

Binaural and/or Monophonic self-listening as a tool for music performance simulation: evaluating immersion and neural responses

Autoescuta binaural e/ou monofónica como ferramenta para simulação da performance musical: avaliação da imersão e respostas neurais



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Abstract: This exploratory study investigated the effects of binaural and monophonic self-listening on musicians' neural responses during the mental simulation of their own music performance. Six musicians participated in two EEG-monitored sessions in which they listened to binaural and monophonic recordings of their performances in a counterbalanced order. We hypothesized that binaural recordings, by preserving head-related transfer functions, would elicit stronger neural activation in spatial perception, emotional engagement, and motor simulation than monophonic recordings. The study examined differences in brain activity related to sensory, motor, and emotional processing between the two conditions. The results indicated that binaural self-listening induced broader cortical activation patterns aligned with sensorimotor and emotional processes. At the same time, monophonic recordings were associated with greater frontal activation linked to cognitive effort and attentional control. These findings suggest that binaural self-listening enhances mental performance simulation and may offer a promising tool for therapeutic interventions targeting music performance anxiety in contexts where physical practice is not feasible. Future research should employ larger samples and inferential analyses to strengthen these initial findings.

Keywords: binaural self-listening, binaural recording, music performance simulation; neural responses; electroencephalogram.

Resumo: Este estudo exploratório investigou os efeitos da autoescuta binaural e monofônica nas respostas cerebrais de músicos durante a simulação mental de suas próprias performances. Seis músicos participaram de sessões com EEG, escutando gravações binaurais e monofônicas de suas interpretações em ordem contrabalançada. Hipotetizou-se que as gravações binaurais, por preservarem as funções de transferência relacionadas à cabeça, induziriam uma ativação neural mais intensa relacionada à percepção espacial, envolvimento emocional e simulação motora. A atividade

cerebral foi comparada entre os domínios sensorial, motor e emocional. A autoescuta binaural induziu uma ativação cortical mais ampla, associada a processos sensório-motores e emocionais, enquanto a autoescuta monofônica apresentou maior ativação frontal, ligada ao esforço cognitivo e controle atencional. Esses resultados sugerem que a autoescuta binaural potencializa a simulação mental da performance e pode apoiar estratégias terapêuticas para ansiedade na performance musical, especialmente quando a prática física não é viável. Investigações futuras devem recorrer a amostras de maior dimensão e a análises inferenciais, de modo a consolidar e testar de forma mais robusta estes resultados preliminares.

Palavras-chave: auto escuta binaural; gravação binaural; simulação da performance musical; respostas neurais; EEG (Eletroencefalograma).

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1 Introduction

Physiological and psychological skills, such as physical control and mental resilience, are essential for effective musical performance, ensuring both technical and emotional quality (Papageorgi; Kopiez, 2012). Meeting personal and collective expectations regarding these skills is a complex and arduous challenge, often triggering feelings of dissatisfaction that result in heightened and debilitating levels of music performance anxiety (MPA) (Ely, 1991; Papageorgi, 2021; Papageorgi; Creech; Welch, 2011; Papageorgi; Welch, 2020). High levels of MPA can trigger stress responses that impair public performance, regardless of the musician's experience (Fernholz et al., 2019; Kenny, Driscoll, Ackermann, 2014b). The prevalence of MPA is estimated to range from 16.5% to 60% among musicians and up to 83.1% among music students (Barros et al., 2022; Fernholz et al., 2019); both studies used self-reported assessments. MPA can be triggered by various factors, including the perception of high personal and collective expectations, which increases the complexity and challenge of delivering a high-quality musical performance (Barros et al., 2022; Castiglione; Rampullo; Cardullo, 2018; Ryan, 2021; Zhukov, 2019). Coping strategies are crucial, as a positive perception of performance can mitigate symptoms (Biasutti; Concina, 2014; Irie; Morijiri; Yoshie, 2023; Macafee; Comeau, 2023). Therefore, the development, exploration, and refinement of strategic approaches to managing MPA levels are vital during performance preparation to minimize its impact on musicians' lives and transform it, through control, into a beneficial factor for performance (Kenny, 2011; Kenny; Davis; Oates, 2004; Kenny; Arthey; Abbass, 2014a; Kokotsaki; Davidson, 2003).

Some training protocols for managing MPA are implemented without the performer playing their instrument, such as biofeedback training (Barros et al., 2018; Rabelo; Mármora; Ribeiro, 2018; Thurber et al., 2010; Wells et al., 2012), which

requires maximum physiological relaxation to prevent movement from altering the monitored parameters (Borges et al., 2023). In this context, mental simulation emerges as a promising tool, since it can be combined with various approaches for the treatment of MPA (Hoffman; Hanrahan, 2012; Schaefer, 2015). Mental simulation has been widely recognized as a valuable strategy in music performance preparation, offering cognitive, emotional, and physiological benefits (Bernardi et al., 2012). It activates neural circuits related to motor planning, auditory imagery, and emotional processing, thereby enhancing technical skills, musical memory, and artistic expressiveness (Clark; Williamon, 2011; Toth et al., 2020). Research shows that mental practice can reduce physical strain, optimize study time, and improve performance confidence, especially when combined with physical practice or tools such as self-listening (Volioti; Williamon, 2016; Hewitt, 2001). Furthermore, mental simulation can serve as an effective coping strategy for music MPA, particularly when it includes mental rehearsal of the performance environment, promoting emotional resilience and reducing stress reactivity (Parreiras Gonçalves; Torchia Zanon, 2022). The use of recordings of one's own performance has also proven to be an effective tool for visualizing and mentally simulating a performance (Clark; Williamon, 2011).

The effectiveness of mental simulation, however, may vary according to individual visualization and concentration skills (Emmelkamp, 2013; Keller, 2012; Kosslyn et al., 1984). Techniques such as listening to recordings of oneself and practicing guided visualization can amplify the benefits, making them more effective (Clark; Williamon, 2011a; Guillot; Collet, 2005; Holmes; Collins, 2001). Self-listening has also proven to be a valuable tool in performance preparation, aiding musical awareness and refining technical and interpretive aspects (Hewitt, 2001; Volioti; Williamon, 2016). Borges et al. (2023) highlight the contribution of self-listening in mental performance simulation.

Based on the premise that binaural recordings capture sound in three dimensions, contributing to the perception of acoustic spatiality, immersive characteristics can be enhanced when the correct head-related transfer function (HRTF), specific to each individual, is considered during the recording process (Everest; Pohlmann, 2022; Rumsey, 2021). This study adopted an exploratory approach, aiming to gain an initial understanding of the neurophysiological effects of binaural listening in musicians. The hypothesis was that binaural recordings could elicit brain responses associated with more realistic and immersive auditory experiences compared to monophonic recordings. At this stage, the study sought to investigate whether these different auditory perspectives—binaural and monophonic—could serve as supportive tools for the mental simulation of musical performance in controlled environments, particularly in contexts where instrumental execution is not possible, such as in therapeutic protocols targeting MPA through neurofeedback and biofeedback techniques. Recent advances in music neuroscience have highlighted the complex neural mechanisms involved in musical listening, particularly through tools such as electroencephalography (EEG) and functional magnetic resonance imaging (fMRI), which allow detailed observation of brain oscillations and connectivity during auditory experiences (Koelsch, 2011; Zatorre; Chen; Penhune, 2007). In this sense, the present study preliminarily explored the neurophysiological responses of musicians during self-listening to their musical performances in two distinct formats: monophonic and binaural, highlighting the particularities of each condition as an auxiliary tool in the mental simulation of musical performance, and contributing to intervention protocols for MPA in which active execution is not possible during treatment.

1.1 Mental Simulation

Understanding the different ways to simulate music performance is crucial for developing effective MPA management strategies. Performance simulation can take

three primary forms: 1) Live Performance — recreating the actual performance situation (Kendrick et al., 1982; Wiederhold; Wiederhold, 2005), which can increase stress and anxiety, making MPA more difficult to control; 2) Virtual Performance Simulation — using technology such as augmented, virtual, and mixed reality, to offer a less intimate but more accessible alternative, with promising results in MPA control (Aufegger Et Al., 2017; Bissonnette et al., 2015, 2016; Fanger et al., 2020; Gorini; Riva, 2008; Orman, 2003, 2004; Owens; Beidel, 2015; Williamon; Aufegger; Eiholzer, 2014); 3) Mental Simulation — mental recreation of the performance that allows musicians to mentally rehearse and improve their skills without the pressure of a live audience, enhancing confidence and reducing MPA (Clark; Williamon, 2011b; Connolly; Williamon, 2004; Hoffman; Hanrahan, 2012; Immonen; Ruokonen; Ruismäki, 2012; Kosslyn et al., 1984; Osborne; Greene; Immel, 2014).

