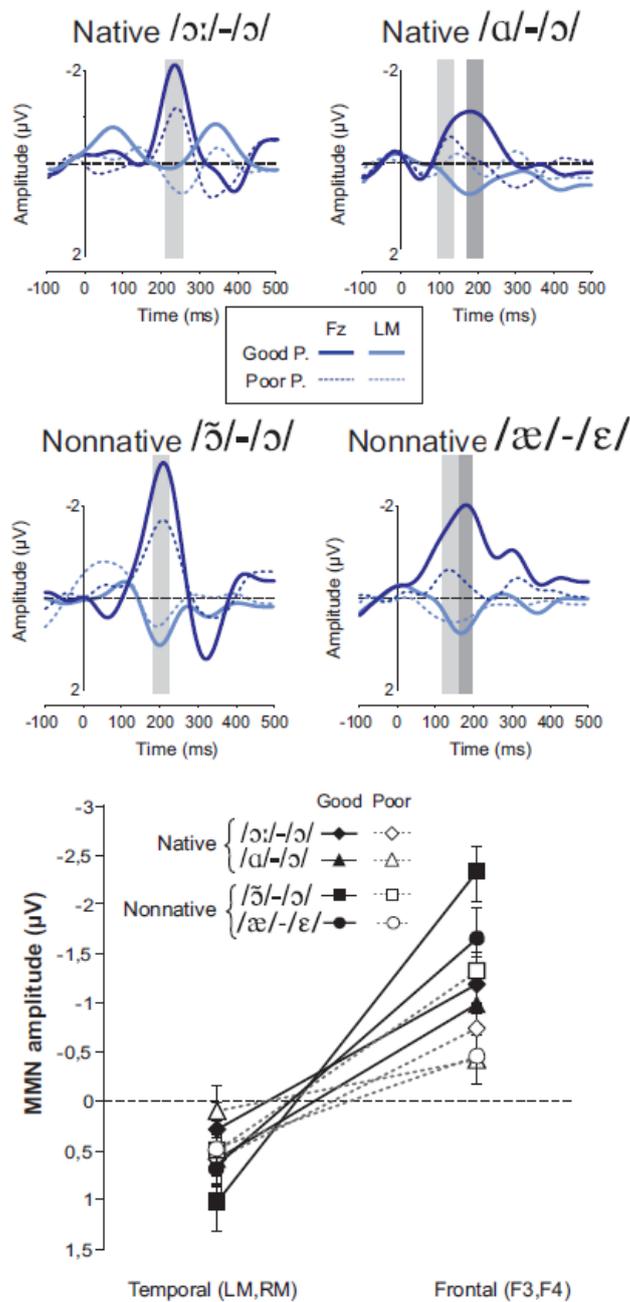## PATTERN

AA/BB - AB/BA

— Good P.
...... Poor P.

Figure 2. *MMNs for the acoustic conditions.* The grand-mean difference waves (responses elicited by the standard stimuli subtracted from that elicited by the deviant stimuli) are displayed for the good and poor perceivers in the acoustic conditions (duration, frequency and pattern conditions) at the C3 electrode. Grey boxes indicate latency windows of the MMNs. For visualization purposes, this data is displayed with a low-pass filter of 8Hz.

factor 'Deviant' ($F_{(2,28)}$ = 6.93, p < .05). *T*-test comparisons showed that the negativity elicited by the deviant non-native phoneme /O~/ was larger from the ones elicited by each native deviant vowel (/O~/ vs. /O:/: $t_{(15)}$ = 3.53, p < .05; /O~/ vs. /A/: $t_{(15)}$ = 2.86, p < .05). The MMN to the native deviant Dutch phonemes /O:/ and /A/ were not different ($t_{(15)}$ < 1) (MMN mean amplitudes: /O:/ = -.97 µV, /A/ = -.74 µV, /O~/ = -

1.91 µV). The factor 'Frontality' also reached significance ($F_{(2,28)}$ = 10.08, p < .05) revealing a central scalp prominence of the MMN (frontal = -1.15 µV, central = -1.51 µV).

