Study of tonotopic brain changes with functional MRI and FDG-PET in a patient with unilateral objective cochlear tinnitus.

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Abbreviations:

BOLD: blood oxygenation level dependant

dB SPL: decibel sound pressure level

EPI: Echo Planar Imaging

FDG-PET: ¹⁸F-fluorodeoxyglucose positron emission tomography

Hz: Hertz

LYSO: lutetium-yttrium-orthosilicate

OHCs: Outer hair cells

OSEM: ordered-subset expectation maximization

PFL: paraflocculus of the cerebellum

SOAEs: Spontaneous otoacoustic emissions

SPM: Statistical parametric mapping

3T fMRI: 3 Tesla functional MRI

7T fMRI: 7 Tesla functional MRI

Abstract:

We studied possible brain changes with functional MRI (fMRI) and fluorodeoxyglucose positron emission tomography (FDG-PET) in a patient with a rare, high-intensity "objective tinnitus" (high-level SOAEs) in the left ear of 10 years duration, with no associated hearing loss. This is the first case of objective cochlear tinnitus to be investigated with functional neuroimaging.

The objective cochlear tinnitus was measured by Spontaneous Otoacoustic Emissions (SOAE) equipment (frequency 9689 Hz, intensity 57 dB SPL) and is clearly audible to anyone standing near the patient. Functional modifications in primary auditory areas and other brain regions were evaluated using 3T and 7T fMRI and FDG-PET.

In the fMRI evaluations, a saturation of the auditory cortex at the tinnitus frequency was observed, but the global cortical tonotopic organization remained intact when compared to the results of fMRI of healthy subjects. The FDG-PET showed no evidence of an increase or decrease of activity in the auditory cortices or in the limbic system as compared to normal subjects.

In this patient with high-intensity objective cochlear tinnitus, fMRI and FDG-PET showed no significant brain reorganization in auditory areas and/or in the limbic system, as reported in the literature in patients with chronic subjective tinnitus.

Keywords: objective tinnitus, human neuroimaging, auditory cortex, tonotopy

1. Introduction

Tinnitus is characterized by the perception of auditory signals in the absence of any external sound source. It affects about 10 to 15% of the population (Rauschecker, et al., 2010), and in 1-3% it adversely affects quality of life. There are two types of tinnitus: objective and subjective. Subjective tinnitus is more common and may contribute to depression, insomnia and anxiety. The pathophysiology of subjective tinnitus is still poorly understood, but is most often associated with sensorineural

hearing loss of various origins. As a consequence of damage to the peripheral auditory system, reduced auditory input reaches the central auditory neurons within the affected frequency range, which in turn leads to compensatory changes in the central auditory system. Therefore, subjective tinnitus is a phantom perception of sound that may be due to aberrant plastic reorganization within the auditory centers of the brain. There is currently no standard treatment for the management of subjective tinnitus, however behavioral therapies are considered the most effective. Earlier studies of human brain imaging using fMRI in patients with subjective tinnitus have suggested reorganization in the auditory cortex (Rauschecker, et al., 2010; Melcher, et al., 2000; Lockwood, et al., 1998; Plewnia, et al., 2007, but see Langers et al., 2012). Furthermore, stress and emotion can modulate tinnitus with brain imaging studies showing functional and anatomical differences in the limbic system between tinnitus patients and controls (Rauschecker, et al., 2010).

Objective tinnitus is much less common and corresponds to a physical noise produced in the body and is audible to others. The etiology is most often vascular and, more rarely, is related to other causes such as palatal myoclonus, myoclonus of the tensor tympani, tubal patency or cochlear damage. Objective tinnitus of cochlear origin is due to a highly localized anomaly of the basilar membrane at high frequencies with spontaneous activity of the outer hair cells (OHCs) and it can be measured by Spontaneous Otoacoustic emissions (SOAEs). Some authors have described the correlation between tinnitus pitch and SOAEs frequency (Prasher, et al., 2001; Roberts, et al., 2010).

This article considers the case of a 30 year-old man who presented with a rare left ear objective cochlear tinnitus, of over 10 years durations, which is clearly audible to others standing near the patients as a continuous high-pitched tone. Using functional

magnetic resonance imaging (fMRI) and ^{18}F -fluorodeoxyglucose positron emission tomography (FDG-PET), we investigated possible reorganization of the auditory cortex and limbic system in this patient, as compared to healthy control subjects. The results of three experiments (3T fMRI, 7T fMRI, and FDG-PET) showed no large-scale reorganization of auditory brain areas, revealing a case of overall resiliency to pathological sound exposure.