Several studies have focused on mental simulation and imagery (Clark; Williamon, 2011b; Connolly; Williamon, 2004; Hoffman; Hanrahan, 2012). Mental simulation focuses on the detailed practice of motor tasks and specific actions, enhancing muscle memory and psychological preparation, which is useful when physical practice is limited. By contrast, imagery involves creating vivid, sensory mental images that engage multiple senses, offering a rich, immersive mental experience (Keller, 2012; Mielke & Comeau, 2019). Mental simulation can help improve focus, manage MPA, and enhance real-life performance (Keller, 2012). Musical imagination activates brain areas involved in physical performance (Kristeva et al., 2003), mainly in auditory and motor regions (Tanaka; Kirino, 2022). However, the effectiveness of mental simulation can vary depending on individual strategies and skills in visualization and concentration (Emmelkamp, 2013; Keller, 2012; Kosslyn et al., 1984).

Self-listening is the process of listening to your own performance, whether live or via an audio recording (Hewitt, 2001). Borges et al. (2023) suggest that self-listening can serve as a “guiding thread,” providing an auditory foundation that facilitates more precise, contextually informed mental simulation. This approach is particularly beneficial in MPA management training programs that do not allow the use of the instrument during treatment sessions, such as biofeedback and neurofeedback training, which require physiological relaxation and participant immobility.

1.2 Binaural Recording and Listening

When considering self-listening as a tool for performance simulation, it is essential to account for spatiality, which can be captured in different ways, such as through stereo or binaural recordings. Using intensity and/or time cues, a two-channel stereo recording uses two channels (left and right) to create a sense of direction, allowing the listener to identify the source of sounds in space (Bartlett, 2016). Binaural recording simulates human hearing by using microphones positioned at the ears to capture sound in three dimensions (left-right, up-down, front-back), providing greater and more realistic immersion (Rumsey, 2021). According to Everest and Pohlmann (2022), the choice between stereo and binaural depends on the desired level of spatial realism, with binaural recording ideal for more immersive simulations, mainly when the performer’s individual HRTFs are used, thereby ensuring greater sound realism.

In recent years, binaural recording has been used in various studies investigating auditory perception and its neurological implications. Brinkmann et al. (2014) evaluated the perceptual authenticity of binaural simulations in reverberant environments, showing that this technology can be indistinguishable from real auditory experiences for certain types of sounds, such as the

human voice. Additionally, research by Cheng and Wakefield (2001) explored the use of moving-sound-source synthesis with HRTFs, expanding the possibilities for spatial reproduction in musical applications and for auditory simulations in general.

The HRTF plays a fundamental role in describing how individual anatomy (head, ears, and torso) modifies sound as it reaches the ears, making it crucial for recreating spatial auditory experiences. The importance of this technology lies in its ability to provide a more immersive and realistic listening experience, making it valuable for areas such as virtual reality, headphone development, and neuroscientific research (Zieliński; Antoniuk; Lee, 2022). This method enables a three-dimensional perception of sound, accurately replicating the way we hear in the real world (Brinkmann et al., 2014). Preserving individual HRTFs in binaural recordings is crucial for creating realistic auditory simulations that are virtually indistinguishable from real experiences (Brinkmann et al., 2014). Studies show that the use of personalized HRTFs significantly improves sound localization perception, reducing spatial errors, especially in the median and front-back planes, compared with generic HRTFs (Grijalva et al., 2018; Kudo et al., 2006; Romigh et al., 2013). The lack of personalization can lead to poor spatial localization, as demonstrated by Yao and Chen (2013), who emphasize the importance of adjusting HRTFs to enhance auditory quality and precision. Moreover, Gan et al. (2017) stress that HRTF personalization is crucial for 3D audio rendering, improving spatial perception in augmented and virtual reality devices. Finally, Cheng and Wakefield (2001) demonstrated that using interpolated HRTFs enables the accurate reproduction of moving sound sources in electroacoustic environments, thereby enhancing spatial perception.

Binaural recordings have gained prominence in the field of musical performance because they create realistic acoustic simulations, allowing musicians to practice and interact with virtual environments. Examples include recreating historical

acoustic spaces and simulating stage conditions to enhance musical perception and training (Chu, 2004; Martens; Woszczyk, 2004), as well as their use in online performances and remote recordings (Hagan, 2022; Mróz et al., 2022). Studies show that musicians respond differently to binaural auditory stimuli compared to non-musicians, suggesting significant potential for this technology in musical training (Ioannou et al., 2015). Furthermore, musicians have shown a preference for binaural audio over stereo due to the increased immersion it provides (Turchet; Tomasetti, 2023), and equalization has been identified as crucial for enhancing spatial perception, making the technology even more effective (Schärer; Lindau, 2009).

Binaural performance monitoring has also been studied and offers significant advantages over stereo: improved instrument localization, facilitating coordination in studios and remote collaborations (Turchet; Tomasetti, 2023); improved sound proximity and localization, benefiting rehearsals and live performances (Griesinger, 2016); a more realistic auditory scene compared to ambisonics (They, 2019); and increased immersion and enhanced musician performance during recording sessions (Bauer et al., 2022).

In general, binaural listening can enhance spatial perception and immersion during mental performance simulation. Zieliński et al. (2022) demonstrated that this technology improves the localization of sound sources in simulations, making it particularly useful for creating immersive experiences. A growing body of research highlights the versatility and impact of binaural self-listening both in simulations and live performance (monitoring). Furthermore, it can be used as an auxiliary tool in instrumental performance simulation during treatment protocols for performance anxiety when playing the instrument is impossible.

1.3 Neural Processing of Music self-listening

Research indicates that musical listening involves multiple neural processes that interact to shape the auditory experience. EEG and functional magnetic resonance imaging (fMRI) have enabled the identification of oscillatory patterns and connectivity between brain regions, thereby clarifying how the human brain processes music (Koelsch, 2011; Zatorre, Chen, Penhune, 2007). Almudena et al. (2021) analyzed the results of electroencephalographic investigations addressing music perception from various technical analytical perspectives. The main themes explored include intrinsic musical features, such as tonality, rhythm, dissonance, and musical syntax, as well as musical emotions, expectations, differences between pleasant and unpleasant music, and the effects of familiarity and musical experience. Thus, research primarily encompasses auditory perception, memory, cognition, emotion, motor processing, harmonic expectation, and sensory integration.

Auditory perception is primarily monitored by the primary and secondary auditory cortices, which analyze fundamental sound characteristics such as frequency, timbre, and intensity (Patterson et al., 2002; Zatorre; Belin, 2001). Furthermore, the secondary auditory cortex is essential for processing more complex patterns, including melody, rhythm, and harmony, facilitating the recognition and organization of musical stimuli (Hyde; Peretz; Zatorre, 2008; Koelsch, 2011).

Musical memory is also a central factor in auditory experience. It is strongly linked to the activity of the hippocampus and prefrontal cortex, regions associated with the storage and retrieval of musical information (Koelsch, 2011). Studies show that repeated exposure to a melody can lead to greater coherence in neural activity, reflecting memory encoding processes and pattern recognition (Zatorre; Chen; Penhune, 2007).

Regarding music cognition, music activates neural mechanisms shared with language, such as Broca's and Wernicke's areas, which process both linguistic syntax and musical syntax (Koelsch, 2011). This neural overlap suggests that the brain analyzes musical organization like that of spoken sentences.

Musical emotion is another widely studied aspect in music neuroscience. Research indicates that the amygdala and orbitofrontal cortex play fundamental roles in regulating emotions evoked by music (Koelsch, 2011). Additionally, variations in alpha (8–12 Hz) and beta (13–30 Hz) band activity have been associated with emotional differences in music perception, reflecting the involvement of these brain oscillations in the affective experience of music (Schaefer; Vlek; Desain, 2011; Lin et al., 2022).

Motor processing is also intrinsically linked to music listening, as motor regions are activated even when an individual is merely listening to music without moving. This activation of the motor cortex suggests that listening to musical rhythms triggers an internal simulation of movement, which may be related to our ability to anticipate and synchronize with rhythmic patterns (Zatorre; Chen; Penhune, 2007).

Harmonic expectation, in turn, involves the prefrontal cortex, which is responsible for predicting and detecting surprises in musical structure (Koelsch, 2011). The brain can anticipate harmonic changes based on previously learned patterns, activating neural networks involved in predicting musical events.

Finally, sensory integration enables listeners to combine auditory, motor, and emotional information to create a coherent and meaningful musical experience. The superior parietal cortex is involved in this process, enabling the fusion of different sensory stimuli and facilitating the understanding of musical structure within the context of the overall auditory experience (Zatorre; Chen; Penhune, 2007).

