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Research Article

Functional Near-Infrared Spectroscopy to Probe State- and Trait-Like Conditions in Chronic Tinnitus: A Proof-of-Principle Study

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Objective. Several neuroscience tools showed the involvement of auditory cortex in chronic tinnitus. In this proof-of-principle study we probed the capability of functional near-infrared spectroscopy (fNIRS) for the measurement of brain oxygenation in auditory cortex in dependence from chronic tinnitus and from intervention with transcranial magnetic stimulation. **Methods.** Twenty-three patients received continuous theta burst stimulation over the left primary auditory cortex in a randomized sham-controlled neuronavigated trial (verum = 12; placebo = 11). Before and after treatment, sound-evoked brain oxygenation in temporal areas was measured with fNIRS. Brain oxygenation was measured once in healthy controls ($n = 12$). **Results.** Sound-evoked activity in right temporal areas was increased in the patients in contrast to healthy controls. Left-sided temporal activity under the stimulated area changed over the course of the trial; high baseline oxygenation was reduced and vice versa. **Conclusions.** By demonstrating that rTMS interacts with auditory evoked brain activity, our results confirm earlier electrophysiological findings and indicate the sensitivity of fNIRS for detecting rTMS induced changes in brain activity. Moreover, our findings of trait- and state-related oxygenation changes indicate the potential of fNIRS for the investigation of tinnitus pathophysiology and treatment response.

1. Introduction

Chronic tinnitus is a highly prevalent condition (5–15%) [1] with no available effective therapy to cure this condition, that is, to turn off the tinnitus. Today, it is evident that chronic tinnitus is related to excitatory-inhibitory dysbalance of neural activity with increased spontaneous activity along the auditory pathway [2, 3]. Subsequent tonotopic reorganisation

in the auditory cortex was also detected in animal models of tinnitus [4]. In humans, divergent findings are evident [5, 6].

In humans, increased activity in the auditory cortex is discussed as neuronal correlate of tinnitus [7, 8]. Early positron-emission tomography (PET) studies showed hints for a left-lateralization of activity of the primary auditory cortex in tinnitus [9]. In most cases longitudinal designs with modulation of the perception of tinnitus by lidocaine injection

or somatic manoeuvres were used. A recent meta-analysis affirmed increased activity in the left primary auditory cortex and showed also increased activity in bilateral auditory cortex [10]. A more recent study contrasting patients and controls [11] challenged this concept by providing evidence that left-lateralization is independent from tinnitus.

In contrast to PET which measures brain metabolism over several minutes, functional magnetic resonance imaging (fMRI) is able to measure resting state and sound-evoked activity. Resting state studies mainly showed increased coherent activity of auditory areas [12, 13] with occasional reports of no changes between patients and controls [14]. Sound-evoked studies showed increased activity in auditory cortex [15] or asymmetric activity in correspondence to the lateralization of the tinnitus [16]. There are also reports of no changes in auditory cortex [17, 18]. One recent study showed decreased connectivity between the inferior colliculi and the auditory cortices [17]. In conclusion, tinnitus seems to be associated with increased activity and connectivity of auditory cortex.

Also studies using electroencephalography (EEG) and magnetoencephalography (MEG) showed changes in auditory evoked potentials and resting state power and connectivity. Most of these studies investigated the association of tinnitus characteristics with physiological markers of brain activity in correlational designs [19].

Because of the pathophysiological involvement of the auditory cortex in chronic tinnitus, the noninvasive brain stimulation method repetitive transcranial magnetic stimulation (rTMS) was introduced as a possible treatment option [20–22]. Despite initial positive treatment effects, it turned out over the course of the last 11 years that effect sizes of clinical trials using rTMS for treating tinnitus are moderate, interindividual variability is high, and there is still no effective cure [23]. Based on demographic and clinical data, there is no clear indicator or biomarker for rTMS treatment response or treatment effects in chronic tinnitus [24]. Neurophysiological and neuroimaging data might represent a more objective and reliable index of the treatment response than subjective ratings. EEG [25] and MEG [26] resting state and auditory evoked fMRI and EEG [27] measures were associated with rTMS treatment effects. Thus, the evaluation of rTMS efficacy by neuroimaging constitutes the next step in tinnitus research [28]. This strategy may increase the efficacy of rTMS in tinnitus by explaining the interindividual variability enabling a more individualized treatment.

Functional near-infrared spectroscopy (fNIRS) is a neuroimaging method which enables the measurement of calot-tear brain activity by measuring blood oxygenation level dependent signals based on principles of optical topography [29–31]. Despite moderate spatial and depth resolution, the big advantages of fNIRS over fMRI are the silent and upright measurement setting, mobility, handiness, little movement artifacts, and high temporal resolution [30]. Particularly in tinnitus, it is of high relevance to have as little machine noise as possible due to interaction effects with auditory cortical activity.

The aim of the present study was to probe the capability of fNIRS as a potential indicator for tinnitus-related activity in the auditory cortex. For this aim we contrasted brain

oxygenation of patients with chronic tinnitus and healthy controls and before and after treatment with rTMS over the left auditory cortex. We expect that sound-evoked activity is increased in chronic tinnitus and that hyperactivity can be decreased by cTBS. As the left temporal hemisphere was stimulated we expect effects to take primarily place on this hemisphere.