2. Materials and Methods

2.1. Case report

The patient in this study was a 30-year-old male, in good general health, with no previous history of acoustic trauma, presenting with a left continuous objective tinnitus of over 10 years duration. The tinnitus was non-pulsatile and of high intensity and frequency. The patient was bothered by the symptom and looked for possible treatment. Validated French versions of the Subjective Tinnitus Severity Scale (STSS) (Meric, et al., 1996), the Hospital Anxiety and Depression Scale (HADS) (Lepine, et al., 1985), and the Tinnitus Handicap Inventory (THI) (Ghulyan-Bédikian, et al., 2010), were used to evaluate the impact of the tinnitus. On the STSS, the patient's score was 8/16 indicating a moderate tinnitus discomfort. On the HADS, the global score was 23/42 revealing moderate anxiety (14/21) and slight depression (9/21) symptoms. The THI score was 74 points indicating a severe handicap (severe is more than 58 points). Finally, using a Visual Analogic Scale (VAS), the patient evaluated at 9/10 the intensity and suffering of his tinnitus.

The otoscopy showed normal tympanic membranes bilaterally. Weber's test was central and Rinne's test was positive on both sides. CT-scan of the temporal bone followed by a cerebral and angio-MRI failed to reveal any abnormality.

Otherwise, the patient underwent a complete neurovestibular evaluation including caloric testing, video-head impulse test and videonystagmography, which were normal.

2.2. Audiological assessment

2.2.1. Audiometry and Impedancemetry

Pure tone audiogram was normal on both sides in all frequencies, except for a rapid drop on the left side up to 50 dB at 8 kHz, probably due to the tinnitus interfering with hearing thresholds. (*Fig. 1*)

Impedancemetry (objective test of middle-ear function evaluating the mobility of the tympanic membrane and ossicular chain) was normal (type A) on both sides.

2.2.2. Otoacoustic emissions recording

Otoacoustic emissions are produced by the outer hair cells activity in the cochlea, either spontaneously (Spontaneous Otoacoustic Emissions – SOAEs) or in response to an acoustic stimulus (Transitory Evoked Otoacoustic Emissions - TEOAEs) and can be recorded with an intra-auricular probe. To measure the otoacoustic emissions, we used the Madsen Celesta 503 device, Madsen Celesta logiciel, NOHA system 3.6, GN Otometrics. SOAEs were recorded bilaterally from 125 to 10,000 Hz and TEOAEs from 1000 to 4000 Hz. In the right ear, physiologic SOAEs were present with small peaks of 8 dB SPL recorded at 5200 Hz and 7500 Hz. In the left ear, pathological SOAEs were present with a highly abnormal significant peak of 57 dB SPL at 9689 Hz, producing an audible sound (*Fig. 1*).

2.2.3. Functional neuroimaging

To test for possible brain reorganization, we performed three neuroimaging experiments with the patient: (1) whole-brain assessment of response to sound

stimulation at the tinnitus frequency with 3T-fMRI, (2) high-resolution assessment of the tonotopic organization of the primary auditory cortex with 7T fMRI, (3) evaluation of whole-brain metabolic activity with FDG-PET.

2.3. 3 Tesla fMRI

We looked for changes in the activity in the auditory cortex bilaterally and in the limbic system by comparing sound stimulation at the SOAEs frequency with rest and with stimulation two octaves lower in a silent event-related design protocol (for details see [Maeder, et al. 2001](#)).