These studies demonstrate that music listening is not an isolated process but rather a complex interaction among distributed neural networks involving perception, cognition, memory, emotion, and motor processing. EEG has been a fundamental tool in mapping these interactions and understanding how music influences brain function at various levels. Most of the studies analyzed by Almudena et al. (2021) involved participants in a resting state, listening to musical excerpts, note sequences, or chords. A significant gap identified is the scarcity of studies investigating music perception in participants during music performance or imagined performance. Additionally, no studies were found that use self-listening of a performance combined with EEG monitoring to analyze musicians' perception.

Based on the immersive properties of binaural recordings and their ability to preserve individualized spatial cues, we hypothesized that binaural self-listening would promote stronger neural activation patterns associated with motor imagery, emotional engagement, and sensory integration, compared to traditional monophonic or usual stereo recordings.

2 Methods

2.1 Participants

The sample consisted of six musicians (three women and three men) from a Portuguese university, including three professors and three students, who played three different instruments. Each of the three instruments (cello, piano, and guitar) was represented by one professor and one student (Tab. 1). The professors held doctoral degrees, and the students were enrolled in bachelor's or master's programs. Given the small sample size ($n=6$), this study is intended as exploratory research. As such, the results should be interpreted with caution and viewed as preliminary evidence rather than definitive conclusions.

Tabela 1 – Participant sample characteristics

| Participant | Role | Instrument | Gender | Age | Experience (years) | Performances per year | Degree |
|-------------|-----------|------------|--------|-----|--------------------|-----------------------|--------|
| P1 | Professor | Cello | Male | 41 | 25 | 20 | PhD |
| P2 | Professor | Piano | Female | 56 | 47 | 30 | PhD |
| P3 | Professor | Guitar | Male | 45 | 30 | 30 | PhD |
| P4 | Student | Cello | Female | 22 | 10 | 2 | BA |
| P5 | Student | Piano | Male | 36 | 27 | 4 | MA |
| P6 | Student | Guitar | Female | 25 | 15 | 4 | BA |

Source: Data from Author (2025).

Description of the image: table displaying sociodemographic and musical characteristics of participants, including role, instrument, gender, age, years of experience, annual performances, and academic degree.

Overall, the sample had a mean age of 37.5 years (SD = 12.7), 25.7 years of musical experience (SD = 12.9), and performed publicly an average of 15 times per year (SD = 13.3). They also self-reported good health and hearing. Participation was voluntary, and all participants were fully informed about the study procedures and provided written informed consent. The study was approved by the Ethics and Deontology Committee of the University of Aveiro (CED-UA), under approval no. 01-CED/2019. Participants also self-reported good health and hearing.

2.2 Procedures

Each participant was invited to prepare a piece of their choice from their own repertoire, with a maximum duration of 6 minutes, appropriate to their technical and interpretative level. The main selection criterion was that the musicians felt prepared, confident, and comfortable performing and recording the piece.

The study was conducted in two phases: 1) simultaneous audio recording of the participants' performances from binaural and stereo perspectives, and 2) EEG recordings while listening to the performance from the perspectives of binaural and monophonic.

2.2.1 Phase 1: audio recording of the performances

The musical performances were recorded simultaneously under two conditions: binaural and stereo. The recordings took place in a concert hall at the university, familiar to all participants and with favorable acoustic conditions, ensuring consistent reverberation time and sound diffusion for all recordings and participants. Before the recordings began, the microphones were calibrated, installed, and tested. Participants were then given 15 minutes to play their instruments freely, allowing them to adapt to the room and to the microphones attached to their ears.

After the adaptation period, the participants performed their pieces. Immediately after each performance, they rated their satisfaction on a scale from 1 (dissatisfied) to 10 (satisfied). Each participant had the opportunity to follow this recording protocol three times consecutively, allowing the researchers to select the best of the three performances for analysis in the study's second phase. The mean satisfaction rating for the most satisfying performance was 9 (SD = 1). No sound compensation was made between the different instruments in terms of volume levels, acoustic characteristics, or spatial positioning.

Recording performance: We used a Sound Devices MixPre 6 II recorder to capture both formats. Binaural recording was conducted using two Primo 173 omnidirectional microphones, each attached to the participants' auricle and cross-channel calibrated with a Bruel & Kjaer Type 4231. For the stereo recording, we used two Rode NT2 microphones configured in an XY coincident pattern, placed in front of the instrument and aligned with its primary sound radiation angle, and subsequently summed to mono, following the principles of this type of recording. It is important to note that a specialist recommended this microphone configuration. The stereo microphones were positioned from an audience perspective, and channel summation to dual mono was adopted to limit spatial-intensity cues and support controlled EEG comparisons.

2.2.2 Phase 2: self-listening under binaural and monophonic conditions monitored by EEG

During the self-listening procedure, all participants completed two listening sessions of their performances, monitored by EEG, corresponding to the two different recording approaches implemented in the first phase. Two groups of three participants were formed to counterbalance years of experience. Each group included professors and students of each instrument, although in different proportions: one group comprised two professors and one student, and the other, two students and one professor. Participants listened to the same two recordings—one in binaural format and the other in mono—but in different orders. Specifically, one group listened to the binaural version first, followed by the monophonic version, while the other group began with the monophonic version and then listened to the binaural version.

Following EEG setup, preparation, and positioning, participants completed a baseline data collection, during which they were monitored for 6 minutes without any stimuli. During this period, they were asked to remain still for 3 minutes with their eyes open, and then for 3 minutes with their eyes closed. Before each listening session, participants were instructed to imagine their performance of the piece while simultaneously listening to the recording, remaining as still as possible to avoid interfering with EEG monitoring and data recording. During the listening sessions, participants were blindfolded to prevent visual input from generating EEG signal artifacts.

After baseline data collection, participants completed the first listening session for their assigned listening condition. At the end of this session, they completed an unrelated distractor task for approximately 30 minutes to minimize any carry-over effects. This task consisted of completing questionnaires on sociodemographic data, interests, and experiences related to wine tourism. Participants then viewed two slide shows (20

images of wine cellars and 20 of vineyards) while their brain activity was recorded via EEG. Finally, the second listening session was conducted, corresponding to the assigned listening condition for their participant group.

EEG recordings: the EEG recordings and data processing were carried out using specific hardware and software to ensure precision and efficiency at every stage. The hardware used was the ANT system with the 32-channel WaveGuard cap (ANT Neuro), offering high spatial resolution, superior signal quality, and adaptable comfort for participants. This technology provides comprehensive scalp coverage while simultaneously capturing electrical activity from multiple brain regions, which is essential for detailed analysis of brain responses. Data acquisition was performed using Asalab software, version 4.10.1, which allows signal recording, initial filtering, and artifact exclusion, using predefined cutoffs of $-150 \mu\text{V}$ to $150 \mu\text{V}$ (Fabiani et al., 2009; Luck, 2014). Asalab provides real-time data control, ensuring the quality of signals captured and seamless integration with other software, such as standardized low-resolution brain electromagnetic tomography (sLORETA).

sLORETA: standardized low-resolution brain electromagnetic tomography (sLORETA) is a 3D source localization method developed to model and localize brain electrical activity from multichannel EEG recordings (Pizzagalli et al., 2001; Northoff et al., 2006; Northoff et al., 2006). sLORETA was developed by Pascual-Marqui and Lehmann (1994), Pascual-Marqui (1999, 2002), and Pascual-Marqui et al. (2002) to estimate the locations of neural generators underlying brain electrical activity using a maximally smooth linear inverse solution. This means that the algorithm assumes a more dispersed and continuous distribution of neural currents to minimize errors in reconstructing EEG signal sources. Validation studies of the sLORETA algorithm have shown that it can accurately estimate the location of neural sources, even in deep brain regions, as demonstrated by combined fMRI-EEG experiments (Vitacco et

al., 2002). Mulert et al. (2004) used a classic oddball paradigm with pure tones to compare brain activations measured by fMRI with those estimated by sLORETA. Their results showed that sLORETA correctly localized the neural correlates of unimodal auditory stimuli, with minimal localization error. This error occurs due to the method's low spatial resolution, but its accuracy is sufficient for exploratory studies and functional EEG analyses.

