Relationship between regional white matter hyperintensities and alpha oscillations in older adults

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ARTICLE INFO

Article history:
Received 2 October 2020
Revised 22 September 2021
Accepted 12 October 2021
Available online 20 October 2021

Keywords:
EEG
MRI
White matter hyperintensity
Aging
Alpha power
Resting-state

ABSTRACT

Aging is associated with increased white matter hyperintensities (WMHs) and with alterations of alpha oscillations (7–13 Hz). However, a crucial question remains, whether changes in alpha oscillations relate to aging per se or whether this relationship is mediated by age-related neuropathology like WMHs. Using a large cohort of cognitively healthy older adults (N = 907, 60–80 years), we assessed relative alpha power, alpha peak frequency, and long-range temporal correlations from resting-state EEG. We further associated these parameters with voxel-wise WMHs from 3T MRI. We found that a higher prevalence of WMHs in the superior and posterior corona radiata as well as in the thalamic radiation was related to elevated alpha power, with the strongest association in the bilateral occipital cortex. In contrast, we observed no significant relation of the WMHs probability with alpha peak frequency and long-range temporal correlations. Finally, higher age was associated with elevated alpha power via total WMH volume. We suggest that an elevated alpha power is a consequence of WMHs affecting a spatial organization of alpha sources.

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One of the most prominent EEG rhythms is alpha oscillations (7–13 Hz), which have been shown to originate from thalamocortical and cortico-cortical interactions (Bazanova and Vernon, 2014; Lopes Da Silva et al., 1997). Importantly, measures of alpha oscillations have been related to many aspects of cognitive function (Klimesch, 1999) and also to endophenotypes of brain aging (Ishii et al., 2018; Knyazeva et al., 2018) either using alpha peak frequency or power. While individual alpha peak frequency has been consistently shown to decrease with age (Ishii et al., 2018; Knyazeva et al., 2018; Mierau et al., 2017), the findings on alpha power remain rather inconsistent. Previous EEG studies showed decreases of alpha power across the lifespan when using relatively large sample sizes (Babiloni et al., 2006a; Lodder and van Putten, 2011; Vysata et al., 2012), yet these age-related reductions in alpha power were either not strongly present within the older age groups (>60 years of age; Lodder and van Putten, 2011) or not replicated (Sahoo et al., 2020; Scally et al., 2018).

Apart from these 2 measures of alpha oscillations, temporal dynamics of the signals can be quantified with autocorrelation showing to what extent the past of the signal relates to its future. A very slow attenuation of the autocorrelation, which can be described with a power law, is also referred to as long-range temporal correlations (LRTC). The presence of LRTC indicates scale-free properties of the signal fluctuation pattern that look similar at different time scales. LRTC in the amplitude envelope of the neuronal oscillations were shown to extend to tens or even hundreds of seconds (Linkenkaer-Hansen et al., 2001; Nikulin and Brismar, 2005). Importantly, the presence of LRTC is consistent with the idea that neuronal networks may operate at a critical state, characterized by a balance between inhibition and excitation (Linkenkaer-Hansen et al., 2001; Nikulin and Brismar, 2005; Palva et al., 2013; Shew and Plenz, 2013). LRTC exponent that represents the decay of the autocorrelation has been linked to functional connectivity measures (Zhigalov et al., 2017), brain maturation (Smit et al., 2011), and different aspects of cognition (Mahjooory et al., 2019; Samek et al., 2016; Smit et al., 2011). Given previous literature and a recent work by Quandt et al. (2020) showing a link between WMHs and altered functional connectivity, it is plausible that age-related structural changes in the brain could manifest in alterations of LRTC.