Scanning methods and analysis:

The MRI protocol was performed on a Siemens 3T Tim Trio scanner and included a sagittal T1-weighted gradient-echo sequence (MPRAGE, 160 contiguous slices, 1mm isotropic voxel, TR=2300 ms, TE=2.98 ms, FoV=256 mm) and the fMRI acquisition. Functional scans were acquired with an EPI sequence (TR=1500 ms, TE=30 ms, flip angle=90°, FoV=256 mm). The 28 axial slices (matrix size 128*128 with 4 mm slice thickness) were aligned with the anterior commissure-posterior commissure line. During the experiment, we collected 20 volumes for each condition. MRI data were pre-processed and analyzed using Statistical Parametric Mapping (SPM8, Wellcome Department of Imaging Neuroscience, London, England; <http://www.fil.ion.ucl.ac.uk/spm/software/spm8/>). Functional images were corrected for motion along the experiment by-means of rigid-body transformations and then co-registered to the high-resolution T1w acquisitions. The anatomical images were then normalized to the MNI T1 template and the normalization parameters were applied to the functional images, which were finally smoothed with a 6-mm Gaussian kernel. All

the pre-processing steps mentioned minimized the non task-related variability. Single-subject statistics were performed according to the General Linear Model and contrasts of interest were thresholded for peak height at $p=.001$ (uncorrected), with an extent threshold (k) of 50 voxels.

Sound stimulation methods:

During the active condition, the patient received binaural stimulation (on/off) for 5 seconds followed by EPI acquisition (silent event-related paradigm, [Hall, et al., 1999](#); [Maeder, et al., 2001](#)) every 15 seconds. The 5 sec stimulation included 16 pulsed sounds (pure tone bursts of 150ms, the amplitude multiplied with half sine-wave function) either at the SOAEs frequency (9689 Hz, 90 dB SPL) or two octaves below (2422 Hz, 75 dB SPL, for a perception of similar loudness to the SOAEs frequency). The tone stimuli were sampled at 44.1 kHz and were delivered by MR-compatible earplug headphones (AudioSystem, Nordic NeuroLab Bergen, Norway). The 5 sec stimulation was repeated 5 times for each condition and followed by five rest acquisitions; this cycle was repeated four times.

The stimulus was not masked by scanner noise. During fMRI, following the sparse sampling technique (Hall, et al. 1999; Edmister, et al., 1999), the stimulus was not presented during the scanner noise and the tinnitus could be heard normally. The use of stimulation earphones did not alter the perception of the tinnitus.

2.4. 7 Tesla fMRI

High-resolution tonotopic maps of the primary auditory cortex were obtained with functional 7T MRI. Normally, neurons of primary auditory cortex are arranged in spatial gradients according to their preferred sound frequencies from low to high.

Previous studies from our laboratory (Da Costa, et al., 2011, Da Costa, et al., 2013) have demonstrated the consistency with which we can measure cortical tonotopic maps with high-resolution 7T imaging in human subjects with normal hearing. The human primary auditory cortex is situated bilaterally on the transverse temporal plane centered on Heschl's gyrus. Moving from the posterior to the anterior end, we normally observe two complete tonotopic gradients ("high to low" followed by "low to high") that correspond to the primary cortical regions A1 and R (*Fig. 2*). Both of these regions correspond to the primary core region and this organization is largely homologous to that observed in non-human primates (Saenz, et al., 2014).

Scanning methods

Blood oxygenation level-dependent (BOLD) functional imaging was performed with an actively shielded 7T Siemens MAGNETOM scanner (Siemens Medical Solutions) located at the Centre d'Imagerie BioMedicale (CIBM) in Lausanne, Switzerland. fMRI data were acquired using an eight-channel head volume rf-coil (RAPID Biomedical) and an EPI pulse sequence with sinusoidal readout (1.5 × 1.5 mm in-plane resolution, slice thickness = 1.5 mm, TR = 2000 ms, TE = 25 ms, flip angle = 47°, slice gap = 0.07 mm, matrix size = 148 × 148, field of view 222 × 222, 30 oblique slices covering the superior temporal plane. The first three EPI images of each BOLD sequence were discarded).

BrainVoyager QX software v2.3 was used for standard fMRI data analysis. The tonotopic maps follow the convoluted cortical surface and are most easily viewed on cortical surface renderings. Cortical surface meshes were generated for the patient based on his own high-resolution 3D anatomical brain scan (1 × 1 × 1 mm, MPR2AGE T1-weighted sequence).

Sound stimulation methods

The tonotopic mapping stimulus follows a “phase-encoded” mapping paradigm. Also known as the “travelling wave” technique, this paradigm has been shown to be highly efficient for mapping visual retinotopy and auditory tonotopy (Engel et al., 1994; Sereno et al., 1995; Talavage, et al., 2004; Engel, 2012). Briefly, the mapping stimulus is designed to generate a wave of response across the tonotopic maps of the cortex, peaking earliest at low frequency endpoints and progressively later in parts of the map preferring higher sound frequencies. Hence, the time-to-peak of the response (i.e. the response phase) reveals the preferred frequency of each responsive voxel.