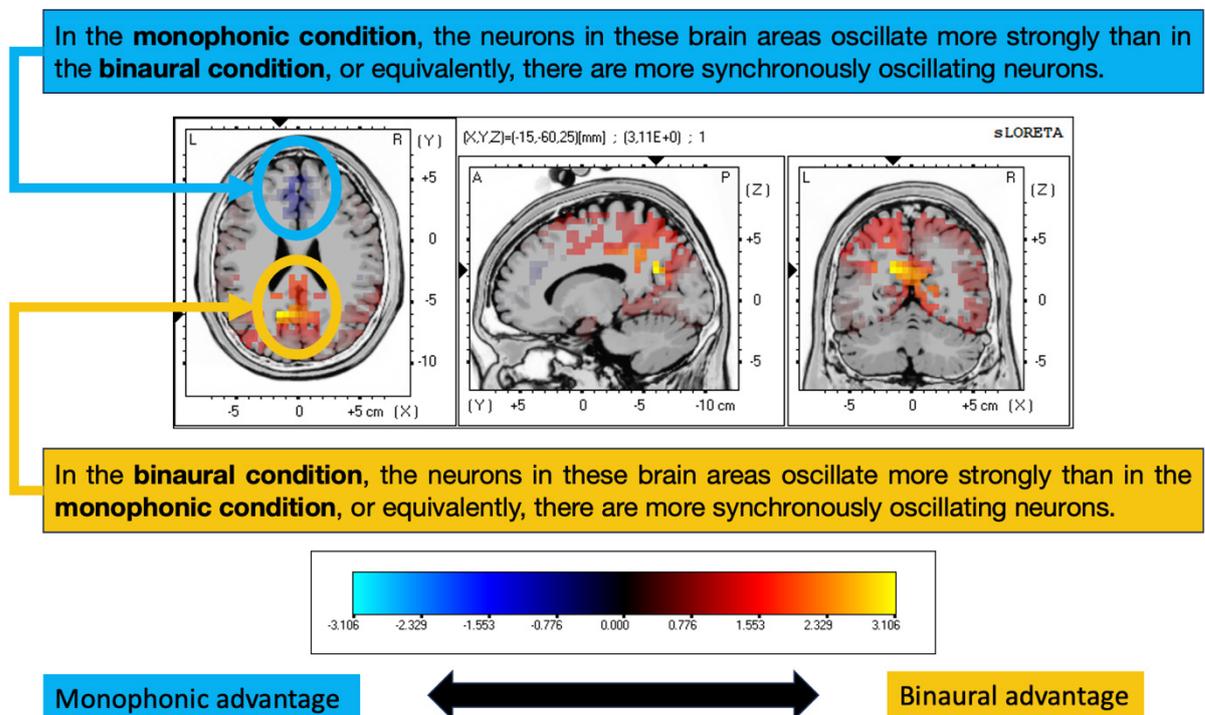
sLORETA for EEG analysis and reporting: the exploratory analysis of brain activations during self-listening to binaural and monophonic auditory stimuli was conducted using sLORETA, allowing the identification of the cortical regions most involved in each listening condition. All reported activations were obtained by comparing participants' baselines, ensuring that the results reflect variations in neural activity induced by the auditory stimuli.

sLORETA software version v20210701 was used for source estimation, cortical distribution analysis, and calculation of activation averages across different brain regions. This software enabled the transformation of the cross-spectral data into files that allowed the evaluation of differences in oscillatory activity between binaural and monophonic listening conditions across all frequency bands: Delta (δ) (0.5–4 Hz); Theta (θ) (4–8 Hz); Alpha 1 (α_1) (8–10 Hz); Alpha 2 (α_2) (10–12 Hz); Beta 2 (β_2) (18–25 Hz); Beta 3 (β_3) (25–30 Hz). Activation averages were calculated directly in sLORETA. For each participant, cross-spectral values in each frequency band were summed across the regions of interest (ROI) and then divided by the total number of analyzed segments, yielding individual activation averages. These individual averages were then integrated to generate the group's overall average, reflecting the combined brain activation of all participants.

The results are presented through images and tables containing Talairach coordinates (X, Y, Z), voxel values, Brodmann areas (BA), cerebral lobes, and the structures most

activated when listening to the stimuli. This information enables a comparative visualization of the different regions recruited under each sound condition. Brain regions highlighted in blue tones represent greater neural activation in the monophonic condition. In comparison, regions in red and yellow tones emphasize a greater neural activation in the binaural condition, as illustrated in Figure 1.

Figure 1 - sLORETA output.



Source: sLORETA/authors (2025)

Description of the image: Figure used as an example of sLORETA output showing the intracranial source differences between the monophonic and binaural conditions.

Data triangulation procedure: to improve understanding of neural dynamics across auditory conditions, a data triangulation process was conducted. This procedure organized cortical

activations by frequency band and stimulus type (binaural and monophonic), identifying convergent and divergent regions. Brain structures were classified according to Brodmann areas, associated lobes, and the frequency bands in which they were activated.

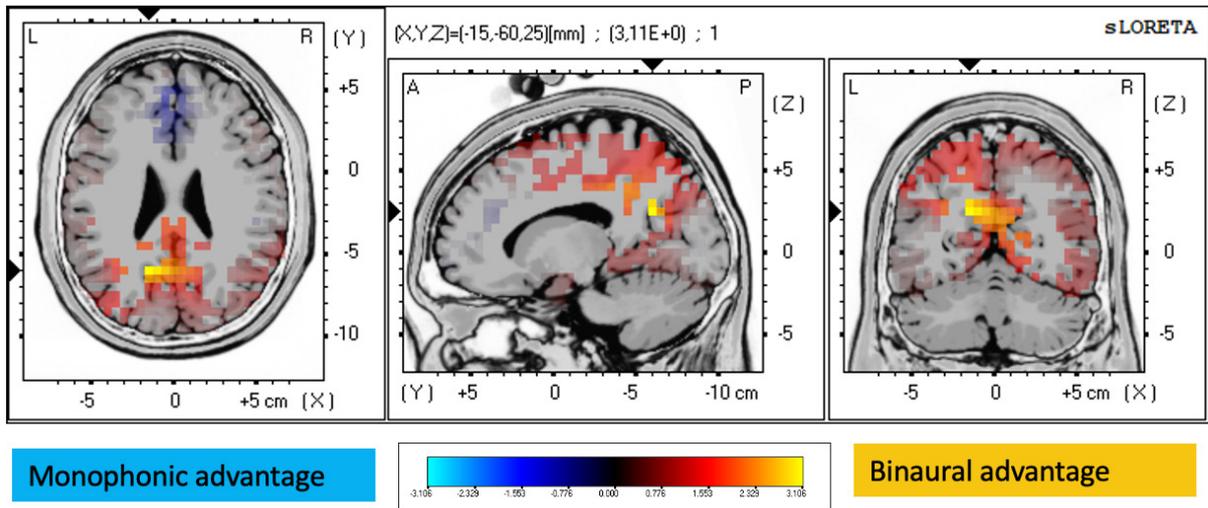
No inferential statistical analyses were conducted due to the exploratory nature of the study and the limited sample size. Visual inspection and descriptive analyses were used to identify trends and inform the direction of future research.

3 Results

The results indicate distinct activation patterns between the binaural and monophonic conditions, suggesting differences in auditory processing mechanisms and neural integration. Specifically, variations were observed in the regions of most significant activation across frequency bands, suggesting that each sound configuration may modulate cortical activity differently in musicians. Below, we present the activation maps generated by sLORETA, along with descriptive tables containing the five central regions activated for each frequency bands—Delta (δ) (0.5–4 Hz), Theta (θ) (4–8 Hz), Alpha 1 ($\alpha 1$) (8–10 Hz), Alpha 2 ($\alpha 2$) (10–12 Hz), Beta 2 ($\beta 2$) (18–25 Hz), and Beta 3 ($\beta 3$) (25–30 Hz)—and self-listening condition (binaural and monophonic).

3.1 Delta band (δ) (0.5–4 Hz).

Figure 2 – Delta band (δ) (0.5–4 Hz) brain scan.



Source: sLORETA/authors (2025).

Description of the image: comparison of delta band (δ) (0,5 - 4 Hz) activations between monophonic and binaural self-listening.

Tabela 2 – Delta band (δ) (0.5–4 Hz) results.