2. Material and Methods

2.1. Samples and Procedures. The study has been approved by the local ethics committee of Wuerzburg (Germany) and has been performed according to the declarations of Helsinki. Twenty-three patients with chronic tinnitus were enrolled in the study after having given written informed consent. Patients with acute or chronic inflammation of the middle ear, Menière disease, sudden idiopathic hearing loss, or fluctuating hearing were excluded. Otologic assessment included micro-otoscopy, pure-tone audiometry, tympanometry, and stapedius reflex measurement to verify normal middle ear function. Mean hearing loss in dB HL was defined as the average pure-tone hearing threshold of both ears at 0.125, 0.250, 0.500, 1, 2, 3, 4, 6, and 8 kHz. Patients with a history of seizures, a suspected diagnosis of organic brain damage, and pregnancy as well as patients with cardiac pacemakers, mobile metal implants, or implanted medication pumps were excluded.

Clinical effects and study design will be published in detail separately. In sum, we conducted a randomised sham-controlled pilot trial (verum: $n = 12$; sham: $n = 11$) using continuous theta burst stimulation of the left Heschl's gyrus targeted with anatomical neuronavigation. Stimulation was performed twice (2×600 pulses with a short break) on five days per week, starting on Monday, with an intermediate break over the weekend between the first and last five days of treatment. Stimulation intensity was set at 30% of the maximum stimulator output. fNIRS measurements were performed before (2 weeks before stimulation) and after treatment (on the day of the last treatment). Healthy controls ($n = 12$) were measured once with fNIRS. Sample characteristics are provided in Table 1.

2.2. Sound-Evoked Activity. We used one block and one event-related design for measurement of oxygenation changes induced by acoustic stimulation. "Comité Consultatif International Télégraphique et Téléphonique" (CCITT) speech noise was presented binaurally by means of insert earphones (E-A-RTONE3A, Aero Company, USA). The tip of the earphones was placed into the auditory canal, guaranteeing an exact adjustment of the sound intensity. Intensity level was set to 70 dB SPL. For the block design, participants listened to 12 blocks of CCITT noise. Each block lasted 20 s and was followed by a 20 s resting period. For the event-related design, stimuli were presented 40 times with a variable interstimulus interval of 12–14 s for 1.75 s.

2.3. Measurement of Brain Activity. Functional near-infrared spectroscopy (fNIRS) has been shown to have high validity

TABLE 1: Sample characteristics.

	Verum patient group [<i>n</i> = 12]	Sham patient group [<i>n</i> = 11]	Healthy control group [<i>n</i> = 12]	Statistics for group contrasts
Age [years]	48.2 ± 10.7	46.5 ± 11.5	43.6 ± 15.0	$F = 0.407$; $df = 2, 32$; $P = 0.669$
Sex [female/male]	5/7	4/7	3/9	$\chi^2 = 0.770$; $df = 2$; $P = 0.680$
Mean hearing loss [dB HL]	20.5 ± 11.0	21.4 ± 6.7	—	$T = 0.223$; $df = 16$; $P = 0.826$
Tinnitus duration [months]	68.9 ± 61.4	96.8 ± 120.4	—	$T = 0.702$; $df = 20$; $P = 0.491$
Tinnitus laterality [unilateral/not unilateral]	2/10	2/8	—	$\chi^2 = 0.041$; $df = 1$; $P = 0.840$
Tinnitus questionnaire [0–84]	39.4 ± 11.8	46.3 ± 15.9	—	$T = 1.181$; $df = 21$; $P = 0.251$
Tinnitus handicap inventory [0–100]	42.3 ± 17.5	50.2 ± 18.7	—	$T = 1.039$; $df = 21$; $P = 0.311$
Beck depression inventory II [0–63]	11.6 ± 8.8	7.0 ± 7.2	—	$T = 1.356$; $df = 21$; $P = 0.190$

[32] and reliability [33–35] and is suited to measure sound evoked activity in the auditory cortex [36]. For the fNIRS measurement, we used a continuous multichannel wave system (ETG4000 Optical Topography System; Hitachi Medical Co., Japan) working with two different wavelengths (695 ± 20 nm and 830 ± 20 nm) and a time resolution of 10 Hz to measure relative changes of absorbed near-infrared light. These changes are transformed into concentration changes of oxygenated (O_2Hb) and deoxygenated haemoglobin (HHb) as indicators for brain activity by means of a modified Beer-Lambert law [37]. The unit is $mmol \times mm$; that is, changes of chromophore concentration depend on the path length of the near-infrared light. We used two identical rectangular probe sets (plastic panels) of optodes (light emitters and detectors) for each side of the head. One probe set consisted of 8 light emitters and 7 detectors with an interoptode distance of 3 cm. A measuring point of activation (channel) was defined as the region between one emitter and one detector. Thus, one probe set consisted of 22 channels and covered an area of 6×12 cm on the scalp. The panels were fastened to the head by elastic straps. The probe sets were placed on the head with regard to the relevant standard positions of the international 10–20 system for EEG electrode placement [38, 39]. The channel over the middle inferior optode was placed over T3/T4 with vertical orientation in direction to C3/C4. The arrangement of the probe set is shown in the top of Figure 1.

2.4. Data Analysis and Statistics. Before statistical analysis of fNIRS data, the high frequency portion of the signal was removed by calculating a moving average with a time window of 5 seconds. During normal physiological activity neurovascular decoupling is accompanied in local increases in O_2Hb and simultaneous decreases in HHb resulting in negative correlations of both chromophores. Correlations that tend to be positive or equal to zero may indicate noise caused by motion or extracerebral or systemic hemodynamic activity unrelated to the experimental condition [37].