The auditory stimuli were pure tones of 16 distinct frequencies presented in increasing order from low to high: 88, 125, 177, 250, 354, 500, 707, 1000, 1414, 2000, 2828, 4000, 5657, 8000, 11,314, and 16,000 Hz (half-octave steps). The transition from low to high frequencies occurred over 32 seconds (2 second presentation of each frequency) followed by a 4 second of silence. This 36 second period was cycled 15 times during a continuous 9 minute scan run. The tone stimuli were generated with a sampling rate of 44.1 kHz and were delivered via MRI-compatible headphones (AudioSystem, Nordic NeuroLab). The sound intensity of each frequency was chosen between 82-97 dB SPL to equate perceived loudness according to standard equal-loudness curves (ISO226, 85 phon). To minimize exposure to scanner noise, the patient (and controls) wore earplugs during the experiment which reduced background noise by about 24dB. This means that a significant amount of scanner noise remains, of which it is difficult to assess the impact. However, it is controlled for in the sense that the background noise is the same for both the patient and the control subjects. According to the patient's self-

report, the tinnitus remained hearable during the scanning session. Subjects (the patient and 5 normal hearing control subjects) were instructed to keep eyes closed during scanning. Linear cross-correlation analysis was used to determine the temporal delay that best fit the measured fMRI response time course of each voxel and to assign a corresponding best frequency. Briefly, the measured fMRI time courses were correlated with model time courses of 16 different delays corresponding to each sound frequency, and frequency-preferences were assigned according to the corresponding delay of highest correlation value. Model time courses were constructed by convolution with a standard two gamma hemodynamic response function (HRF) assuming a 5 sec time to response-peak in the BOLD response (Glover, 1999). Response amplitudes (in % fMRI signal change) were determined as the maximal signal change of voxels assigned to each sound frequency, as in Da Costa et al., 2015. Response amplitudes were compared between the patient and controls.

2.5. FDG-PET

A fasting period of more than 6 hours was respected by the patient before the PET study. The patient received an intravenous injection of 200 MBq of ^{18}F -FDG after resting comfortably in a quiet, dimmed room. He was instructed not to speak or move. The patient wore earplugs and phonic isolation headphones during the initial resting phase (20 minutes) before radiotracer injection and PET acquisition starting immediately after. A PET/CT scanner (Discovery-690 time-of-flight scanner. GE Healthcare, Milwaukee, WI) combining lutetium-yttrium-orthosilicate (LYSO) block detectors with a 64-slice CT scanner was used for image acquisition and 3D iterative reconstruction was performed using ordered-subset expectation maximization

(OSEM) using a 256x256 pixels matrix, with a final image resolution of about 4–6 mm FWHM. The PET data was analyzed with the 3D-Stereotactic Surface Projections Neurostat Software (3D-SSP, Department of Radiology, University of Washington, Seattle, WA, USA, <http://128.208.140.75/~Download/>). A subsequent analysis compared to a database of 20 healthy persons of the same mean age using SPM8 with a p-value < 0.001, corrected for cluster size (Guedj E, et al., 2010).

3. Results

In the patient, we identified pathological SOAEs on the left ear with a significant peak of 57 dB SPL recorded at 9689 Hz. This highly abnormal spontaneous activity of OHCs generates a continuous tonal tinnitus at 57 dB SPL that is clearly audible to others standing near the patient.

First, we looked for brain reorganization in the auditory pathways and the limbic system with 3T fMRI. Results did not reveal any activation of primary auditory cortex when comparing stimulation at the SOAE's frequency with rest. There was no asymmetry of the response to the stimulus at the tinnitus frequency since no response at all could be obtained. The comparison of the lower frequency (two octaves lower, see Method for details) with the SOAEs frequency showed bilateral activation of the middle Heschl's gyrus with a slight predominance in the left hemisphere (*Fig. 3*). No activation in the limbic areas was observed.