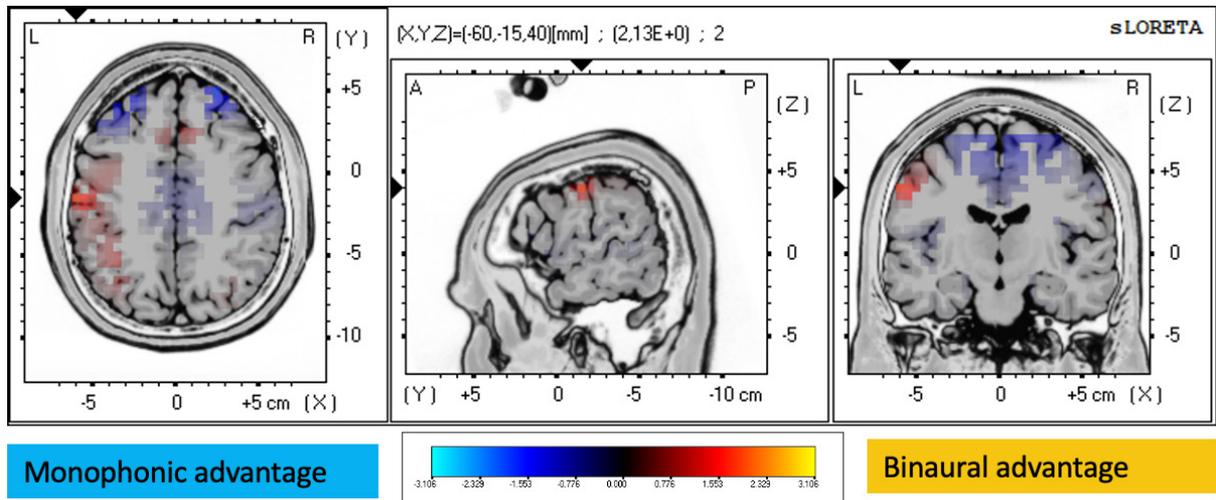
| | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
|--------------------------|--------|--------|---------|-------------|---------------|----------------|----------------------|
| Delta band Monophonic | 45 | -8 | 37 | -1.32080 | 6 | Frontal Lobe | Precentral Gyrus |
| | 0 | 45 | 25 | -1.28187 | 9 | Frontal Lobe | Medial Frontal Gyrus |
| | -5 | 45 | 21 | -1.24312 | 9 | Frontal Lobe | Medial Frontal Gyrus |
| | 45 | -8 | 33 | -1.22889 | 6 | Frontal Lobe | Precentral Gyrus |
| | 45 | -12 | 42 | -1.20153 | 4 | Frontal Lobe | Precentral Gyrus |
| Delta band Binaural | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
| | -15 | -57 | 26 | 3.10566 | 31 | Limbic Lobe | Precuneus |
| | -10 | -57 | 26 | 2.99041 | 31 | Occipital Lobe | Precuneus |
| | -15 | -57 | 30 | 2.92863 | 31 | Limbic Lobe | Precuneus |
| | -50 | -33 | 20 | 2.91854 | 13 | Sub-lobar | Insula |
| -35 | -42 | 34 | 2.85009 | 40 | Parietal Lobe | Sub-Gyral | |

Source: sLORETA/authors (2025).

Description of the image: five brain areas showing the strongest synchronous neuronal activity in the delta band during monophonic and binaural self-listening.

3.2 Theta band (θ) (4–8 Hz)

Figure 3 – Theta band (θ) (4–8 Hz) brain scan.



Source: sLORETA/authors (2025).
 Description of the image: comparison of theta band (θ) (4–8 Hz) activations between monophonic and binaural self-listening.

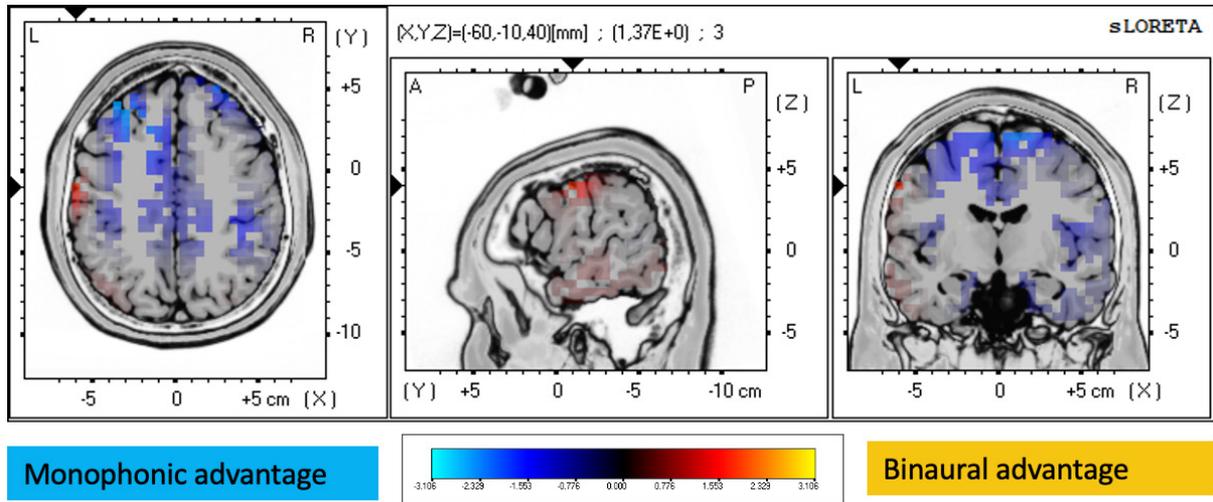
Tabela 3 – Theta band (θ) (4–8 Hz) results.

| | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
|-----------------------|--------|--------|---------|-------------|--------------|------------------|------------------------|
| Theta band Monophonic | -50 | 15 | 8 | -3.62649 | 44 | Frontal Lobe | Precentral Gyrus |
| | -50 | 20 | 4 | -3.50839 | 45 | Frontal Lobe | Inferior Frontal Gyrus |
| | -54 | 19 | -1 | -3.47897 | 47 | Frontal Lobe | Inferior Frontal Gyrus |
| | -50 | 15 | -1 | -3.46982 | 47 | Frontal Lobe | Inferior Frontal Gyrus |
| | -45 | 15 | 8 | -3.04628 | 44 | Frontal Lobe | Precentral Gyrus |
| Theta band Binaural | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
| | -59 | -13 | 37 | 2.13435 | 4 | Frontal Lobe | Precentral Gyrus |
| | -54 | -13 | 37 | 1.93863 | 4 | Frontal Lobe | Precentral Gyrus |
| | -54 | -18 | 38 | 1.80261 | 3 | Parietal Lobe | Postcentral Gyrus |
| | -54 | -13 | 33 | 1.75852 | 4 | Frontal Lobe | Precentral Gyrus |
| -50 | -13 | 37 | 1.58895 | 4 | Frontal Lobe | Precentral Gyrus | |

Source: sLORETA/authors (2025).
 Description of the image: five brain areas showing the strongest synchronous neuronal activity in the theta band during monophonic and binaural self-listening.

3.3 Alpha 1 band (α_1) (8–10 Hz)

Figure 4 – Alpha 1 band (α_1) (8–10 Hz) brain scan.



Source: sLORETA/authors (2025).
 Description of the image: comparison of alpha 1 band (α_1) (8 - 10 Hz) activations between binaural and monophonic self-listening.

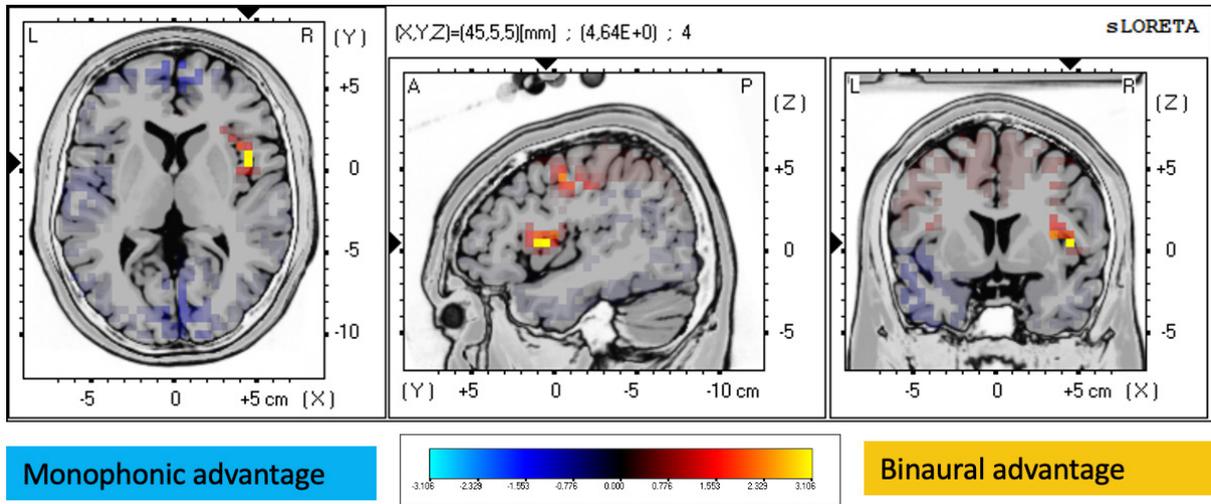
Tabela 4 – Alpha 1 band (α_1) (8–10 Hz) results.

| | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
|----------------------------|---------|---------|---------|-------------|----|--------------|------------------------|
| Alpha band 1 Monophonic | -30 | 36 | 40 | -2.36748E | 8 | Frontal Lobe | Middle Frontal Gyrus |
| | -35 | 45 | 30 | -2.32785E | 9 | Frontal Lobe | Superior Frontal Gyrus |
| | -25 | 36 | 40 | -2.20038E | 8 | Frontal Lobe | Middle Frontal Gyrus |
| | -25 | 31 | 40 | -2.13616E | 8 | Frontal Lobe | Middle Frontal Gyrus |
| | -40 | 45 | 25 | -2.13109E | 10 | Frontal Lobe | Middle Frontal Gyrus |
| | X(TAL), | Y(TAL), | Z(TAL), | Voxel Value | BA | Lobe | Structure |
| Alpha band 1 Binaural | -59 | -8 | 37 | 1.36793E | 6 | Frontal Lobe | Precentral Gyrus |
| | -59 | -13 | 37 | 1.25898E | 4 | Frontal Lobe | Precentral Gyrus |
| | -59 | -13 | 33 | 1.15576E | 4 | Frontal Lobe | Precentral Gyrus |
| | -54 | -13 | 33 | 1.06633E | 4 | Frontal Lobe | Precentral Gyrus |
| | -59 | -18 | 38 | 1.04848E | 4 | Frontal Lobe | Precentral Gyrus |

Source: sLORETA/authors (2025).
 Description of the image: five brain areas showing the strongest synchronous neuronal activity in the alpha 1 band (α_1) (8–10 Hz) during monophonic and binaural self-listening.