A correlation based signal improvement algorithm was used to filter out spikes and to improve signal quality based on the assumed negative correlation between O_2Hb and HHb [40]. The applied correction results in one single parameter of brain oxygenation which is a mathematical combination of O_2Hb and HHb.

For the block design measurement, slow drifts in the measurement were excluded by the use of a linear fitting resulting from calculation of the mean of the 10 s right before the auditory stimulation and the mean of the interval between the 10th and 20th second after the stimulation. A mean trajectory for each condition was calculated by averaging the 12 repetitions. For statistical analyses, we used the mean amplitude of oxygenation during the stimulation period. For the event-related design, slow drifts in the signal were excluded by the use of a high pass filter with six discrete cosine basis functions. We used the general linear model approach according to fMRI and fNIRS literature [41–43]. Association of channels and brain areas were deducted from the work of Okamoto and colleagues [39], which interrelates EEG positions and fNIRS probe sets. Analyses were done by home-made scripts on MatLab (The MathWorks Inc., USA).

For statistical analyses of imaging data we used two-sided Student *t*-tests and analyses of variance (ANOVAs) for each channel of each probe set. Firstly, for reasons of plausibility we contrasted the activity (amplitude for the block and beta values for the event-related design) of each channel against zero to test for region-specific activity of auditory areas. These basal brain activations were indicated by two-sided Student *t*-tests against zero and by according T-maps interpolated for all channels over the whole probe set (Figure 1). Secondly, we analysed overall group differences for the baseline measurements and contrasted the group of patients and the group of controls with unpaired *t*-tests. Thirdly, to evaluate intervention specific pre-post effects we conducted ANOVAs with the independent factor group (verum versus sham) and the dependent factor time

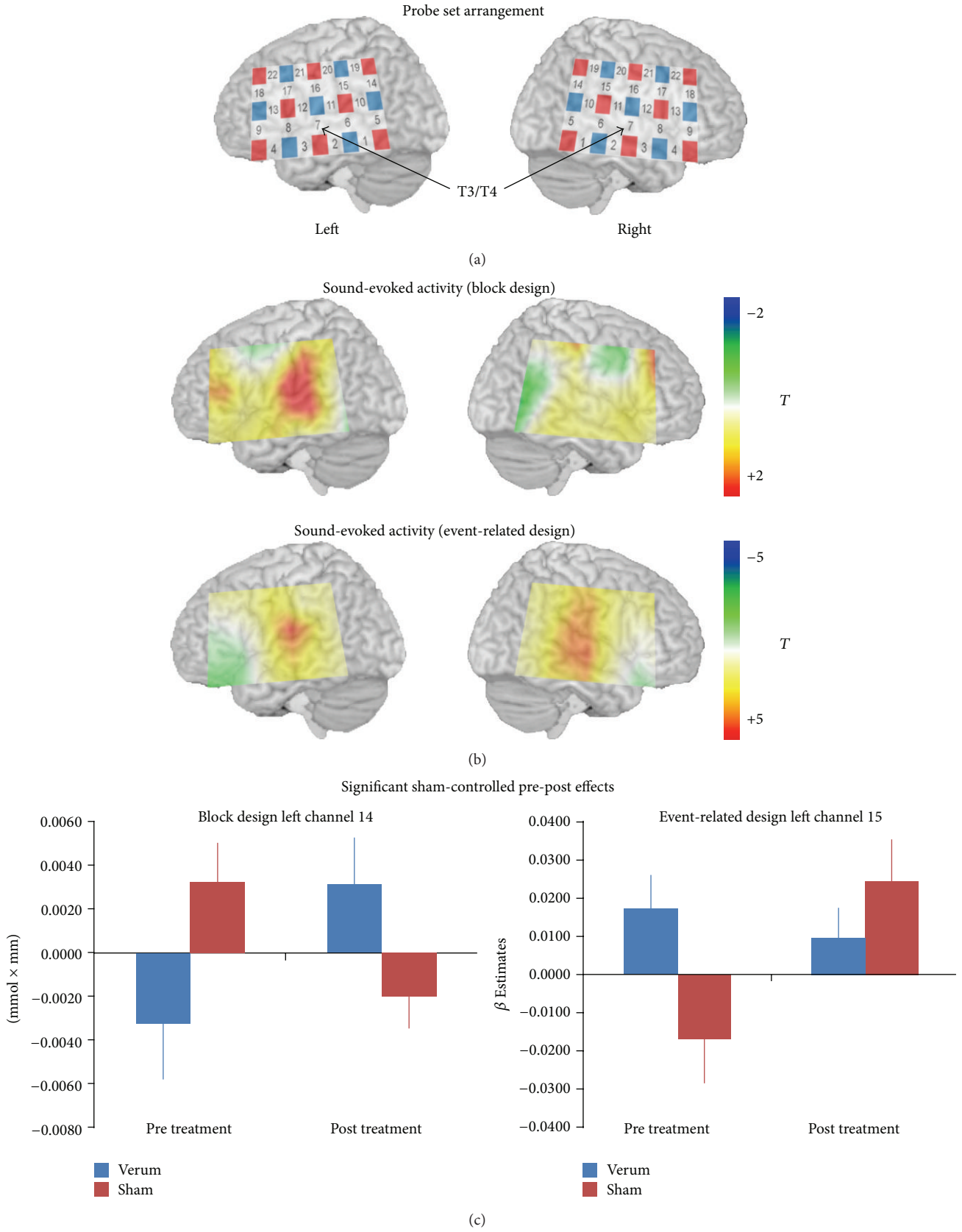


FIGURE 1: Localisation of light emitters (red squares) and detectors (blue squares) and measurement channels (numbers) for near-infrared spectroscopy (a). Sound-evoked activity for all subjects at baseline ($n = 35$; tinnitus patients and healthy controls) as indicated by brain oxygenation (b). Channels with significant group by time interaction effects (c).