The triple interaction 'Deviant' x 'Laterality' x 'Group' was also significant ($F_{(2,28)}$ = 4.81, p < .05). *T*-test comparisons between the two groups were performed separately for each deviant and electrode. GPs had larger MMNs than PPs at right electrodes for the native deviant /A/ (at F4: $t_{(14)}$ = 2.32, p < .05, GP: -1.10 µV, PP: -0.25 µV) and for the non-native deviant /O~/ (at F4: $t_{(14)}$ = 2.17, p < .05, GP: -2.53 µV, PP: -1.29 µV; at C4: $t_{(14)}$ = 2.23, p < .05, GP: -2.94 µV, PP: -1.34 µV). A further ANOVA was carried out to investigate the role of the frontal and temporal MMN generators in the group differences (Figure 3). There was a main effect of the factor 'Generator' ($F_{(1,14)}$ = 59.82, p < .001) due to the reversal of the frontal negativity at the mastoid electrodes (frontal = -1.15 µV, mastoids = .51 µV). The factor 'Group' did not reach significance ($F_{(1,14)}$ = 2.22, p > .05) but the interaction 'Group' x 'MMN generator' was marginally significant ($F_{(1,14)}$ = 4.36, p = .056). Subsequent analyses showed significant differences between the two groups at the frontal electrodes ($t_{(14)}$ = 2.36, p < .05; GP = -1.49 µV, PP = -.82 µV) but not at the temporal ones ($t_{(14)}$ < 1; GP = .63 µV, PP = .40 µV). That is, the amplitude of the MMN was larger for GPs than for PPs only at the frontal electrodes. As in the previous ANOVA, the triple interaction 'Deviant' x 'Laterality' x 'Group' reached significance ($F_{(2,28)}$ = 3.98, p < .05). Further *t*-test comparisons between the groups for each deviant and electrode separately showed, as in the previous ANOVA, a larger MMN for GPs at the F4 electrode for the native deviant /A/ and the unknown deviant /O~/.

Figure 3. *MMNs for the speech conditions.* The grand-mean difference waves (responses elicited by the standard stimuli subtracted from that elicited by the deviant stimuli) are displayed for the good and poor perceivers in the speech conditions (speech-1 and speech-2 conditions) at the C3 electrode. Grey boxes indicate latency windows of the MMNs. When the latency of the MMN peak maxima is different between the groups, MMN latency windows for good perceivers are in dark grey boxes and the ones for poor perceivers are in light grey boxes (upper panel). For visualization purposes, this data is displayed with a low-pass filter of 8Hz. Mean amplitude values for both groups at frontal (F3, F4) and temporal (LM, RM) electrodes for all deviants in the speech conditions. Error bars depict standard errors (bottom panel).

MMN elicited in the Speech-2 condition

The deviant L2 phoneme /æ/ elicited a reliable MMN only for the GP group (see Table 3). The ANOVA revealed a main effect of 'Group' ($F_{(1,14)} = 5.24$, $p < .05$) with a larger MMN in the GP group than in the PP group (GP = -1.61 µV, PP = -.47 µV) (Figure 3). This result is in line with the behavioral differences in the discrimination abilities of the two groups for the L2 contrast /æ/ - /ε/, on the basis of which these groups were formed. The main effect of 'Laterality' was also significant ($F_{(1,14)} = 13.29$, $p < .05$) revealing a larger MMN at left than at right-hemisphere electrodes (right = -.89 µV, left = -1.18 µV).

An ANOVA comprising the factors 'MMN generator', 'Laterality', and 'Group' again revealed a generator effect ($F_{(1,14)} = 19.29$, $p < .001$) caused by the change in polarity at temporal sites (frontal = -1.05 µV, mastoids = .59 µV) (Figure 3). The interaction between the factors 'Group' and 'MMN generator' was marginally significant ($F_{(1,14)} = 3.46$, $p = .084$). Further analyses showed that the MMN was larger for the GP than the PP group at the frontal generator ($t_{(14)} = 2.49$, $p < .05$; GP = -1.65 µV, PP = -.45 µV), while no differences were observed at the temporal one between the two groups ($t_{(14)} < 1$; GP = .69 µV, PP = .49 µV).