As both static (i.e., power, individual alpha peak frequency) and dynamic (i.e., LRTC) measures of alpha oscillations might be affected by microstructural deteriorations, due to the disconnection among neural cells and damage to cortico-cortical and cortico-subcortical pathways (Madden et al., 2017), WMHs-associated alterations of EEG rhythms are plausible. However, there are only a few EEG studies that have directly investigated the relationship between alpha oscillations and WMHs or integrity (Babiloni et al., 2011, 2008a; Valdés-Hernández et al., 2010; van Straaten et al., 2012). Previously, local and global disturbances of brain anatomy like white matter microstructure (Hinault et al., 2020; Hendriks et al., 2015; Minami et al., 2020; Valdés-Hernández et al., 2010) have been found to be related to alpha rhythm affecting its peak frequency and power. For instance, a previous study with 222 subjects using the Cuban Human Brain Mapping Project (Valdés-Hernández et al., 2010) provides evidence that alpha peak frequency can be associated with both decrease and increase (depending on the region) in the microstructure of thalamocortical or corticothalamic fibers assessed by Fractional Anisotropy using diffusion tensor imaging. Interestingly, so far only a few studies have investigated the relationship between alpha power and WMHs (Babiloni et al., 2009, 2008b,2008a). For instance, it has been observed that higher alpha power was associated with higher scores of the prevalence of WMHs in individuals with mild cognitive impairment (Babiloni et al., 2008a). Similarly, a recent study (Quandt et al., 2020) reported that higher WHM lesion load was related to reduced EEG alpha connectivity measures in healthy older adults (N = 35). However, to our knowledge, no link between voxel-wise whole-brain WMHs and different parameters of alpha oscillations has been investigated using a large sample of healthy older adults. Moreover, a crucial question still remains unresolved, for example, whether changes in alpha oscillations relate to normal aging per se or rather they represent the impact of age-related neuropathology, for instance, WMHs. In this study, using a large population-based sample, we investigated neurophysiological links between age, WMHs and alpha oscillations. More precisely, we investigated the association between age and parameters of alpha oscillations, and whether this relationship was mediated by WMHs. We further explored the association of WMHs with parameters of alpha oscillations in a topographically specific manner taking into account the location of the lesioned white matter tracts.

2. Methods

2.1. Participants

Participants were drawn from the population-based Leipzig Research Center for Civilization Diseases LIFE-Adult study (Loefler et al., 2015). All participants provided written informed consent, and the study was approved by the ethics committee of the medical faculty at the University of Leipzig, Germany. The study was performed in agreement with the Declaration of Helsinki. A subset of participants underwent a 3-Tesla MRI head scan and resting-state (rs)EEG recordings on 2 separate assessment days. We selected participants above 60 years of age and with additional brain pathology or history of stroke, multiple sclerosis, epilepsy, Parkinson’s disease, intracranial hemorrhage, or brain tumors. We further excluded individuals whose rsEEG recordings were not temporally close to the MRI acquisition time and participants for whom alpha peak could not be identified. The details about the time differences between EEG and MRI measurement days can be found in Supplementary Figure 1 (M = 23.4 in absolute days). This resulted in a final sample of 907 participants (M = 69.49 ± 4.63 years of age, 380 female) for the rsEEG sensor space analysis. After excluding individuals with failed T1-weighted segmentation and head-modeling, the final sample for the rsEEG source analysis was 855 (M = 68.89 ± 4.66 years of age, 360 female). For a detailed overview of the selection process, see Fig. 1.

2.2. MRI acquisition and processing

All MRI scans were performed at 3 Tesla on a MAGNETOM Vario scanner (Siemens, Erlangen, Germany). The body coil was used for radiofrequency (RF) transmission and a 32-channel head coil was used for signal reception. T1-weighted MPRAGE and FLAIR images were acquired as part of a standardized protocol: MPRAGE (flip angle [FA] = 9°, relaxation time [TR] = 2300 ms, inversion time [TI] = 900 ms, echo time [TE] = 2.98 ms, 1-mm isotropic resolution, acquisition time [AT] = 5.10 minutes); FLAIR (TR = 5000 ms, TI = 1800 ms, TE = 395 ms, 1 × 0.49 × 0.49-mm resolution, AT = 7.02 minutes).