3.4 Alpha 2 band (α_2) (10–12 Hz)

Figure 5 – Alpha 2 band (α_2) (10–12 Hz) brain scan.



Source: sLORETA/authors (2025).

Description of the image: comparison of alpha 2 band (β_2) (10 - 12 Hz) activations between binaural and monophonic self-listening.

Tabela 5 – Alpha 2 band (α_2) (10–12 Hz) results.

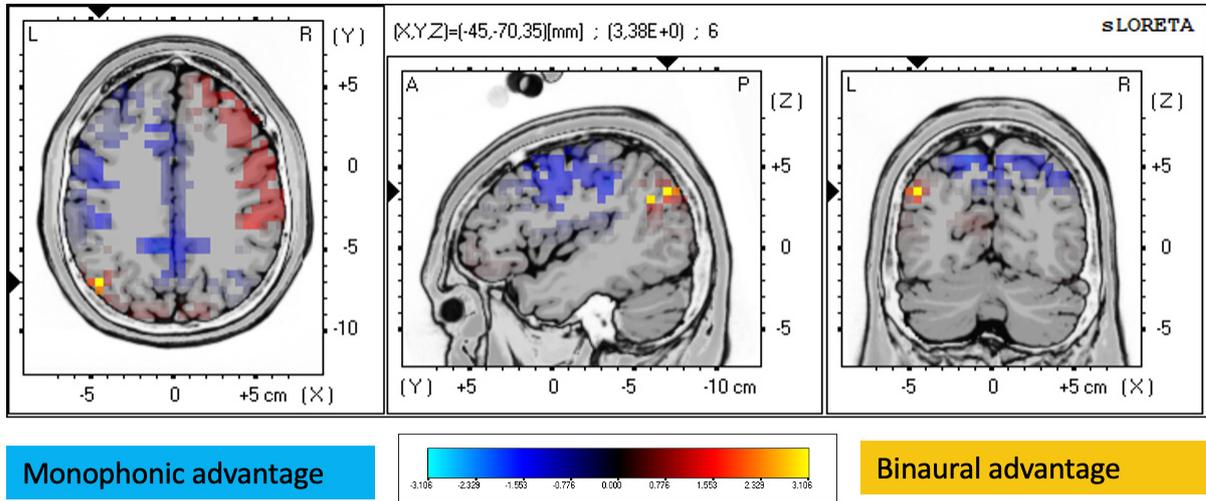
| | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
|----------------------------|--------|--------|---------|-------------|--------------|------------------|----------------------|
| Alpha band 2 Monophonic | 5 | 64 | 11 | -2.37594 | 10 | Frontal Lobe | Medial Frontal Gyrus |
| | 5 | 59 | 11 | -2.35781 | 10 | Frontal Lobe | Medial Frontal Gyrus |
| | 10 | 64 | 11 | -2.32757 | 10 | Frontal Lobe | Medial Frontal Gyrus |
| | 5 | 64 | 15 | -2.32204 | 10 | Frontal Lobe | Medial Frontal Gyrus |
| | 5 | 59 | 6 | -2.29986 | 10 | Frontal Lobe | Medial Frontal Gyrus |
| Alpha band 2 Binaural | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
| | 45 | 5 | 4 | 4.64078 | 13 | Sub-lobar | Insula |
| | 45 | 10 | 4 | 4.54368 | 13 | Sub-lobar | Insula |
| | 40 | 0 | 9 | 3.91919 | 13 | Sub-lobar | Insula |
| | 35 | 5 | 9 | 3.52575 | 13 | Sub-lobar | Insula |
| 45 | -3 | 42 | 3.48467 | 6 | Frontal Lobe | Precentral Gyrus | |

Source: sLORETA/authors (2025).

Description of the image: five brain areas showing the strongest synchronous neuronal activity in the alpha 2 band during monophonic and binaural self-listening.

3.5 Beta 2 band (β_2) (18–25 Hz)

Figure 6 – Beta 2 band (β_2) (18–25 Hz) brain scan.



Source: sLORETA/authors (2025).

Description of the image: comparison of beta 2 band (β_2) (18–25 Hz) activations between binaural and monophonic self-listening.

Tabela 6 – Beta 2 band (β_2) (18–25 Hz) results.

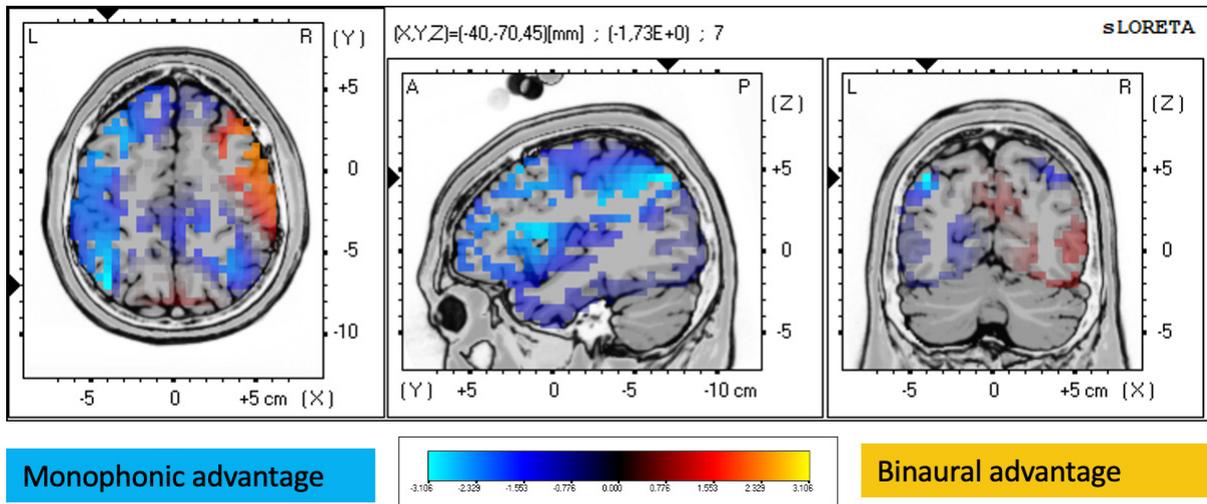
| | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
|---------------------------|--------|--------|--------|-------------|----|---------------|--------------------------|
| Beta band 2 Monophonic | -10 | -56 | 49 | -2.23235 | 7 | Parietal Lobe | Precuneus |
| | -50 | -36 | 48 | -2.22845 | 40 | Parietal Lobe | Inferior Parietal Lobule |
| | -50 | -37 | 43 | -2.22530 | 40 | Parietal Lobe | Inferior Parietal Lobule |
| | -10 | -56 | 53 | -2.17543 | 7 | Parietal Lobe | Precuneus |
| | -15 | -56 | 49 | -2.17462 | 7 | Parietal Lobe | Precuneus |
| Beta band 2 Binaural | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
| | -45 | -66 | 36 | 3.38287 | 39 | Parietal Lobe | Angular Gyrus |
| | -45 | -57 | 30 | 3.23414 | 39 | Temporal Lobe | Superior Temporal Gyrus |
| | -50 | -61 | 40 | 2.59142 | 39 | Parietal Lobe | Inferior Parietal Lobule |
| | -45 | -71 | 36 | 2.57049 | 39 | Parietal Lobe | Angular Gyrus |
| | -40 | -57 | 26 | 2.51595 | 39 | Temporal Lobe | Superior Temporal Gyrus |

Source: sLORETA/authors (2025).

Description of the image: five brain areas showing the strongest synchronous neuronal activity in the beta 2 band during monophonic and binaural self-listening.