(pre versus post). Post hoc analyses were done for significant interactions by using *t*-tests. To take baseline differences into account, we repeated the ANOVAs with significant interaction effects by including the baseline values as covariate (ANCOVAs). Due to 22 calculations per probe set we used a Bonferroni correction by using the Dubey/Armitage-Parmar alpha boundary taking into account spatial correlation among the fNIRS channels [44–47]. Based on the baseline measurements the mean correlation coefficient was 0.539 resulting in an adjusted alpha level of 0.0123. Thus, we just report *P* values below 1.23%. For post hoc tests and the ANCOVAs we abstained from using Bonferroni correction. All statistical analyses were performed with SPSS 20.0.0.1 (SPSS Inc., USA).

3. Results

For all measured subjects (verum and sham patients and control group) both designs showed activity over auditory areas. Topographies of brain oxygenation in the form of T-maps are shown in Figure 1. Statistical values were higher for the event-related design in contrast to the block-design experiment as indicated by the color bar ranges of Figure 1. In line, only for the event-related design we found significant activity (left hemisphere: channel 11: $P < 0.001$; channel 16: $P = 0.010$; right hemisphere: channel 2: $P = 0.009$; channel 6: $P = 0.005$; channel 7: $P < 0.001$; channel 11: $P < 0.001$; channel 16: $P < 0.001$; channel 20: $P = 0.007$). Nonsignificant activity can be seen only over the left auditory areas for the block design.

In the next step we analysed overall group differences for the baseline measurements. *t*-tests between patients and controls showed group differences in the right auditory cortex as indicated by significant differences in right channels 2 ($P < 0.001$) and 7 ($P = 0.006$) for the block and in the right channel 12 for the event-related design ($P = 0.004$). The significant findings of the block design were related to auditory areas; patients had higher oxygenation changes in contrast to controls. The significant channel of the event-related design was related to frontal areas showing decreased activity in the patient group. To sum up the baseline findings, oxygenation in the right hemisphere differed between patients and controls; that is, patients showed higher auditory oxygenation during the block design and more focused activation during the event-related design as mirrored by decreased activation in frontal areas. In the left hemisphere there were no clear differences between patient groups and the control group.

Pre-post changes showed no significant group \times time interaction effects on the right hemisphere—neither for the block nor for the event-related design. For the block design, channel 14 showed a significant interaction effect ($P = 0.005$) (Figure 1). During baseline, the sham group had higher oxygenation in contrast to the verum group ($P = 0.057$). At the last treatment visit, this relation was reversed ($P = 0.061$). The verum group displayed higher oxygenation than the sham group. Oxygenation decreased in the sham group over the course of the trial ($P = 0.059$) and increased for the

verum group ($P = 0.039$). The effect was stable by controlling for baseline differences using an ANCOVA with baseline values as covariate ($P = 0.041$).

In the event-related design, left channel 15 showed a significant interaction effect in the opposite direction as compared to the effect in the block design ($P = 0.005$). The verum group had higher oxygenation during the baseline in contrast to the sham group ($P = 0.025$) with no oxygenation differences during the last treatment visit ($P = 0.294$). Oxygenation increased in the sham group over the course of the trial ($P = 0.012$) with no changes for the verum group ($P = 0.476$). The effect disappeared by controlling for baseline differences using an ANCOVA with baseline values as covariate ($P = 0.135$). To sum up the pre-post effects, the activity of the right hemisphere was not sensitive to specific treatment effects. For the left hemisphere, increased oxygenation during baseline decreased over the course of the treatment and lowered oxygenation during the baseline increased over the course of the treatment with reversed patterns between the verum and sham group and between the block and event-related design. Findings of the ANCOVA indicate that these interaction effects are highly dependent from the baseline values.

4. Discussion

In line with a former study [36], we were able to measure sound-evoked activity in temporal areas with fNIRS. A finding of the present study was a reversed pattern in brain oxygenation between the block and event-related design. For the right hemisphere, activity differences between patients and controls were mirrored by increased activity for the block design and rather focused activity in the event-related design in the patients. For the left hemisphere, oxygenation was changing during treatment with reversed patterns for the verum and the sham group. For the block design, the sham group showed reductions from an increased activity level over the course of treatment; the verum group showed the opposite pattern. This interaction was reversed in the event-related design. Differences between the block and event related design have already been proposed as a potential explanation for the limited reliability in fMRI research [48]. Our findings suggest that short sounds within the range of 1-2 s in contrast to long-lasting sound stimulation within the range of several seconds might lead to differences in neural activation. Differences in brain activation dependent on stimulus duration seem to be reasonable. In contrast to a short salient sound, the same sound becomes irrelevant if presented for a longer time and is presumably inhibited via top-down mechanisms. This might be in accordance with models of prepulse inhibition and sensory gating for which a preceding stimulus decreases the novelty of the incoming information [49, 50]. The sensitivity of fNIRS for top-down mechanisms in the auditory domain has been documented in an earlier fNIRS study which detected an influence of the emotional valence on sound induced temporal oxygenation [36]. Whether the observed differences between the event-related design and the block design are a general

phenomenon or specific for tinnitus has to be elucidated by further studies.