Location of WMH. The automated assessment of WMHs was computed in a previous study (Lampe et al., 2019). All images were checked by a study physician for incidental findings. A computer-based WMHs segmentation algorithm was then used to automatically determine WMH volume on T1-weighted MPRAGE and FLAIR.
images (Shiee et al., 2010) and inspected visually for segmentation errors. Binary WMH maps of all participants were non-linearly co-registered to a standardized MNI template (1-mm isometric) with ANTs (Avants et al., 2011). In standard space, binary subject-wise WMH maps were grand-averaged to create a population WMH frequency map (Jenkinson et al., 2012) and to further compute the voxel-wise statistics. As previously implemented (Lampe et al., 2019), to segregate the periventricular (pv)WMH and deep (d)WMH, a default distance of 10 mm to the ventricular surface was used (DeCarli et al., 2005). Every voxel of WMH located within this border was classified as pvWMH; voxels outside the border were classified as dWMH.

**WMH Volume.** Regional WMH volume was calculated separately for the deep and periventricular white matter. Following Lampe et al. (2019), we added a constant value 1 to every participant’s regional dWMH volume because there were participants without lesions in the deep WM. We then calculated the ratio of dWMH and pvWMH (dWMH/pvWMH) as localized WMH volume. Total, deep and periventricular WMH volumes were further normalized to head size by total intracranial volume. Total and localized WMH (dWMH/pvWMH) volume were log-transformed for further statistical analyses.

**2.3. EEG acquisition and preprocessing**

rsEEG activity was recorded in an electrically and acoustically shielded room using an EEG cap with 34 passive Ag/AgCl electrodes (EasyCap, Brain Products GmbH, Germany). Thirty-one scalp electrodes were placed according to the extended international 10–20 system. The signal was amplified using a QuickAmp amplifier, frequency range: DC-280 Hz (Brain Products GmbH, Germany). Two electrodes recorded vertical and horizontal eye movements while 1 bipolar electrode was used for electrocardiography. The rsEEG activity was referenced against common average and sampled at 1000 Hz. Impedances were kept below 10 kΩ. RsEEG data were preprocessed using EEGLAB toolbox (version 14.1.1b) and scripts were custom written in Matlab 9.3 (Mathworks, Natick, MA, USA). We filtered data between 1 and 45 Hz and applied a notch filter at 50 Hz. We then down-sampled the data to 500 Hz and ran a semiautomatic pipeline for artifact rejection: different noise threshold levels to mark bad time segments were used for the signal filtered in higher frequency (15–45 Hz) and lower frequency (1–15 Hz) ranges. The noise threshold for higher frequencies was set to 40 µV since noise at this range (i.e., induced by muscle activity) is typically lower in amplitude. The noise threshold for the lower frequency range was set to 0 ± 3SD over the mean amplitude of a filtered signal between 1 and 15 Hz. To control for the accuracy of automatically marked bad segments, we compared them to the noisy segments marked by another research group (Jawinski et al., 2017). Whenever these segments did not overlap by more than 10 s or they exceeded 60 s of total bad-segment duration, we inspected those datasets visually (~10% of cases) to confirm whether they indeed were contaminated by noise. We further visually assessed power spectral densities (PSD) for data quality and used it to identify broken channels. Next, using independent component analysis (Infomax; Bell and Sejnowski, 1995), activity associated with the confounding sources — namely eye-movements, eye-blinks, muscle activity, and residual heart-related artifacts — was removed.