Secondly, we tested using 7T fMRI for a change in the tonotopic maps of primary auditory cortex, with the hypothesis that a long-term noise at the same frequency could create a change in the auditory area. We observed that tonotopic mapping results of the patient were in the normal range (*Fig. 4a*) based on qualitative comparison to 20 individual hemispheres of healthy young adults without tinnitus (Da

Costa, et al., 2011). In both left and right brain hemispheres of the patient, mappings of sound frequency preference (“high-to-low” followed by “low-to-high”) corresponding to primary auditory regions A1 and R appeared normal (see Da Costa, et al., 2011). These tonotopic maps appeared normal in terms of location (centered on Heschl’s gyrus of the temporal lobe of each hemisphere) and orientation (overall posterior-to-anterior). *Fig. 4b* displays the preferred frequency as a function of distance along the cortical surface in each hemisphere. Note the distinct “V-shape” indicative of the two continuous and gradual tonotopic transitions. We did not observe a gross overrepresentation of sound frequencies in any frequency range. The x,y,z Talairach coordinates of the patient’s estimated left and right map endpoints (left: [-39 -22 5] , [-45 -25 3]; right: [40 -21 6], [46 -29 12]) were in the range of five normal-hearing control subjects (mean±sd; left: [-38.8±4.0 -21.4±2.7 3.0±6.8], [-44.0± 5.6 -27.2±4.8 3.8±3.3]; right: [41.2±4.1 -18.2±4.7 4.4±3.8] , [49.0±4.7 -28.2±6.2 9.2±4.1]). Figure 5 shows response amplitudes to each sound frequency in the patient compared to the five normal-hearing control subjects. The response amplitudes of the patient were within the range of the control subjects and were not significantly different from controls at any frequency (*t-tests*, $p>0.05$). Finally, we investigated for evidence of hypermetabolism in the brain with the FDG-PET. This examination revealed no abnormalities. Images were compared to a database of normal, age-matched control subjects using the 3D-Stereotactic surface projections software 3DSSP Neurostat, which showed no significant abnormalities. Furthermore, The PET data were subsequently compared to a database of 20 healthy persons of the same mean age using SPM8 and $p\text{-value} < 0.001$, corrected for cluster size (Guedj E, et al., 2010). This analysis did not further identify any area of abnormally increased or decreased glucose metabolism.

4. Discussion

Spontaneous otoacoustic emissions (SOAEs) are thought to arise as a response to random perturbations in cochlear mechanisms due to irregularities in the OHC arrangement or due to a highly localized anomaly of the basilar membrane at high frequencies (Prasher, et al., 2001). Small peaks across the frequency spectrum in the cochlea (generally not exceeding 20 dB SPL) can be recorded, inconsistently, in normal hearing subjects. Very rarely and exceptionally, more severe SOAEs can be recorded as audible sounds classifiable as objective non-pulsatile tinnitus. Only nine cases of objective tinnitus measured by SOAE equipment have been previously described in the literature (Glanville, et al., 1971; Huizing, et al., 1973; Mathis, et al., 1991; Yamamoto, et al., 1987), but the tinnitus frequency and loudness were lower than in our patient. In Glanville et al., there was a familial history of high frequency tinnitus and the authors suggested a genetic defect in the cochlea associated to a venous abnormalities at the base of the skull. The others studies described 6 people with high-intensity SOAEs measured in normally hearing ears and emission failed to synchronize to external click stimuli (TEOAEs). They hypothesized that high-intensity SOAEs are associated with damage to the cochlea. In these studies, the associated hearing loss observed in five patients was explained by a masking effect due to the high intensity SOAEs.

Here we studied a rare case of objective tinnitus associated with clearly audible SOAEs. Given the continuous tonal quality of our patient's objective tinnitus, we hypothesize that it is generated by activity of the outer hair cells, in contrast to myogenic or vascular origins which produce pulsatile, non-tonal sounds. In the cochlea, only outer hair cells are contractile and would appear to be capable of

emitting pure tone sounds. To our knowledge, this is the first case report of objective tinnitus to be studied with neuroimaging. We considered the possibility of changes in auditory brain centers, particularly in the cortex and in the limbic system, based on the findings of previous studies of auditory plasticity and of subjective tinnitus in both humans and animal models.

Firstly, animal studies of auditory plasticity have shown that repeated exposure to a particular sound frequency can alter the tonotopic maps of auditory cortex, resulting in overrepresentation of that frequency (Recanzone, et al., 1993). However, the map reorganization depended on pairing the sounds with an attentive task requiring the animal to focus on the frequency content (Recanzone, et al., 1993; Polley, et al., 2006). Tonotopic map alterations may also be possible following long-term passive sound exposure (Noreña, et al. 2006). In our patient, the duration of sound exposure largely exceeds those studies and the specific attentional state afforded to the sound is unknown.