A further notion is that our data did not clearly indicate a favourable sound-evoked design to measure activity specific for tinnitus and/or specific for rTMS interventions over the auditory cortex. The event-related design showed higher baseline T -values and clear bilateral activity in contrast to the block design with smaller T -values and only left-sided activity for the whole group (Figure 1). On the other hand rTMS related changes in the block design seem to be more stable for baseline differences in pre-post designs.

One advantage of fNIRS is the possibility to easily combine it with noninvasive brain stimulation to measure TMS induced activity changes [51, 52]. In the present trial rTMS and fNIRS were performed successively because of the interest in the neurobiological change induced by a therapeutic intervention. The present findings showed trait related increases of brain oxygenation in the right auditory areas as indicated by increased activity in the block design and more focused activity in the event-related design. Increased activity in patients is in line with several studies using sound-evoked fMRI [16, 53] or tinnitus-related alterations in oxygenation or metabolic brain activity [10]. This increased activity is considered to represent the tinnitus percept or at least one aspect of the tinnitus percept [54, 55]. The quiet measurement setting of fNIRS affirms the fMRI findings. The latter method might have been confounded by scanner background noise due to the radio waves and cooling pumps, which limits the interpretation of fMRI findings in tinnitus [56].

Our results further suggest that left auditory cortex oxygenation may reflect state-like effects. Based on the baseline level (block design: sham > verum; event-related design: verum > sham) the intervention led to a decrease in the group with increased baseline activity and vice versa (block design: verum > sham; event-related design: sham > verum). The importance of these pre-post effects is highlighted by the ANCOVAs. The interpretation of these findings is challenging because of the difference in baseline activity between these two groups. Thus it is difficult to disentangle, to which extent the inverse changes during treatment depend on the stimulation procedure (verum versus sham) or on the difference in baseline measurements. Nevertheless our findings are in accordance with a recent fMRI study in tinnitus showing the same neural pattern [27]. Our findings of treatment related changes in temporal blood flow are also comparable to findings of temporoparietal rTMS for the treatment of auditory hallucinations in schizophrenia. In twelve medication-resistant patients, PET revealed a reduction in symptoms and left temporoparietal metabolism after low frequency of the corresponding area [57]. In another study cerebral blood flow (arterial spin labeling) in the left superior temporal gyrus predicted treatment response [58]. Open questions for future work are the meaning of negative baseline values during baseline and if the pretreatment differences are due to differences in samples. These might be resolved by more rigorous characterization of the samples, by increasing the sample size, by repeated baseline measurements, and by correlation of baseline values with tinnitus characteristics.

Similar like in sound evoked fMRI or PET studies, we have to acknowledge that it still has to be elucidated, how the observed alterations of sound evoked activity in tinnitus relate to its neuronal correlates. Alterations in sound evoked activity can only provide an indirect hint for the neuronal changes underlying tinnitus.

Altogether, even if we cannot provide final explanations for all aspects of our fNIRS results, the data confirm the usefulness of fNIRS for neuroimaging of auditory function. Future studies including the same design in control subjects should shed even more light to trait- and state-like effects of rTMS on auditory cortex activity in tinnitus. Open questions in tinnitus research might be evaluable with this method such as association of tinnitus characteristics (e.g., laterality) with temporal activity [11, 16] or combination with EEG to verify source localisation of power changes [59, 60]. Also for the investigation of other areas such as the orbitofrontal cortex fNIRS has a higher suitability because of susceptibility artifacts in fMRI for these areas [61]. Generally, other fields of phantom perception such as acoustic hallucinations might be investigable with fNIRS [62].

5. Conclusions

The present proof-of-principle study showed that fNIRS is able to measure brain oxygenation changes of the auditory cortices in relation to tinnitus. Furthermore, the combination of noninvasive brain stimulation and neuroimaging offers the detailed investigation of trait- and state-related biomarkers for tinnitus. As fNIRS measures calotte-near brain oxygenation changes which means rather secondary than primary auditory cortex and as rTMS findings also indicate secondary and temporoparietal areas as potential treatment targets [63, 64], the question arises if fNIRS might serve as indicator for functional neuronavigation which is the stimulation of the nonprimary auditory cortical areas with sound-evoked related activity.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

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References

- [1] H. J. Hoffman and G. W. Reed, "Epidemiology of tinnitus," in *Tinnitus: Theory and Management*, J. B. Snow Jr., Ed., pp. 16–42, BC Decker, Hamilton, Canada, 2004.
- [2] J. J. Eggermont and L. E. Roberts, "The neuroscience of tinnitus," *Trends in Neurosciences*, vol. 27, no. 11, pp. 676–682, 2004.
- [3] J. J. Eggermont and L. E. Roberts, "The neuroscience of tinnitus: understanding abnormal and normal auditory perception," *Frontiers in Systems Neuroscience*, vol. 6, article 53, 2012.