4. Discussion

The present study compared auditory discrimination sensitivity of speech and non-speech sounds by good (GPs) and poor (PPs) perceivers of an L2 speech contrast, as indexed by the event-related potential MMN. Comparable MMN responses were elicited by the two groups of participants in three acoustic conditions (involving tones varying in frequency, duration, and presentation pattern) suggesting that the groups did not differ in the processing of non-speech stimuli. The lack of reliable MMN signatures

for deviant stimuli with a small magnitude in both the Duration and the Frequency conditions shows that our paradigm was good at examining the limits of the participants' auditory system, because both rougher and finer discriminatory abilities were evaluated (similar results were found previously for early bilinguals with the same experimental design, Díaz et al., 2008).

The two groups differed in their MMN responses to phonetic stimuli. GPs exhibited larger MMNs than PPs during the processing of phonetic changes, indicating greater discrimination sensitivity of speech sounds in GPs as compared to PPs. Crucially, the distinct sensitivity of the two groups was present no matter what type of speech information changed (i.e., duration, spectrum, and nasality) and their previous experience with the phonemes (i.e., native, L2, and unknown). This result suggests that all speech sounds, regardless of their familiarity, are processed by the same cognitive mechanism and that variations in the capability of this mechanism originate individual differences in the learning of second languages. The present findings agree with previous studies showing that individual variability between L2 phoneme learners is present in the perception of speech sounds but not of pure tones (Díaz et al., 2008; Golestani & Zatorre, 2009). An open question is whether the individual differences in speech discrimination are driven by the linguistic status of the speech sounds or the acoustic complexity of speech. To the best of our knowledge, this question has yet to be investigated.

When comparing the scalp distribution of the group differences to the speech sounds, a marginally significant interaction was found between the scalp distribution of the MMN and the groups. Posterior t-test showed that GPs had larger MMN at frontal electrodes as compared to PPs, whereas the MMN was similar between the groups at the mastoids. The frontal scalp distribution of the group differences is analogous to the

findings of Díaz et al. (2008). The present findings are in line with those of a previous magnetic resonance imaging study in which GPs and PPs early bilinguals were found to have different white matter volumes at frontal, but not temporal regions (Sebastián-Gallés et al., 2012). Sebastián-Gallés et al. (2012) found larger white matter volume in the right insulo/fronto-opercular region for PPs, as compared to GPs, and, importantly, the white matter volume in this frontal region was correlated with the MMN amplitude (the less negative MMN, the more white matter volume). These convergent results suggest differences between GPs and PPs in frontal MMN generators. Distinct functional contributions have been attributed to each MMN generator (Deouell, 2007; Shalgi & Deouell, 2007). The temporal generator is claimed to be related to the comparison of the incoming auditory sensory input and a memory trace. Although one cannot accurately identify MMN generators with ERPs, several studies support that the activity of the MMN generators can be inferred from the amplitude, latency and scalp distribution of the MMN (Escera et al., 1998; Giard et al., 1990; Näätänen, 1990; Yago et al., 2001). Therefore, the absence of differences in the activity generated at mastoids could be interpreted as both groups being equally able to represent and integrate the incoming phonemic auditory information. The frontal generator is claimed to sustain the re-orienting of attention to changes in the auditory signal. Hence, the differences observed at frontal electrodes suggest that the two groups may differ in the way the disparity between an incoming mismatching phoneme and the standard phoneme neural representation triggers involuntary attention. Following this rationale, GPs and PPs seem to differ in the domain general, cognitive mechanism that allows them to reallocate attentional resources to novel and rare speech sounds. This result implies that the learning of new phonemes requires not only a sensory distinction between

phonemes, but also attentional processes to successfully represent the new sounds in long-term storage.

The present results are also relevant for the issue of age differences in language learning. It has been proposed that the neural mechanism that supports speech learning changes with development. The fact that early exposure to an L2 has a beneficial impact on the mastery of second languages has led to the claim that distinct brain learning mechanisms sustain language learning at different ages (Johnson & Newport, 1989; Lenneberg, 1967; Oyama, 1976; Patkowski, 1980; Pulvermüller & Schumann, 1994; Weber-Fox & Neville, 1996). Nevertheless, recent neuroimaging data indicate that there is a common brain network for language processing, regardless of the age of initial learning, and that the age of initial learning of the L2 had only a modulatory effect on the strength of the brain activations (Perani et al., 1998; for a meta-analysis see Indefrey, 2006; for a review see Perani & Abutalebi, 2005; for opposite findings see Kim, Relkin, Lee & Hirsch, 1997; Wartenburger, Heekeren, Abutalebi, Cappa, Villringer & Perani, 2003). The present results are in line with the conclusion that shared neural mechanisms are involved in the processing of languages, regardless of the age of initial exposure to an L2 (see also Sebastián-Gallés & Díaz, 2012). The present study on late bilinguals replicates previous findings in early bilinguals (Díaz et al., 2008): namely, GPs and PPs differed in their discrimination accuracy of all phonemes – L1, L2, and unknown – but not in their sensitivity to general-acoustic changes. The similarity in the results on early and late bilinguals[3] suggests that the perception and processing of native and L2 phonemes (and even of unknown phonemes) are sustained by the same brain mechanism, regardless of the age at which a language is learned or the previous experience with the language.