**2.4. EEG sensor space analysis**

**2.4.1. Parameters of alpha oscillations**

For rsEEG analysis, we used the first 10 minutes of a recording to avoid the potential effect of participants’ drowsiness. We individually adjusted the alpha band frequency range by locating a major peak between 7 and 13 Hz on Welch’s PSD with 4 s Hanning windows. Thus, we determined individual alpha peak frequency in every channel and defined a bandwidth not exceeding 3 Hz around the peak. We then calculated relative alpha power for the individually adjusted alpha frequency range dividing it by the broadband power calculated in the 3–45 Hz frequency range. LRTC were calculated using detrended fluctuation analysis on the amplitude envelope (calculated with Hillbert transform) of alpha band oscillations in time windows ranging from 3 to 50 seconds (while respecting the boundaries where the bad segments had been cut) based on the previously published procedure (Hardstone et al., 2012). Here, we briefly repeat the main steps: (1) a cumulative sum of the amplitude envelope is calculated, (2) the signal is then divided into predefined window sizes (τ), (3) the linear trend is removed in a given window. Fluctuation function F(τ) for all time windows of a given size τ is calculated as the root-mean-square of the detrended signal. In the case of a power-law relationship, we have F(τ) ∝ τ^ν, where ν is a scaling exponent (measuring LRTC) which can be obtained as a slope of a linear fit in log-log plot between F(τ) and τ. An exponent of 0.5 reflects uncorrelated signals (i.e., resembling white noise), ν < 0.5 indicates anticorrelations, while an exponent between 0.5 < ν < 1 shows persistent autocorrelation where large fluctuations are likely to be followed by large fluctuations (Hardstone et al., 2012). This range of 0.5 < ν < 1 is a typical range for many EEG and MEG studies.

The illustration of parameters of alpha oscillations are shown in Fig. 2.

To reduce data dimensionality of rsEEG sensor space data used for the whole-brain voxel-wise inference analyses, we further grouped EEG channels into 6 coarser brain regions (frontal, central, temporal, parietal, and occipital) (Fig. 3A).

**2.5. EEG source space analysis**

To reconstruct sources of the rsEEG signal, we calculated lead-field matrices based on individual brain anatomies and standard
envelope of alpha oscillations in each voxel. The cortex surface mantle was divided into 68 regions of interest (ROIs) based on the Desikan-Killiany atlas (Desikan et al., 2006). These were further combined into 5 coarser ROIs (frontal, parietal, temporal, occipital, and cingulate) for the right and left hemispheres following a standard parcellation atlas (Fig. 3B). Alpha band measures (power, peak frequency, and LRTC) values were then averaged across each ROI.

2.6. Statistical analyses

2.6.1. Correlation of age with total WMH volume and alpha oscillations

Pearson correlations were calculated to examine the relationship between age and (i) total or localized WMH volume (dWMH/pvWMH) and (ii) the parameters of alpha oscillations in 6 regions at sensor space. Differences between correlations were assessed with Fisher’s r-to-z transformation implemented in R version 3.5.2 (http://www.R-project.org/). To correct for multiple comparisons, p values were then adjusted using the False Discovery Rate (FDR; Benjamini and Hochberg, 1995).

2.6.2. Topographical relevance analyses of WMHs for alpha oscillations at sensor space

To identify regions in which WMHs robustly correlated with alpha oscillations, we performed whole-brain voxel-wise regressions. More precisely, we applied general linear models (GLMs) in which individual values of relative alpha power, alpha peak frequency, and LRTC were used as predictors for the topographical occurrence of WMHs, adjusting for effects of age, sex, and intracranial volume as covariates of no interest. 3D voxel-wise binary lesion maps were analyzed using randomise function, implemented in FSL (Winkler et al., 2014). For each statistical analysis, positive and negative contrasts were computed. The significance of results was based on threshold-free cluster enhancement (TFCE, N = 10,000 permutations) with family-wise-error (FWE) corrected p values of 0.05. We further reported statistical results for the more conservative FWE threshold of p < 0.005.

2.6.3. Topographical relevance analyses of WMHs and alpha oscillations at source space

To assess the association between alpha oscillations at the EEG source level and whole-brain regional WMHs, we implemented GLMs separately for 10 ROIs with parameters of alpha oscillations covariate of interest, and age, sex, and total intracranial volume as covariates of no interest. All statistical analyses were further reported with multiple comparisons using TFCE based permutation testing (N = 10,000) at FWE level of p< 0.05, as well as with a conservative threshold of p< 0.005.