- [4] N. D. Engineer, J. R. Riley, J. D. Seale et al., “Reversing pathological neural activity using targeted plasticity,” *Nature*, vol. 470, no. 7332, pp. 101–106, 2011.
- [5] W. Mühlnickel, T. Elbert, E. Taub, and H. Flor, “Reorganization of auditory cortex in tinnitus,” *Proceedings of the National Academy of Sciences of the United States of America*, vol. 95, no. 17, pp. 10340–10343, 1998.
- [6] D. R. M. Langers, “Assessment of tonotopically organised subdivisions in human auditory cortex using volumetric and surface-based cortical alignments,” *Human Brain Mapping*, vol. 35, no. 4, pp. 1544–1561, 2014.
- [7] C. P. Lanting, E. de Kleine, and P. van Dijk, “Neural activity underlying tinnitus generation: results from PET and fMRI,” *Hearing Research*, vol. 255, no. 1–2, pp. 1–13, 2009.
- [8] P. Adjamian, M. Sereda, and D. A. Hall, “The mechanisms of tinnitus: perspectives from human functional neuroimaging,” *Hearing Research*, vol. 253, no. 1–2, pp. 15–31, 2009.
- [9] M. Schecklmann, M. Landgrebe, T. B. Poepl et al., “Neural correlates of tinnitus duration and Distress: a positron emission tomography study,” *Human Brain Mapping*, vol. 34, no. 1, pp. 233–240, 2013.
- [10] J.-J. Song, D. de Ridder, P. van de Heyning, and S. Vanneste, “Mapping tinnitus-related brain activation: an activation-likelihood estimation metaanalysis of PET studies,” *Journal of Nuclear Medicine*, vol. 53, no. 10, pp. 1550–1557, 2012.
- [11] L. I. Geven, E. de Kleine, A. T. M. Willemsen, and P. van Dijk, “Asymmetry in primary auditory cortex activity in tinnitus patients and controls,” *Neuroscience*, vol. 256, pp. 117–125, 2014.
- [12] A. Maudoux, P. Lefebvre, J.-E. Cabay et al., “Auditory resting-state network connectivity in tinnitus: a functional MRI study,” *PLoS ONE*, vol. 7, no. 5, Article ID e36222, 2012.
- [13] S. A. Schmidt, K. Akrofi, J. R. Carpenter-Thompson, and F. T. Husain, “Default mode, dorsal attention and auditory resting state networks exhibit differential functional connectivity in tinnitus and hearing loss,” *PLoS ONE*, vol. 8, no. 10, Article ID e76488, 2013.
- [14] J. Davies, P. E. Gander, M. Andrews, and D. A. Hall, “Auditory network connectivity in tinnitus patients: a resting-state fMRI study,” *International Journal of Audiology*, vol. 53, no. 3, pp. 192–198, 2014.
- [15] A. M. Leaver, L. Renier, M. A. Chevillet, S. Morgan, H. J. Kim, and J. P. Rauschecker, “Dysregulation of limbic and auditory networks in tinnitus,” *Neuron*, vol. 69, no. 1, pp. 33–43, 2011.
- [16] M. Smits, S. Kovacs, D. de Ridder, R. R. Peeters, P. van Hecke, and S. Sunaert, “Lateralization of functional magnetic resonance imaging (fMRI) activation in the auditory pathway of patients with lateralized tinnitus,” *Neuroradiology*, vol. 49, no. 8, pp. 669–679, 2007.
- [17] K. Boyen, E. de Kleine, P. van Dijk, and D. R. Langers, “Tinnitus-related dissociation between cortical and subcortical neural activity in humans with mild to moderate sensorineural hearing loss,” *Hearing Research*, vol. 312, pp. 48–59, 2014.
- [18] C. P. Lanting, E. de Kleine, H. Bartels, and P. van Dijk, “Functional imaging of unilateral tinnitus using fMRI,” *Acta Oto-Laryngologica*, vol. 128, no. 4, pp. 415–421, 2008.
- [19] S. Vanneste and D. de Ridder, “The auditory and non-auditory brain areas involved in tinnitus. An emergent property of multiple parallel overlapping subnetworks,” *Frontiers in Systems Neuroscience*, vol. 6, article 31, 2012.
- [20] T. Kleinjung, P. Eichhammer, B. Langguth et al., “Long-term effects of repetitive transcranial magnetic stimulation (rTMS) in patients with chronic tinnitus,” *Otolaryngology—Head and Neck Surgery*, vol. 132, no. 4, pp. 566–569, 2005.