An important difference between the population of early bilinguals studied previously (Díaz et al., 2008) and the present study is the distribution of the participants' accuracy across the behavioral tasks used to evaluate the mastery of an L2 vowel contrast. In both studies bilingual participants performed three tasks that aimed to evaluate their mastery of a difficult L2 vowel contrast across several phonological processes: from acoustic-phonetic analysis (categorizing isolated vowels) to lexical access (detecting vowel mispronunciations within words). In the Díaz et al. (2008) study, the GPs performed similar to a group of native listeners in all three tasks, while the PPs performed below the native listeners' range in all three tasks. Therefore, the two groups represented the extreme endpoints of non-native phoneme perception. In the present study, the difference between GPs and PPs was not so sharp: that is, participants with native-like accuracy in one task sometimes had low scores in another task. Because of the less clear division between skilled and less skilled listeners, they were classified on their scores in the categorization task, which assessed the most basic phonological processes (i.e., acoustic-phonetic analysis, Díaz et al., 2012). Importantly, despite the different selection criteria between the present study and the Díaz et al. (2008), the MMN assessed in the present study in response to speech was again different between the two groups of late bilinguals.

The present results also show that the MMN is a reliable measure to capture group differences in speech perception. Firstly, the MMN relates to behavioural discrimination of difficult L2 phoneme contrasts (Amenedo & Escera, 2000; Näätänen, 2001). Accordingly, in the present study, only the GPs showed a reliable MMN in response to the deviant phoneme /æ/. This result reinforces the validity of the criteria established for selecting the participants. Secondly, the agreement between the behavioural and MMN results for the L2 contrast speaks for the extrapolation of the

MMN results with synthetic, isolated phonemes to natural speech. GPs were more accurate than PPs in discriminating the L2 phonemic contrast in the word identification task for which sentences recorded by a native speaker were presented. In line with this finding, GPs displayed a larger MMN to isolated, synthetic L2 vowels than PPs. Thirdly, the MMN allows the measurement of subtle differences in native perception in healthy individuals that, otherwise, are highly difficult to assess with behavioural studies because of the robustness of native processing.

This is the first study that assesses the MMN elicited by a nasal vowel in non-native listeners. Nasal vowels are easy to discriminate from oral vowels (Hawkins & Stevens, 1985) and should elicit a MMN in non-native listeners (Nenonen et al., 2005). Unexpectedly, the MMN to the non-native nasal deviant was significantly larger than to the other speech deviants suggesting that a change in nasality is much more salient than a change in formant frequency or duration (perhaps because it is acoustically a more prominent contrast than the other ones or perhaps because non-native listeners interpret the nasal vowels as an oral vowel followed by a nasal consonant as /n/ or /m/; Lahiri & Marslen-Wilson, 1991).

Whether the findings of the present study can be generalized to phoneme training studies is an open question. In contrast to the present findings, training studies usually attribute variability in phoneme learning to general-acoustic capabilities (Golestani et al., 2007; Lengeris & Hazan, 2010; Wong et al., 2007; Wong et al., 2008; for speech-specific origin of learning success in a training study, see Golestani & Zatorre, 2009). A plausible reason for the opposite results between this and training studies is that natural and training learning experiences trigger distinct types of learning strategies and processes (Goldinger, 2007; McClelland et al., 1995). For example, differential activation of striatal regions has been shown for training involving mere

passive observation compared with feedback of a non-native contrasts (the English /r-l/ contrast in Japanese listeners) (Tricomi, Delgado, McCandliss, McClelland & Fiez, 2006). The many differences between the populations studied and the distinct procedures to assess the relation between individual variability in language learning and in auditory perception make it highly difficult to discern the cause of the distinct results found by natural and training learning studies.