2.7. Sensitivity analyses

2.7.1. Confounding factors

Given that different cardiovascular risk factors including body mass index (BMI), systolic blood pressure (SBP), smoking, and diabetes are associated with WMHs (Habes et al., 2016; Lampe et al., 2019; Ryu et al., 2014), we further considered these factors as potential confounders (as covariates of no interest) for the voxel-wise associations between parameters of alpha oscillations and probability of WMH occurrence in the overall sample (N = 907). To assess a degree of collinearity between the regressors used in GLMs, we additionally computed variance inflation factor in R. All predictors had a variance inflation factor below 2, therefore, we concluded that models showed acceptably low multicollinearity.
2.7.2. Medication
We implemented the voxel-wise inference analyses between parameters of alpha oscillations and WMHs excluding participants taking medications affecting the central nervous system (opioids, hypnotics, and sedatives, antiparkinsonian drugs, anxiolytics, an- tipsychotics, antiepileptic drugs). The resulting sample included 801 individuals ($M = 68.96 \pm 4.58, 323$ female).

2.8. Control analyses

2.8.1. Absolute alpha power
To assess the robustness of our results, we further applied voxel-wise inference analyses between the probability of WMH occurrence and absolute alpha power in the left and right occipital region at EEG source space, using age, sex, and total intracranial volume as covariates of no interest. Absolute power in both regions was log-transformed to normalize the distribution of the data for statistical analyses.

2.8.2. Median split
A median split was performed based on the log-transformed total WMH volume using dichot function in R (N: higher WMH = 454, N: lower WMH = 453). We further computed Spearman correlations between total WMH volume (log-transformed, see: Supplementary Figure 2E) and relative alpha power separately for each group and each ROIs.

2.9. Mediation analyses
We performed mediation analyses using mediation package (Tingley et al., 2014) in R to test the association between a predictor (X), and an outcome (Y) which can be transmitted through a mediator (M) (Hayes and Rockwood, 2017). Here, we examine whether a total or localized WMH volume (M) mediates the relationship between age as a predictor (X) and parameters of alpha oscillations at sensor space as an outcome variable (Y). Bootstrapping (n = 5000) with 99% confidence intervals (CI) was used for testing the indirect effect because it does not assume normality in sampling distribution (Hayes and Rockwood, 2017). While the indirect effect shows whether age was associated with the parameters of alpha oscillations through a mediator, a total effect is the sum of indirect and direct effect. The indirect effect was considered significant if the corresponding 99% bootstrap CIs did not include zero.

2.10. Cognition
The Trail Making Test (TMT) is a cognitive test measuring executive function, including processing speed and mental flexibility (Reitan, 1955; Reitan and Wolfson, 1995). In the first part of the test (TMT-A) participants are asked to connect numbers in an ascending order, while in the second part (TMT-B), participants need to alternate between numbers and letters. In both TMT-A and B, the time to complete the task quantifies the performance, and lower scores indicate better performance.

We ran mediation analyses with 99% bootstrap CIs using relative alpha power in different regions as a predictor, total WMH volume as a mediator, and the task completion time in TMT-A or TMT-B as an outcome variable. The TMT data was available for 899 participants at the EEG sensor and 848 individuals at the EEG source space.

The regression coefficients ($\beta$) and their 95% CIs were estimate. Mediation analysis tests whether the association between a predictor (X), and an outcome (Y) is transmitted through a mediator (M). Mediation analysis tests whether the association between a predictor (X), and an outcome (Y) is transmitted through a mediator (M).

3. Results

3.1. Sample characteristics
Details about the demographic, anthropometric, cardiovascular measures, as well as WMH volume, and alpha oscillations can be found in Table 1. Histograms of total WMH volume, averaged relative alpha power, its peak frequency, and LRTC can be found in Supplementary Figure 2A–E.