3.6 Beta 3 band (β_3) (25–30 Hz)

Figure 7 – Beta 3 band (β_3) (25–30 Hz) brain scan.



Source: sLORETA/authors (2025).

Description of the image: comparison of beta 3 band (β_3) (25–30 Hz) activations between binaural and monophonic self-listening.

Tabela 7 – Beta 3 band (β_3) (25–30 Hz) results.

| | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
|---------------------------|--------|--------|---------|-------------|-----------|---------------|------------------------|
| Beta band 3 Monophonic | -40 | -66 | 45 | -1.72677 | 7 | Parietal Lobe | Inferior Parietal |
| | -40 | -61 | 45 | -1.70036 | 40 | Parietal Lobe | Inferior Parietal |
| | -35 | 6 | 14 | -1.69341 | 13 | Sub-Lobar | Insula |
| | -35 | 6 | 18 | -1.65012 | 13 | Sub-Lobar | Insula |
| | -35 | 10 | 13 | -1.64821 | 13 | Sub-Lobar | Insula |
| Beta band 3 Binaural | X(TAL) | Y(TAL) | Z(TAL) | Voxel Value | BA | Lobe | Structure |
| | 35 | 15 | -1 | 1.59617 | 13 | Sub-Lobar | Insula |
| | 55 | 19 | -1 | 1.58762 | 47 | Sub-Lobar | Extra-Nuclear |
| | 40 | 15 | -1 | 1.57985 | 13 | Sub-Lobar | Insula |
| | 45 | 19 | -1 | 1.57704 | 47 | Frontal Lobe | Inferior Frontal Gyrus |
| 40 | -14 | -5 | 1.57387 | 47 | Sub-Lobar | Insula | |

Source: sLORETA/authors (2025).

Description of the image: five brain areas showing the strongest synchronous neuronal activity in the beta 3 band during monophonic and binaural self-listening.

3.7 Convergent and divergent activations between self-listening stimuli

Triangulation of the data revealed patterns of convergent and divergent cortical activations between binaural and monophonic self-listening conditions. Table 8 systematically

organizes this information, facilitating interpretation and enabling the decoding of the functional specificities associated with each type of self-listening.

Tabela 7 – Cortical structures activated across frequency bands during binaural and monophonic self-listening.

| Structure | Brodmann Area (BA) | Frequency Band | Lobe | Stimulus Type |
|--------------------------|--------------------|----------------|-----------------|---------------|
| Precentral Gyrus | BA 4, 6 | Delta | Frontal | Both |
| Insula | BA 13 | Beta 3 | Sub-lobar | Both |
| Precuneus | BA 31 | Delta | Parietal/Limbic | Binaural |
| Sub-Gyral | BA 40 | Delta | Parietal/Limbic | Binaural |
| Precentral Gyrus | BA 4 | Theta | Frontal | Binaural |
| Postcentral Gyrus | BA 3 | Theta | Parietal | Binaural |
| Precentral Gyrus | BA 4, 6 | Alpha 1 | Frontal | Binaural |
| Insula | BA 13 | Alpha 2 | Sub-lobar | Binaural |
| Precentral Gyrus | BA 6 | Alpha 2 | Frontal | Binaural |
| Superior Temporal Gyrus | BA 39 | Beta 2 | Temporal | Binaural |
| Angular Gyrus | BA 39 | Beta 2 | Parietal | Binaural |
| Inferior Parietal Lobule | BA 39 | Beta 2 | Parietal | Binaural |
| Extra-Nuclear | BA 47 | Beta 3 | Sub-lobar | Binaural |
| Medial Frontal Gyrus | BA 9 | Delta | Frontal | Monophonic |
| Precentral Gyrus | BA 44 | Theta | Frontal | Monophonic |
| Inferior Frontal Gyrus | BA 45, 47 | Theta | Frontal | Monophonic |
| Medial Frontal Gyrus | BA 10, 8 | Alpha 1 | Frontal | Monophonic |
| Superior Frontal Gyrus | BA 9 | Alpha 1 | Frontal | Monophonic |
| Medial Frontal Gyrus | BA 10 | Alpha 2 | Frontal | Monophonic |
| Precuneus | BA 7 | Beta 2 | Parietal/Limbic | Monophonic |
| Inferior Parietal Lobule | BA 40 | Beta 2 | Parietal | Monophonic |
| Inferior Parietal | BA 40, 7 | Beta 3 | Parietal | Monophonic |
| Inferior Frontal Gyrus | BA 47 | Beta 3 | Frontal | Monophonic |

Source: sLORETA/authors (2025).

Description of the image: Cortical structures activated across frequency bands (Delta, Theta, Alpha 1, Alpha 2, Beta 2, Beta 3) during binaural and monophonic self-listening. The type of stimulus indicates whether activation was exclusive to one condition or common to both.

Common activations were identified in the precentral gyrus (BA 4, 6) in the delta band, and in the insula (BA 13) in the beta 3 band. Binaural self-listening showed predominant activations in more widespread regions such as the precuneus (BA 31), insula (BA 13), angular gyrus (BA 39), and superior temporal gyrus (BA 39). In contrast, monophonic self-listening

revealed greater recruitment of frontal regions, including the medial frontal gyrus (BA 9, 10), superior frontal gyrus (BA 9), and inferior frontal gyrus (BA 45, 47). These findings are further detailed and discussed in the following section.

4 Discussion

This exploratory study investigated whether binaural self-listening of a performance, by preserving more realistic spatial characteristics, could elicit neurophysiological responses more closely aligned with actual music performance situations, thereby contributing more effectively to the mental simulation process of music performance than monophonic self-listening. The results provide preliminary evidence supporting this hypothesis, indicating that binaural listening is associated with a more distributed cortical activation pattern that is functionally aligned with motor simulation, sensory integration, and emotional engagement. In contrast, monophonic listening revealed greater involvement of frontal regions associated with cognitive effort and executive control.

Analysis of cortical activations by structure revealed distinct patterns between binaural and monophonic listening stimuli. The precentral gyrus (BA 4 and 6) emerged as one of the regions most responsive to auditory stimuli, with activations distributed across various frequency bands. In the binaural condition, BA 6 was activated in the alpha 1 and alpha 2 bands, suggesting its involvement in motor imagery processes, action planning, and attentional coordination. Studies indicate that reduced alpha power in motor regions during motor imagery tasks — a phenomenon known as event-related desynchronization (ERD) — is widely recognized as an indicator of functional cortical activation (Stecklow; Infantosi; Cagy, 2007; Deiber et al., 2012). Guided auditory tasks have also shown that increased alpha activity may be associated with efficient sensorimotor engagement, especially in immersive contexts such as binaural listening (Zhou et al., 2022). BA 4 was activated in the theta

and alpha 1 bands, a pattern consistent with its role in motor readiness and simulation of voluntary movement. Although classically associated with physical execution, fMRI studies show that BA 4 can also be recruited during imagined motor planning, involving subregions 4a and 4p (Sharma et al., 2008).

Furthermore, EEG recordings show alpha-band desynchronization in the sensorimotor region during motor imagery tasks, indicating functional activation even in the absence of actual movement (Formaggio et al., 2010; Yuan et al., 2010). Conversely, in monophonic conditions, activation of the same gyrus was observed in the delta band, particularly in BA 6, whose role in motor simulation under high cognitive load and in inhibitory control mechanisms is supported by EEG and fMRI research (Piano et al., 2017; Tanaka et al.; Honda et al.; Sadato et al., 2005). BA 44 was recruited explicitly in the theta band, consistent with its involvement in speech motor planning, executive functions, and directed auditory attention (Pardo et al., 1990). The combined presence of delta and theta activity in this region may reflect a greater cognitive effort during the simulation of musical performance under monophonic conditions (Harmony, 2013). In monophonic or spatially co-located conditions, the reduced availability of spatial/binaural cues is associated with increased listening effort and attentional demands during auditory scene analysis, compared with conditions that preserve binaural spatial cues (Rennies; Kidd Junior, 2018; Krueger et al., 2022; Koelewijn et al., 2015; Feuerstein, 1992; Bremen; Middlebrooks, 2013; Litovsky, 2012; Fintor, 2022).

These findings suggest that while binaural listening enhances neural mechanisms related to efficient motor simulation and sensorimotor engagement, monophonic listening requires greater executive resource recruitment and cognitive control during simulation. This implies that binaural self-listening facilitates automatic motor simulation, whereas monophonic listening requires more cognitive effort.