The present results suggest that the mastery of the L2 sounds relates to both native and unknown phoneme discrimination abilities for several types of speech information in late bilinguals. Because similar results were reported for early bilinguals (Díaz et al., 2008), we conclude that variability in the learning of non-native speech-sounds stems from variability in a uniquely speech-specific capability, regardless of the age of onset of L2 learning, and a continuity of the mechanism for L2 learning through lifespan. Hence, the assessment of speech discrimination capabilities of speech sounds, regardless of the familiarity, is a general index of linguistic abilities and can predict the successful learning of non-native phonemes.

References

Amenedo, E., & Escera, C. (2000). The accuracy of sound duration representation in the human brain determines the accuracy of behavioural perception. *The European Journal of Neuroscience, 12*, 2570–2574.

Archila-Suerte, P., Bunta, F., & Hernandez, A. E. (in press). Speech sound learning depends on individuals' ability, not just experience. *International Journal of Bilingualism.*

Atienza, M., Cantero, J. L., Grau, C., Gomez, C., Dominguez-Marin, E., & Escera, C. (2003). Effects of temporal encoding on auditory object formation: A mismatch negativity study. *Cognitive Brain Research, 16*, 359–371.

Best, C. T. (1995). A direct realist view of cross-language speech perception. In W. Strange (ed.), *Speech perception and linguistic experience: Issues in cross-language research,* pp. 171–204. Baltimore: York Press.

Best, C. T., & Tyler, M. D. (2007). Non-native and second-language speech perception: Commonalities and complementarities. In O. S. Bohn, & M. J. Munro (eds.), *Language experience in second language speech learning: In honor of James Emil Flege,* pp. 13–34. Amsterdam: John Benjamins.

Birdsong, D. (1999). *Second language acquisition and the critical period hypothesis.* Mahwah, NJ: Lawrence Erlbaum.

Boersma, P. (2001). Praat, a system for doing phonetics by computer. *Glot International, 5*, 341–345.

Bongaerts, T. (1999). Ultimate attainment in foreign language pronunciation: The case of very advanced late foreign language learners. In D. Birdsong (ed.), *Second language acquisition and the critical period hypothesis*, pp. 133–159. Mahwah, NJ: Erlbaum.

Broersma, M. (2005). Perception of familiar contrasts in unfamiliar positions. *Journal of the Acoustical Society of America, 117*, 3890–3901.

Broersma, M. (2012). Increased lexical activation and reduced competition in second-language listening. *Language and Cognitive Processes, 27*, 1205-1224.

Broersma, M., & Cutler, A. (2008). Phantom word activation in L2. *System: An International Journal of Educational Technology and Applied Linguistics, 36*, 22-34.

Broersma, M., & Cutler, A. (2011). Competition dynamics of second-language listening. *Quarterly Journal of Experimental Psychology, 64*, 74–95.

Caramazza, A., Yeni-Komshian, G., Zurif, E., & Carbone, E. (1973). The acquisition of a new phonological contrast: The case of stop consonants in French-English bilinguals. *Journal of the Acoustic Society of America, 54*, 421–428.

Corbera, S., Corral, M. J., Escera, C., & Idiazábal, M. A. (2005). Abnormal speech sound representation in developmental stuttering. *Neurology, 65,* 1246–1252.

Cutler, A., Weber, A., Smits, R., & Cooper, N. (2004). Patterns of English phoneme confusions by native and non-native listeners. *Journal of the Acoustical Society of America, 116*, 3668–3678.

Deouell, L. Y. (2007). The frontal generator of the mismatch negativity revisited. *Journal of Psychophysiology, 21*, 188–203.

Díaz, B., Baus, C., Escera, C., Costa, A., & Sebastián-Gallés, N. (2008). Brain potentials to native phoneme discrimination reveal the origin of individual differences in learning the sounds of a second language. *Proceedings of the National Academy of Sciences of the United States of America, 105*, 16083–16088.

Díaz, B., Mitterer, H., Broersma, M., & Sebastián-Gallés, N. (2012). Individual differences in late bilinguals' L2 phonological processes: From acoustic-phonetic analysis to lexical access. *Learning and Individual Differences, 22,* 680–689.