3.2. Topography and characteristics of alpha oscillations
The relative alpha power at sensor space showed a maximum over the occipital channels, with a mean value of 0.66 ± 0.17 (%) (Supplementary Figure 3A). Similarly, the relative alpha power at source space showed a maximum over the bilateral occipital cortex, including cuneus and lateral occipital regions with a mean value of 0.59 ± 0.18 (%) (Supplementary Figure 3D). The grand-average peak frequency was 9.40 ± 0.49 Hz, showing larger values at occipital regions (Supplementary Figure 3B and 3E). The average scaling exponent ($v$) was 0.72 ± 0.017. Similarly, topographies of the scaling exponent had higher values at occipital and parietal areas as well as frontal regions (Supplementary Figure 3C and 3F).

3.3. Association of age with WMH volume and alpha oscillations
We found a correlation between age and total WMH volume ($r = 0.374, p < 0.001$, Supplementary Figure 4), but not with the dWMH/prWMH ($r = 0.03, p > 0.05$, Supplementary Figure 5). Regarding parameters of alpha oscillations, we found that higher age was associated with decreased alpha peak frequency all EEG ROIs ($r$ from -0.13 to -0.17, $p_{FWE} < 0.05$), while no correlations between age and relative alpha power or LRTC were found (all $p_{FDR} > 0.05$). A full report of these correlations for the entire sample and by sex are provided in Supplementary Figures 6–8.

3.4. Topographical association between WMHs and alpha oscillations

3.4.1. Sensor space
The voxel-wise inference analyses revealed that higher relative alpha power (%) in the frontal region was associated with higher WMH probabilities in the right body of corpus callosum ([116, -26, 32], $T = 3.76, k = 653$). Higher relative alpha power in the central region was associated with higher WMH probabilities in the right anterior thalamic radiation extending to the posterior corona radiata ([22, -49, 37], $T = 4.44, k = 2744$), while higher relative AP in the right temporal region was linked to higher WMHs in the right superior longitudinal fasciculus ([22, -49, 37], $T = 4.52, k = 6893$) extending to the left inferior fronto-occipital fasciculus ([131, -53, 32], $T = 4.00, k = 4210$). Furthermore, higher relative alpha power in the parietal region was associated with higher WMHs in the right superior corona radiata ([18, -19, 37], $T = 4.05, k = 4474$). Similarly, for relative alpha power in the occipital region, we observed a higher prevalence of WMHs in the bilateral superior corona radiata through the body of the corpus callosum to the anterior corona radiata, including the right anterior thalamic radiation ([18, -19, 37], $T = 4.39, k = 9450$). Accordingly, higher voxel-wise WMH probabilities were associated with higher relative alpha power independent of age, sex, and brain size, as shown in Fig. 4 (TFCE, $p < 0.05$, FWE-corrected). Note that using a more stringent TFCE, FWE rate of $p < 0.005$, the correlation between the probability of WMH occurrence and relative alpha power was only evident for the occipital region ([18, -19, 37], $T = 4.39, k = 904$). Finally, no voxel-wise associations between regional WMHs and al-
### Table 1
Sample characteristics

|                | Mean or n | Min. | Max. | SD  |
|----------------|-----------|------|------|-----|
| Age            | 69.49     | 60.15| 80.03| 4.63|
| Female / Male  | 380 / 527 |      |      |     |
| BMI (kg/m²)    | 27.59     | 18.68| 42.26| 3.97|
| SBP (mm Hg)    | 133.71    | 92.00| 200.5| 16.31|
| DBP (in mm Hg) | 74.54     |      | 43.5 | 120 |
| Never / former / active smokers | 517 / 319 / 71 |        |      |     |
| Diabetes (yes / no / unknown) | 143 / 748 / 16 |    |      |     |
| WMH volume (mm³) | 3935 | 127 | 78509 | 6676.76 |
| Normalized total WMH Volume | 0.0093 | 0.0003 | 0.170 | 0.015 |
| dWMH/pvWMH (%) | 0.439     | 0.011| 3.635 | 0.402|
| Intracranial volume (mm³) | 172981.1 | 1297219 | 2466529 | 147492.5 |
| Mean relative alpha power (%) | 0.55 | 0.21 | 0.88 | 0.15 |
| Mean alpha peak frequency (Hz) | 9.4 | 7.34 | 12.01 | 0.86 |
| Mean Scaling Exponent (β) | 0.73 | 0.53 | 1.14 | 0.093 |
| TMT A (s)      | 41.33     | 17.00| 126  | 13.32|
| TMT B (s)      | 89.29     | 25.00| 300  | 43.49|