- [21] P. Eichhammer, B. Langguth, J. Marienhagen, T. Kleinjung, and G. Hajak, “Neuronavigated repetitive transcranial magnetic stimulation in patients with tinnitus: a short case series,” *Biological Psychiatry*, vol. 54, no. 8, pp. 862–965, 2003.
- [22] C. Plewnia, M. Reimold, A. Najib, G. Reischl, S. K. Plontke, and C. Gerloff, “Moderate therapeutic efficacy of positron emission tomography-navigated repetitive transcranial magnetic stimulation for chronic tinnitus: a randomised, controlled pilot study,” *Journal of Neurology, Neurosurgery and Psychiatry*, vol. 78, no. 2, pp. 152–156, 2007.
- [23] B. Langguth and D. de Ridder, “Tinnitus: therapeutic use of superficial brain stimulation,” *Handbook of Clinical Neurology*, vol. 116, pp. 441–467, 2013.
- [24] A. Lehner, M. Schecklmann, M. Landgrebe et al., “Predictors for rTMS response in chronic tinnitus,” *Frontiers in Systems Neuroscience*, vol. 6, article 11, 2012.
- [25] D. De Ridder, J.-J. Song, and S. Vanneste, “Frontal cortex TMS for tinnitus,” *Brain Stimulation*, vol. 6, no. 3, pp. 355–362, 2013.
- [26] T. Hartmann, I. Lorenz, N. Müller, B. Langguth, and N. Weisz, “The effects of neurofeedback on oscillatory processes related to tinnitus,” *Brain Topography*, vol. 27, no. 1, pp. 149–157, 2014.
- [27] J.-P. Lefaucheur, P. Brugières, F. Guimont et al., “Navigated rTMS for the treatment of tinnitus: a pilot study with assessment by fMRI and AEPs,” *Neurophysiologie Clinique*, vol. 42, no. 3, pp. 95–109, 2012.
- [28] B. Langguth, M. Schecklmann, A. Lehner et al., “Neuroimaging and neuromodulation: complementary approaches for identifying the neuronal correlates of tinnitus,” *Frontiers in Systems Neuroscience*, vol. 6, article 15, 2012.
- [29] F. Scholkmann, S. Kleiser, A. J. Metz et al., “A review on continuous wave functional near-infrared spectroscopy and imaging instrumentation and methodology,” *NeuroImage*, vol. 85, part 1, pp. 6–27, 2014.
- [30] A.-C. Ehlis, S. Schneider, T. Dresler, and A. J. Fallgatter, “Application of functional near-infrared spectroscopy in psychiatry,” *NeuroImage*, vol. 85, part 1, pp. 478–488, 2014.
- [31] M. Ferrari and V. Quaresima, “A brief review on the history of human functional near-infrared spectroscopy (fNIRS) development and fields of application,” *NeuroImage*, vol. 63, no. 2, pp. 921–935, 2012.
- [32] T. J. Huppert, R. D. Hoge, S. G. Diamond, M. A. Franceschini, and D. A. Boas, “A temporal comparison of BOLD, ASL, and NIRS hemodynamic responses to motor stimuli in adult humans,” *NeuroImage*, vol. 29, no. 2, pp. 368–382, 2006.
- [33] M. M. Plichta, M. J. Herrmann, C. G. Baehne et al., “Event-related functional near-infrared spectroscopy (fNIRS): are the measurements reliable?” *NeuroImage*, vol. 31, no. 1, pp. 116–124, 2006.
- [34] M. M. Plichta, M. J. Herrmann, C. G. Baehne et al., “Event-related functional near-infrared spectroscopy (fNIRS) based on craniocerebral correlations: reproducibility of activation?” *Human Brain Mapping*, vol. 28, no. 8, pp. 733–741, 2007.
- [35] M. Schecklmann, A.-C. Ehlis, M. M. Plichta, and A. J. Fallgatter, “Functional near-infrared spectroscopy: a long-term reliable tool for measuring brain activity during verbal fluency,” *NeuroImage*, vol. 43, no. 1, pp. 147–155, 2008.
- [36] M. M. Plichta, A. B. M. Gerdes, G. W. Alpers et al., “Auditory cortex activation is modulated by emotion: a functional near-infrared spectroscopy (fNIRS) study,” *NeuroImage*, vol. 55, no. 3, pp. 1200–1207, 2011.