Escera, C., Alho, K., Winkler, I., & Näätänen, R. (1998). Neural mechanisms of involuntary attention to acoustic novelty and change. *Journal of Cognitive Neuroscience, 10*, 590–604.

Flege, J. E. (1995). Second language speech learning: Theory, findings and problems. In W. Strange (ed.), *Speech perception and linguistic experience,* pp. 233–272. Baltimore, MD: York Press.

Giard, M. H., Perrin, F., Pernier, J., & Bouchet, P. (1990). Brain generators implicated in the processing of auditory stimulus deviance : A topographic event-related potential study. *Psychophysiology, 27*, 627–640.

Goldinger, S. D. (2007). A complementary-systems approach to abstract and episodic speech perception. In J. Trouvain, & W. J. Barry (eds.), *Proceedings of the 16th International Congress of Phonetic Sciences*, pp. 49–54. Dudweiler: Pirrot.

Golestani, N., Molko, N., Dehaene, S., LeBihan, D., & Pallier, C. (2007). Brain structure predicts the learning of foreign speech sounds. *Cerebral Cortex, 17*, 575–582.

Golestani, N., Paus, T., & Zatorre, R. J. (2002). Anatomical correlates of learning novel speech sounds. *Neuron, 35*, 997–1010.

Golestani, N., & Zatorre, R. J. (2009). Individual differences in the acquisition of second language phonology. *Brain and Language, 109*, 55–67.

Goto, H. (1971). Auditory perception by normal Japanese adults of the sounds "l" and "r". *Neuropsychologia, 9*, 317–323.

Hara, K., Ohta, K., Miyajima, M., Hara, M., Iino, H., Matsuda, A., Watanabe, S., Matsushima, T., Maehara, T., & Matsuura, M. (2012). Mismatch negativity for speech sounds in temporal lobe epilepsy. *Epilepsy & Behavior, 23*, 335-341.

Hawkins, S., & Stevens, K. N. (1985). Acoustic and perceptual correlates of the non-nasal-nasal distinction for vowels. *The Journal of the Acoustical Society of America, 77*, 1560–1575.

Indefrey, P. (2006). A meta-analysis of hemodynamic studies on first and second language processing: Which suggested differences can we trust and what do they mean? *Language Learning, 56*, 279–304.

Jääskeläinen, I. P., Pekkonen, E., Hirvonen, J., Sillanaukee, P., & Näätänen, R. (1996). Mismatch negativity subcomponents and ethyl alcohol. *Biological Psychology, 43*, 13–25.

Johnson, J. S., & Newport, E. L. (1989). Critical period effects in second language learning: The influence of maturational state on the acquisition of English as a second language. *Cognitive Psychology, 21*, 60–99.

Kim, K., Relkin, N., Lee, K. M., & Hirsch, J. (1997). Distinct cortical areas associated with native and second languages. *Nature, 388*, 171–174.

Klatt, D. (1980). Software for a cascade/parallel formant synthesizer. *Journal of the Acoustical Society of America, 67*, 971–995.

Kuhl, P. K., Tsao, F. M., & Liu, H. M. (2003). Foreign-language experience in infancy: Effects of short-term exposure and social interaction on phonetic learning. *Proceedings of the National Academy of Sciences, 100*, 9096–9101.

Kujala, T., & Näätänen, R. (2010). The adaptive brain: a neurophysiological perspective. *Progress in Neurobiology, 91,* 55–67.

Lahiri, A., & Marslen-Wilson, W. (1991). The mental representation of lexical form: A phonological approach to the recognition lexicon. *Cognition, 38*, 254–294.

Lavikainen, J., Huotilainen, M., Pekkonen, E., Ilmoniemi, R. J., & Näätänen, R. (1994). Auditory stimuli activate parietal brain regions: a whole-head MEG study. *NeuroReport, 6*, 182–184.

Lengeris, A., & Hazan, V. (2010). The effect of native vowel processing ability and frequency discrimination acuity on the phonetic training of English vowels for native speakers of Greek. *The Journal of the Acoustical Society of America, 128*, 3757–3768.

Lenneberg, E. H. (1967). *Biological foundations of language*. New York: Wiley.