Abbreviations: BMI, body mass index; DBP, diastolic blood pressure; dWMH/pvWMH, the ratio of deep/periventricular white matter hyperintensities; SBP, systolic blood pressure; SD, standard deviation; TMT, Trail Making Test; WMH, white matter hyperintensity.

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**Fig. 4.** Association between white matter hyperintensities (WMHs) and relative alpha power at EEG sensor space (N = 907). (A) Schematic depiction of the significant association between regional WMHs and relative alpha power: thicker lines indicate higher t-values. (B) We implemented nonparametric permutation testing based on whole-brain voxel-wise analysis to investigate the association between WMHs and relative alpha power (%). The brain WMH clusters show significant relation with the EEG frontal region (purple), central region (orange), right temporal region (yellow), parietal region (green), and occipital region (blue), respectively (TFCE, FWE-corrected, p < 0.05 corrected for age, sex, and total intracranial volume). (C) Scatter plots show the association between WMH probability (x-axis) extracted from clusters based on significant whole-brain voxel-wise inference analyses and elevated relative alpha power (y-axis) in different EEG regions. The resulting statistical images (P-map) were further thresholded at 0.05 and binarized (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article). Abbreviations: A, anterior; L, left; P, posterior; R, right.

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The peak frequency or LRTC were observed (TFCE, p < 0.05, FWE-corrected).

#### 3.4.2. Source space

We found that a higher probability of WMH occurrence was associated with higher relative alpha power (%) in all EEG regions except for the left frontal region (Supplementary Table 2, TFCE, p < 0.05, FWE-corrected, Fig. 5). With the stricter FWE-level of p < 0.005, the association between the occurrence of WMHs and relative alpha power was evident for left [[18, -19, 37], T = 4.29, k = 192] and right occipital regions [[18, -19, 37], T = 4.45, k = 845]. We further observed a negative correlation of regional WMHs with individual alpha peak frequency in right temporal [-26, 29, 35], T = 4.25, k = 2086], left frontal [-26, 28, 35], T = 3.87, k = 1143], left cingulate [-26, 28, 35], T = 3.99, k = 821], right [-31, 49, 29], T = 4.89, k = 222] and left occipital regions [[31, 49, 29], T = 4.77, k = 119] (TFCE, p < 0.05, FWE-corrected). However, these associations were no longer significant after correcting with the conservative FWE-level of p < 0.005. Finally, confirming sensor space analyses, no voxel-wise associations of regional WMHs with LRTC were observed (TFCE, p < 0.05, FWE-corrected).

#### 3.5. Sensitivity analyses

#### 3.5.1. Confounding factors

Voxel-wise inference analyses after controlling for age, sex, intracranial volume, BMI, SBP, diabetes, and smoking status yielded a similar relationship between higher WMH probability and elevated relative alpha power in the following regions: central [[22, -49, 37], T = 4.46, k = 5417], right temporal [[22, -49, 37], T = 4.52,
k = 5417), left temporal ([122, −49, 37], T = 4.59, k = 4772), parietal ([118, −19, 37], T = 3.68, k = 231), and occipital ([118, −19, 37], T = 4.08, k = 4018) EEG regions across the overall sample. Note that with TFCE, FWE-corrected, p < 0.005, we did not find any clusters. Lastly, no WMH clusters were related to alpha peak frequency or LRTC (TFCE, p > 0.05, FWE-corrected).