Second, previous studies of subjective tinnitus in both animal models and humans report a number of associated changes in the auditory cortex. Subjective tinnitus is often associated with sensorineural hearing loss. Nerve deafferentation due to the destruction of hair cells induces changes in the central auditory pathways leading to the perception of a phantom tinnitus (Rauschecker, et al., 1999). Usually, the pitch of the tinnitus corresponds to the hearing loss range (Roberts, et al., 2011; Mühlau, et al., 2006; Melcher, et al., 2000, 2009). Animal models of subjective tinnitus consistently show elevated responses throughout the central auditory system including elevated spontaneous and stimulus driven activity, increased bursting and neural synchrony, as well as down-regulation of inhibition (Noreña, et al., 2003; Mulders, et al., 2009; Seki, et al., 2003). Others have reported tonotopic changes

including expansion of lesion-edge frequency areas (Seki, et al. 2003, Noreña, et al., 2005) or more global distortion pervading the entire tonotopic map organization (Engineer, et al., 2011; Yang, et al., 2011). Earlier studies of human brain imaging with fMRI and FDG-PET have shown abnormal responses in central auditory brain regions in patients suffering from chronic subjective tinnitus (Rauschecker, et al., 2010; Melcher, et al., 2000; Lockwood, et al., 1998; Plewnia, et al., 2007) with some suggesting cortical tonotopic map distortion (Muhlnickel, et al, 1998; Wienbruch, et al., 2006). However, in most studies of subjective tinnitus it is difficult to disentangle the effects of tinnitus from more general consequences of hearing loss. Notably, Langers, et al., 2012 tested subjective tinnitus patients with only minimal hearing loss and, like us, reported a lack of macroscopic changes in the tonotopic maps. Finally, neuroimaging studies have also described changes in the limbic-related brain areas in subjective tinnitus patients, including alterations in some parts of the limbic networks and significant grey matter decreases in the right inferior colliculus and in the left hippocampus, when compared to controls (Lockwood, et al., 1998; Mühlau, et al., 2006; Landgrebe, et al., 2009; Schlee, et al., 2009). Leaver, et al., 2011, suggested that interactions between the limbic corticostriatal network and primary auditory cortex may be key to understanding the pathophysiology of chronic subjective tinnitus. This interaction may be an important gateway to centers mediating emotional control and memory. In addition, the limbic system may participate in the suppression of the tinnitus signal. If this “noise cancellation system” works properly, the tinnitus is transient (Rauschecker, et al., 2010). If this mechanism fails, patients become aware of their tinnitus. Recently, functional imaging in rats with tinnitus showed evidence of elevated neural activity in the paraflocculus of the

cerebellum (PFL), which was not observed in normal rats. This suggests that plastic changes in the PFL may also serve as a tinnitus generator (Bauer, et al., 2013).

Based on these observations in animal models and in patients with subjective tinnitus, we investigated whether our objective tinnitus patient would have aberrant brain reorganization in the auditory cortex, reorganization in the tonotopy of the auditory cortex and/or activation in non-auditory areas (i.e. limbic and para-limbic areas), which would reflect an emotional reaction to tinnitus. Functional imaging and FDG-PET in this patient failed to reveal any major changes. The 3T fMRI experiment showed no activation in the primary auditory cortex when comparing stimulation at the SOAEs frequency with rest, but revealed bilateral activation of the Heschl's gyrus when comparing the SOAEs frequency with lower frequencies. This observation is compatible with a saturation of the measurable auditory cortex response at the tinnitus frequency due to continuous SOAE stimulation. Saturation effects have been previously described in fMRI responses in auditory cortex (Talavage and Edmister, 2004). Except for this probable saturation effect in the primary auditory cortex at the SOAEs frequency, neither gross reorganization of the central auditory system nor changes in the limbic-related brain areas were observed.