McClelland, J. L., McNaughton, B. L., & O'Reilly, R. C. (1995). Why there are complementary learning systems in the hippocampus and neo-cortex: Insights from the successes and failures of connectionists models of learning and memory. *Psychological Review, 102*, 419–457.

Mitterer, H., & McQueen, J. M. (2009). Foreign subtitles help but native-language subtitles harm foreign speech perception. *PLoS ONE, 4*, e7785.

Näätänen, R. (1990). The role of attention in auditory information processing as revealed by event-related potentials and other brain measures of cognitive function. *Behavioral and Brain Sciences, 13*, 201–288.

Näätänen, R. (2001). The perception of speech sounds by the human brain as reflected by the mismatch negativity (MMN) and its magnetic equivalent (MMNm). *Psychophysiology, 38*, 1–21.

Näätänen, R., Gaillard, A. W .K., Mäntysalo, S. (1978). Early selective-attention effect on evoked potential reinterpreted. *Acta Psychologica, 42*, 313–29.

Näätänen, R., Lehtokoski, A., Lennes, M., Cheour, M., Huotilainen, M., Iivonen, A., Vainio, M., Alku, P., Ilmoniemi, R., Luuk, A., Allik, J., Sinkkonen, J. & Alho, K. (1997). Language-specific phoneme representations revealed by electric and magnetic brain responses. *Nature, 385*, 432–434.

Nenonen, S., Shestakova, A., Huotilainen, M., & Näätänen, R. (2005). Speech-sound duration processing in a second language is specific to phonetic categories. *Brain and Language, 92*, 26–32.

Oldfield, R. C. (1971). The assessment and analysis of handedness: The Edinburgh inventory. *Neuropsychologia, 9*, 97–113.

Oyama, S. (1976). A sensitive period for the acquisition of a non-native phonological system. *Journal of Psycholinguitic Research, 5*, 261–285.

Paavilainen, P., Mikkonen, M., Kilpeläinen, M., Lehtinen, R., Saarela, M., & Tapola, L. (2003). Evidence for the different additivity of the temporal and frontal generators of Mismatch Negativity: A human auditory event-related potential study. *Neuroscience Letters, 349,* 79–82.

Pallier, C., Bosch, L., & Sebastián-Gallés, N. (1997). A limit on behavioral plasticity in speech perception. *Cognition, 64*, B9–B17.

Pallier, C., Colomé, A., & Sebastián-Gallés, N. (2001). The influence of native-language phonology on lexical access: Exemplar-based versus abstract lexical entries. *Psychological Science,* 12, 445–449.

Patkowski, M. S. (1980). The sensitive period for the acquisition of syntax in a second language. *Language Learning, 30*, 449–468.

Perani, D., & Abutalebi, J. (2005). The neural basis of first and second language processing. *Current Opinion in Neurobiology, 15*, 202–206.

Perani, D., Paulesu, E., Sebastián-Gallés, N., Dupoux, E., Dehaene, S., Bettinardi, V., Cappa, S., Fazio, F., & Mehler, J. (1998). The bilingual brain - proficiency and age of acquisition of the second language. *Brain, 121*, 1841–1852.

Poulisse, N., Bongaerts, T., & Kellerman, E. (1990). *The use of compensatory strategies by Dutch learners of English*. Dordrecht, the Netherlands: Foris.

Pulvermüller, F., & Schumann, J. H. (1994). Neurobiological mechanisms of language acquisition. *Language Learning, 44*, 681–734.

Reinisch, E., & Sjerps, M. J. (2013). The uptake of spectral and temporal cues in vowel perception is rapidly influenced by context. *Journal of Phonetics, 41,* 101-116.

Rietveld, T., & van Heuven, V. (1997). *Algemene fonetiek [general phonetics].* Bussum: Coutinho.Sato, Y., Hirooki, Y., Tomiharu, H., Takeyuki, S., Naoko, S., Tadayoshi, N., & Sunao, K. (2000). The effect of deviant stimulus probability on the human mismatch process. *NeuroReport, 11*, 3703–3708.

Sebastián-Gallés, N., & Baus, C. (2005). On the relationship between perception and production in L2 categories. In A. Cutler (ed.), *Twenty-first century psycholinguistics: Four cornerstones,* pp. 279–292. New York: Erlbaum.