- [37] H. Obrig and A. Villringer, "Beyond the visible—imaging the human brain with light," *Journal of Cerebral Blood Flow and Metabolism*, vol. 23, no. 1, pp. 1–18, 2003.
- [38] H. H. Jasper, "The ten-twenty electrode system of the International Federation," *Electroencephalography and Clinical Neurophysiology*, vol. 10, pp. 371–375, 1958.
- [39] M. Okamoto, H. Dan, K. Sakamoto et al., "Three-dimensional probabilistic anatomical cranio-cerebral correlation via the international 10–20 system oriented for transcranial functional brain mapping," *NeuroImage*, vol. 21, no. 1, pp. 99–111, 2004.
- [40] X. Cui, S. Bray, and A. L. Reiss, "Functional near infrared spectroscopy (NIRS) signal improvement based on negative correlation between oxygenated and deoxygenated hemoglobin dynamics," *NeuroImage*, vol. 49, no. 4, pp. 3039–3046, 2010.
- [41] K. J. Friston, A. P. Holmes, J.-B. Poline et al., "Analysis of fMRI time-series revisited," *NeuroImage*, vol. 2, no. 1, pp. 45–53, 1995.
- [42] M. L. Schroeter, M. M. Bücheler, K. Müller et al., "Towards a standard analysis for functional near-infrared imaging," *NeuroImage*, vol. 21, no. 1, pp. 283–290, 2004.
- [43] M. M. Plichta, S. Heinzel, A.-C. Ehlis, P. Pauli, and A. J. Fallgatter, "Model-based analysis of rapid event-related functional near-infrared spectroscopy (NIRS) data: a parametric validation study," *NeuroImage*, vol. 35, no. 2, pp. 625–634, 2007.
- [44] S. V. Tupak, T. Dresler, A. Guhn et al., "Implicit emotion regulation in the presence of threat: neural and autonomic correlates," *NeuroImage*, vol. 85, pp. 372–379, 2014.
- [45] M. M. Plichta, M. J. Herrmann, C. G. Baehne et al., "Event-related functional near-infrared spectroscopy (fNIRS) based on cranio-cerebral correlations: reproducibility of activation?" *Human Brain Mapping*, vol. 28, no. 8, pp. 733–741, 2007.
- [46] A. Sassaroli, Y. Tong, C. Benes, and S. Fantini, "Data analysis and statistical tests for near-infrared functional studies of the brain," in *Multimodal Biomedical Imaging III*, vol. 6850 of *Proceedings of SPIE*, January 2008.
- [47] A. J. Sankoh, M. F. Huque, and S. D. Dubey, "Some comments on frequently used multiple endpoint adjustment methods in clinical trials," *Statistics in Medicine*, vol. 16, no. 22, pp. 2529–2542, 1997.
- [48] C. M. Bennett and M. B. Miller, "fMRI reliability: influences of task and experimental design," *Cognitive, Affective, & Behavioral Neuroscience*, vol. 13, no. 4, pp. 690–702, 2013.
- [49] Y. du, X. Wu, and L. li, "Differentially organized top-down modulation of prepulse inhibition of startle," *The Journal of Neuroscience*, vol. 31, no. 38, pp. 13644–13653, 2011.
- [50] A.-C. Ehlis, T. M. Ringel, M. M. Plichta, M. M. Richter, M. J. Herrmann, and A. J. Fallgatter, "Cortical correlates of auditory sensory gating: A simultaneous near-infrared spectroscopy event-related potential study," *Neuroscience*, vol. 159, no. 3, pp. 1032–1043, 2009.
- [51] Y. Noguchi, E. Watanabe, and K. L. Sakai, "An event-related optical topography study of cortical activation induced by single-pulse transcranial magnetic stimulation," *NeuroImage*, vol. 19, no. 1, pp. 156–162, 2003.
- [52] R. H. Thomson, J. J. Maller, Z. J. Daskalakis, and P. B. Fitzgerald, "Blood oxygenation changes resulting from trains of low frequency transcranial magnetic stimulation," *Cortex*, vol. 48, no. 4, pp. 487–491, 2012.
- [53] J. W. Gu, C. F. Halpin, E.-C. Nam, R. A. Levine, and J. R. Melcher, "Tinnitus, diminished sound-level tolerance, and elevated auditory activity in humans with clinically normal hearing sensitivity," *Journal of Neurophysiology*, vol. 104, no. 6, pp. 3361–3370, 2010.
- [54] D. de Ridder, S. Vanneste, N. Weisz et al., "An integrative model of auditory phantom perception: tinnitus as a unified percept of interacting separable subnetworks," *Neuroscience and Biobehavioral Reviews*, vol. 44, pp. 16–32, 2013.
- [55] D. De Ridder, A. B. Elgoyhen, R. Romo, and B. Langguth, "Phantom percepts: tinnitus and pain as persisting aversive memory networks," *Proceedings of the National Academy of Sciences of the United States of America*, vol. 108, no. 20, pp. 8075–8080, 2011.
- [56] J. R. Melcher, R. A. Levine, C. Bergevin, and B. Norris, "The auditory midbrain of people with tinnitus: abnormal sound-evoked activity revisited," *Hearing Research*, vol. 257, no. 1-2, pp. 63–74, 2009.
- [57] J. Horacek, M. Brunovsky, T. Novak et al., "Effect of low-frequency rTMS on electromagnetic tomography (LORETA) and regional brain metabolism (PET) in schizophrenia patients with auditory hallucinations," *Neuropsychobiology*, vol. 55, no. 3-4, pp. 132–142, 2007.
- [58] P. Homan, J. Kindler, M. Hauf, D. Hubl, and T. Dierks, "Cerebral blood flow identifies responders to transcranial magnetic stimulation in auditory verbal hallucinations," *Translational Psychiatry*, vol. 2, article e189, 2012.
- [59] N. Weisz, S. Moratti, M. Meinzer, K. Dohrmann, and T. Elbert, "Tinnitus perception and distress is related to abnormal spontaneous brain activity as measured by magnetoencephalography," *PLoS Medicine*, vol. 2, no. 6, article e153, 2005.
- [60] W. Schlee, T. Hartmann, B. Langguth, and N. Weisz, "Abnormal resting-state cortical coupling in chronic tinnitus," *BMC Neuroscience*, vol. 10, article 11, 2009.
- [61] F. B. Haeussinger, S. Heinzel, T. Hahn, M. Schecklmann, A.-C. Ehlis, and A. J. Fallgatter, "Simulation of near-infrared light absorption considering individual head and prefrontal cortex anatomy: implications for optical neuroimaging," *PLoS ONE*, vol. 6, no. 10, Article ID e26377, 2011.
- [62] P. Allen, F. Larøi, P. K. McGuire, and A. Aleman, "The hallucinating brain: a review of structural and functional neuroimaging studies of hallucinations," *Neuroscience and Biobehavioral Reviews*, vol. 32, no. 1, pp. 175–191, 2008.
- [63] E. M. Khedr, J. C. Rothwell, M. A. Ahmed, and A. El-Atar, "Effect of daily repetitive transcranial magnetic stimulation for treatment of tinnitus: Comparison of different stimulus frequencies," *Journal of Neurology, Neurosurgery and Psychiatry*, vol. 79, no. 2, pp. 212–215, 2008.
- [64] C. Plewnia, R. Vonthein, B. Wasserka et al., "Treatment of chronic tinnitus with theta burst stimulation: a randomized controlled trial," *Neurology*, vol. 78, no. 21, pp. 1628–1634, 2012.