Testing with 7T fMRI revealed overall normal tonotopic maps of primary auditory cortex in terms of location and orientation as shown in *Figure 4*. While these findings do not exclude the possibility of finer scale reorganization, they do suggest that the global tonotopic organization of the patient's primary auditory cortex remained intact. It should be noted that a methodological limitation in our tonotopic mapping is that that sound frequencies were presented only in increasing order. Previous applications of phase-encoded mapping emphasize the importance of presenting mapping stimuli also in the reverse direction, to confirm responses do not depend on

stimulus order (Talavage, et al., 2004). In particular, responses at map endpoints could be influenced by sound offsets and onsets, as opposed to strict frequency-dependence. In our previous tonotopic mapping study, both stimulus directions were applied and we observed that the two major gradients of primary auditory cortex were largely stable, irrespective of stimulus order (Da Costa et al., 2013, Fig. 3). Future tonotopic mapping experiments should ideally apply both stimulus directions each time.

The FDG-PET examination did not identify any abnormal metabolic activity in the auditory cortex and limbic areas, also suggested an absence of brain changes in this patient. A possible limitation of FDG-PET imaging is its intrinsic spatial resolution (4-6 mm), where low-intensity metabolic changes in small-sized brain structures could be missed due to partial volume effects. The difference in findings between our patient and those with subjective tinnitus may be due to the fact that objective cochlear tinnitus is peripheral in origin, producing a continuous neural excitation from the labyrinth, while subjective tinnitus might be maintained by cortical reorganization as a consequence of sensorineural hearing loss.

5. Conclusions

Functional and metabolic imaging in our patient with chronic cochlear objective tinnitus showed no gross distortions in auditory cortical organization and/or in limbic system responses, as observed in patients with chronic subjective tinnitus, despite the presence of chronic objective tinnitus of over 10 years duration. Subjective tinnitus seems to be maintained by cortical reorganization as a consequence of

sensorineural hearing loss. On the contrary, objective cochlear tinnitus is peripheral in origin and, while finer-scale neural changes may exist, the results of our three experiments suggest an overall resilience of the central auditory system, despite long-term pathological sound exposure.

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Figures

Fig. 1. Pure tone audiogram: Normal hearing on both sides except for a rapid fall in the very high frequency range on the left ear up to 50 dB at 8kHz (Masking by the high-level SOAE is an explanation).

Spontaneous otoacoustic emissions (SOAEs): left ear shows a pathological peak of 57 dB SPL at 9689 Hz (arrow) corresponding to the audible tinnitus.

Fig. 2. Tonotopic mapping in auditory cortex with 7T fMRI

(a) Sound stimuli were pure tone bursts presented in cycled progressions from low to high frequencies: 88 to 16,000 Hz in half-octave steps. Each 32 s progression from low to high (red-to-blue color scale) was followed by a 4 s stimulus pause. Sound stimuli were designed to induce a traveling wave of response across cortical tonotopic maps: fMRI responses peak sooner in map regions preferring low frequencies and progressively later in regions preferring higher frequencies.

(b and c) Resulting color-coded frequency maps are projected onto the subject's cortical surface meshes, as shown here in a normal hearing control subject.

Surfaces were minimally inflated to expose the auditory cortex on the temporal plane (adapted from Da Costa, et al., 2011).

Fig. 3. 3T fMRI results. Image represents clusters more active during stimulation at 2422 Hz (two octaves lower than SOAE frequency) than at 9689 Hz (SOAE frequency). The slice is located at z=12mm in MNI space. Map is thresholded at $p < 0.001$, $k = 20$ voxels. Colorbar represents T-values.

Fig. 4. (a) Tonotopic maps in the patient with objective tinnitus: tonotopic maps sound frequency preference (“high-to-low” followed by “low-to-high”) corresponding

to primary auditory regions A1 and R were observed in each brain hemisphere. These tonotopic maps appeared normal in terms of location (centered on Heschl's gyrus on the temporal plane of each hemisphere) and orientation (overall posterior-to-anterior), suggesting that the global tonotopic organisation of the patient remains intact.

(b) Plots of preferred frequency within a posterior-to-anterior strip across the surface Heschl's gyrus in right and left hemispheres, respectively: the distinct "V-shape" shows two continuous and gradual tonotopic transitions in each hemisphere corresponding to regions A1 and R.

Fig. 5. Plot of response amplitudes (in % fMRI signal change) to each sound frequency in the patient compared to normal-hearing control subjects (no difference at any frequency, $p > 0.05$). Overall, the results suggest that the global tonotopic organization of the patient remains within the normal range.

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