The precuneus was engaged in both conditions, albeit across different frequency bands. During binaural listening, BA 31 was activated in the delta band, associated with consciousness, mental imagery, autobiographical memory, and the functioning of the default mode network (DMN) (Cavanna, 2007). This region also plays a crucial role in integrating internal and external processes, as shown by its functional connectivity with sensory, emotional, and executive networks in highly trained musicians, especially during introspective and self-referential states (Tanaka; Kirino, 2016). Delta band activity in the precuneus is recognized as an electrophysiological marker of conscious states and automatic environmental monitoring (Danielson; Guo; Blumenfeld, 2011; Bonfiglio et al., 2013), reinforcing the idea that binaural listening promotes deep sensory immersion. In the monophonic condition, activation of BA 7 in the beta 2 band was observed, consistent with its role in integrating spatial information and visuomotor coordination, particularly in tasks involving bimanual action planning and spatial cognition (Le et al., 2017). Beta band activity (including beta 2) is also linked to visuomotor attention and motor preparation, especially in imagery and visual-motor control tasks (Baravalle; Rosso; Montani, 2018). Hence, BA 7 activation during monophonic listening may indicate compensatory efforts in spatial attention and motor control to support performance simulation in the absence of natural spatial cues.

The insula (BA 13) was activated in both conditions in the beta 3 band, consistent with its role in transforming auditory signals into emotional experiences, particularly in its anterior portion (Zhang et al., 2019). However, only binaural listening elicited additional activation in the alpha 2 band—a frequency associated with internal engagement during emotional tasks, possibly reflecting heightened sensory immersion (Schreiter; Chmielewski; Beste, 2018). This activation pattern indicates that binaural self-listening supports a more vivid and self-referential simulation of the musical experience.

The inferior frontal gyrus (BA 45, 47), predominantly recruited during monophonic listening, plays a key role in executive functions such as inhibitory control, focused attention, and response regulation under high cognitive load (Penfold et al., 2015). Activity in the theta and beta 3 bands within these areas underscores their involvement in inhibitory modulation (Dippel et al., 2017a, 2017b) and in managing perceptual and response conflicts (Daniel et al., 2023). In the context of monophonic listening — characterized by the absence of natural spatial cues — such conflict may stem from discrepancies between the musician's auditory expectations and the perceived stimulus, requiring greater engagement of the inferior frontal gyrus to preserve the coherence of mental simulation. This executive coordination appears to be mediated by communication between prefrontal regions and the subthalamic nucleus via theta and beta oscillations (Zavala et al., 2018), suggesting that monophonic listening imposes a heavier cognitive burden on sustaining music performance simulation.

The inferior parietal lobule displayed distinct activation profiles across listening conditions, reflecting divergent processing of auditory stimuli in spatial and sensory dimensions. In the binaural condition, activation of BA 39 was noted, encompassing the angular gyrus and the inferior parietal lobule, areas linked to multisensory integration and symbolic spatial representation (Caspers et al., 2008), as well as spatial cognition and manipulation of abstract mental constructs (Seghier, 2013). Such activation may reflect enhanced engagement of the attentional network and the construction of a more accurate spatial representation during self-listening. By contrast, monophonic listening elicited activity in BA 40 and BA 7, associated with reflexive attention and automatic detection of relevant stimuli in peripersonal space (Chambers et al., 2007). These areas are involved in auditory localization and three-dimensional sound processing, including orientation and spatial sound perception (Sato et al., 2011; Lewald et al., 2011).

However, these activations were less consistent — or even negative — than in the binaural condition, possibly indicating a less integrated and spatially impoverished sensory experience (Koizumi; Amano; Komatsu, 2019; Fox et al., 2005).

Studies such as Heine et al. (2021) and Callan et al. (2013) have demonstrated that spatialized sounds generated through HRTFs — by convolution or individual modeling — provoke activation in auditory and parietal areas associated with spatial perception and auditory attention (e.g., posterior superior temporal gyrus, BA 7, insula, and precuneus). The present study builds on this by examining binaural music self-listening, recorded from the performer's auditory perspective. Unlike earlier studies that relied on artificial stimuli and simple auditory tasks, the current protocol employed realistic, complex, and personally meaningful musical content, simulating a performance setting with high ecological validity—an element often absent from conventional auditory perception research (Koelsch, 2011; Almudena et al., 2021).

Although individualized HRTFs were not employed, the study used in situ binaural recordings with microphones attached to participants' ears, thereby preserving the natural spatial and reverberation characteristics of the listening experience. This ecologically grounded method aligns with findings that highlight the engagement of distributed neural systems — including auditory, motor, emotional, and cognitive networks — during music perception (Koelsch, 2011; Zatorre; Chen; Penhune, 2007).

Specifically, the activation of regions such as the precentral and inferior frontal gyri may reflect the motor simulation of performance, supported by evidence showing that passive music listening can recruit motor regions, suggesting internal rehearsal or predictive coding of rhythmic patterns (Zatorre; Chen; Penhune, 2007). The insula and the inferior parietal lobule are also part of a network responsible for sensory integration and spatial attention, previously implicated in music listening,

which involves emotional and multimodal processing (Schaefer; Vlek; Desain, 2011; Zatorre; Chen; Penhune, 2007). Moreover, the involvement of the precuneus and parietal cortices may reflect the integration of sensory, motor, and spatial features into a coherent musical experience, consistent with models of music cognition and memory that emphasize the roles of these structures (Koelsch, 2011; Hyde; Peretz; Zatorre, 2008; Almudena et al., 2021).

Our results, therefore, not only support previous findings on spatial auditory processing using HRTF-based stimuli, but also contribute new evidence that ecologically valid, self-referential musical stimuli, such as binaural recordings of one's own performance, may more fully engage the multisensory and sensorimotor networks involved in music perception, memory encoding, and emotional processing. These findings suggest that binaural self-listening may be integrated into mental practice protocols for musicians, particularly in situations where instrument use is restricted, such as biofeedback and neurofeedback training for music MPA. By enhancing the realism and emotional resonance of auditory imagery, binaural recordings could help musicians engage more deeply in mental simulation, potentially improving performance quality and reducing anxiety levels.

Despite the study's interesting and promising findings, it presents several methodological limitations that should be considered when interpreting the results. The most significant of these concerns the lack of inferential statistical testing of cortical activation data. The analysis was based on visual inspection of distribution images generated by sLORETA and the identification of the top five activation foci per frequency band, without applying quantitative comparisons between conditions. Additionally, while sLORETA is useful for initial estimations of cortical localization, it has limited spatial resolution, particularly for subcortical structures or adjacent cortical regions (Pascual-Marqui, 2002; Vitacco et al., 2002; Mulert et al., 2004). Finally,

although the use of personalized binaural recordings represents an advance in ecological realism, it also imposes a practical limitation: the applicability of the results depends on the feasibility of using recordings captured with each participant's individual HRTF. These limitations underscore the exploratory nature of the study and highlight essential directions for methodological refinement in future research.

The findings suggest that binaural self-listening of music performances promotes a cortical activation pattern more closely aligned with mental performance simulation processes, such as motor imagery, sensory integration, and emotional engagement. In turn, monophonic listening was associated with greater activation in executive control regions, indicating a potentially more cognitively demanding auditory experience. These results highlight the potential of binaural self-listening as a complementary tool for mental practice, particularly for addressing MPA in contexts where instrument use is restricted, such as in biofeedback and neurofeedback training. However, the exploratory nature of the study, marked by the absence of inferential statistical analyses and the limited spatial resolution of sLORETA, underscores the need for methodological refinement. Future research should expand the sample size, apply appropriate statistical tests (e.g., paired t-tests or permutation tests with FDR correction), and incorporate subjective measures such as perceived realism, presence, and attentional effort to better relate cortical responses to conscious experience. Building on the present findings, future studies with larger samples should explicitly compare monophonic, stereo, and binaural listening conditions, with a targeted focus on the specific brain regions identified here, enabling a more precise and hypothesis-driven mapping of their neurophysiological correlates. Additionally, comparisons between different spatialization methods, such as real binaural recordings, synthesized HRTFs, and binaural beats, and the integration of other neuroimaging techniques like fMRI could enhance both the accuracy and applicability of findings.

Overall, this study broadens the scope of personalized spatial sound in musical performance research and opens promising avenues for its use in therapeutic and educational settings.

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Authorship contribution

Nery Borges: Conceptualization; Methodology; Investigation; Data curation; Formal analysis; Writing – original draft; Writing – review & editing; Project administration. Pedro Bem-Haja:

Conceptualization; Methodology; Formal analysis; Validation; Writing – review & editing. Isaac Raimundo: Conceptualization; Data acquisition; Software; Technical support; Methodology; Writing – review & editing. Helena Marinho: Supervision; Methodology; Writing – review & editing. Marcos Vinícius Araújo: Supervision; Methodology; Writing – review & editing. Anabela Pereira: Supervision; Validation; Writing – review & editing. Isabel M. Santos: Supervision; Formal analysis; Data acquisition; Validation; Writing – review & editing.

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