Sebastián-Gallés, N., & Díaz, B. (2012). First and second language speech perception: Graded learning. *Language Learning, 62,* 131–147.

Sebastián-Gallés, N., Echeverría, S., & Bosch, L. (2005). The influence of initial exposure on lexical representation: Comparing early and simultaneous bilinguals. *Journal of Memory and Language, 52*, 240–255.

Sebastián-Gallés, N., Soriano-Mas, C., Baus, C., Díaz, B., Ressel, V., Pallier, C., Costa, A., & Pujol, J. (2012). Neuroanatomical markers of individual differences in native and non-native vowel perception. *Journal of Neurolinguistics*, 25, 150–162.

Sebastián-Gallés, N., & Soto-Faraco, S. (1999). Online processing of native and non-native phonemic contrasts in early bilinguals. *Cognition, 72*, 111–123.

Shalgi, S., & Deouell, L. Y. (2007). Direct evidence for differential roles of temporal and frontal components of auditory change detection. *Neuropsychologia, 45*, 1878–1888.

Sussman, E., & Winkler, I. (2001). Dynamic sensory updating in the auditory system. *Cognitive Brain Research, 12*, 431–439

Tremblay, K., Kraus, N., & McGee, T. (1998). The time course of auditory perceptual learning: Neurophysiological changes during speech-sound training. *Neuroreport, 9*, 3557–3560.

Tricomi, E., Delgado, M. R., McCandliss, B. D., McClelland, J. L., & Fiez, J. A. (2006). Performance feedback drives caudate activation in a phonological learning task. *Journal of Cognitive Neuroscience, 18*, 1029–1043.

Wartenburger, I., Heekeren, H. R., Abutalebi, J., Cappa, S. F., Villringer, A., & Perani, D. (2003). Early setting of grammatical processing in the bilingual brain. *Neuron, 37*, 159–170.

Weber, A., & Cutler, A. (2004). Lexical competition in non-native spoken-word recognition. *Journal of Memory and Language, 50*, 1-25.

Weber-Fox, C. M., & Neville, H. J. (1996). Maturational constraints on functional specialization for language processing: ERP and behavioral evidence in bilingual speakers. *Journal of Cognitive Neuroscience, 8*, 231–256.

Winkler, I., Kujala, T., Tiitinen, H., Sivonen, P., Alku, P., Lehtokoski, A., Czigler, I., Csépe, V., Ilmoniemi, R. J., & Näätänen, R. (1999). Brain responses reveal the learning of foreign language phonemes. *Psychophysiology, 36*, 638–642.

Wong, P. C., Perrachione, T. K., & Parrish, T. B. (2007). Neural characteristics of successful and less successful speech and word learning in adults. *Human Brain Mapping, 28*, 995–1006.

Wong, P. C., Warrier, C. M., Penhune, V. B., Roy, A. K., Sadehh, A., Parrish, T. B., & Zatorre, R. J. (2008). Volume of left Heschl's gyrus and linguistic pitch learning. *Cerebral Cortex, 18*, 828–836.

Yago, E., Escera, C., Alho, K., & Giard, M. H. (2001). Cerebral mechanisms underlying orienting of attention towards auditory frequency changes. *Neuroreport, 12*, 2583–2587.

FOOTNOTES

[1] Table 7, http://www.unesco.org/new/en/culture/resources/report/the-unesco-world-report-on-cultural-diversity/

[2] The phoneme /O/ corresponds to the sound in the English word "lord" (/loːɹd/, with /ɹ/ pronounced in most American but only few British accents) and /ɑ/ to the sound in "car" (/kɑɹ/). Together with these vowels, the phonetic symbol " ː " denotes long duration and the symbol " ~ " nasality.

[3] One could argue that our participants were not actually late learners of English because the Netherlands television broadcasts shows and films in English. That means that our participants may have had access to native English speech before receiving formal English language education. We believe that the influence of spoken English during the participants' early childhood must have been minimal. Firstly, learning the speech sounds of a language requires social interaction; mere exposure to that particular language may not be enough to learn its speech sounds (Kuhl, Tsao & Liu, 2003), at least in early ages; and secondly, English spoken contents on Dutch television are accompanied by Dutch subtitles, and this combination of auditory presented foreign language with visually presented native language has been showed to be detrimental for foreign speech perception (Mitterer & McQueen, 2009).