3.5.2. Medication

Voxel-wise inference analyses excluding individuals taking central nervous system medication (N = 801) still indicated the association between higher prevalence of WMHs and increased relative alpha power at sensor space in the following regions: frontal ([17, 9, 31], T = 4.42, k = 6880), central ([20, −30, 35], T = 4.46, k = 9063), right temporal ([20, −48, 35], T = 4.57, k = 12098), left temporal ([22, −49, 37], T = 4.61, k = 9408), parietal ([14, −8, 31], T = 4.61, k = 9054), and occipital ([18, −19, 37], T = 4.44, k = 12,885) EEG regions. Importantly, with TFCE, FWE-corrected, p < 0.005, we identified WMHs clusters (k > 2000) for occipital, left temporal, right temporal, and a small cluster (k > 200) for parietal and central EEG regions. Additional voxel-wise inference analyses revealed that higher WMHs resulted in decreased alpha peak frequency in right temporal ([17, −27, 33], T = 4.00, k = 138) and left temporal regions ([17, −27, 33], T = 4.12, k = 503). Lastly, no WMHs clusters in the brain were related to LRTC (TFCE, p > 0.05, FWE-corrected).

3.6. Control analyses

3.6.1. Absolute alpha power

Voxel-wise inference analyses with absolute alpha power similarly indicated that higher probability of WMH occurrence was associated with elevated absolute alpha power in right ([−23, 0, 36], T = 3.98, k = 5633) and left occipital regions ([−23, 0, 36], T = 4.05, k = 5358) (TFCE, p < 0.05, FWE-corrected).

3.6.2. Median split

The correlation analyses based on the median split similarly revealed a positive association between relative alpha power and total WMH, the correlation values were between 0.07 and 0.14. The scatter plots of each EEG ROIs for higher and lower WMH groups can be found in Supplementary Figure 8.

3.7. Mediation analyses

We examined whether total or localized (dWMH/pvWMH) WMH volume could mediate the relationship between age and relative alpha power in all cortical ROIs. Investigating the relationship between age and relative alpha power, we observed a significant indirect effect of total WMH volume in most of the cortical regions defined at sensor space (Table 2). The direct effect was not significant in any of the ROIs (99% CI > 0), and only in the right temporal region at sensor space did the total effect of age on relative alpha power appear to be significant (Table 2). Further, we confirmed the indirect effects of total WMH volume for relative alpha power at EEG source space for left parietal (β = 0.0012, CI = [0.00006–0.0022]), left (β = 0.0014, CI = [0.00013–0.0022]) and right occipital (β = 0.0014, CI = [0.00015–0.0028]) regions, but not for the left and right temporal regions. Finally, our results revealed that neither total nor localized WMH volume mediated the association of age with alpha peak frequency and LRTC at sensor space (all p > 0.05).

3.8. Cognition

Compared to population-based norms (Hobert et al., 2011; Tombaugh, 2004), our sample shows similar TMT scores (Table 1), indicating good to intermediate cognitive performance. We then investigated the question of whether the relationship between relative alpha power and cognition measured by task completion time in TMT-A and B is mediated by total WMH volume. After controlling for age and sex, we found a significant indirect effect of total WMH volume on the association between TMT-A and relative alpha power only in the right temporal region (β = 1.071, CI = [0.123–2.539]). In TMT-B, we observed a significant indirect effect of total WMH volume for the frontal region (β = 3.399, CI = [0.252–7.896]), as shown in the Supplementary Table 2. At EEG source space, however, we did not confirm these findings. More precisely, we did not observe any indirect effects of alpha power in the right temporal region on TMT-A and in the left and right frontal region on TMT-B. Further, in all analyses with EEG-source space, the direct and total effects were not significant.

4. Discussion

The main goal of this study was to investigate whether regional WMHs affect parameters of alpha oscillations independent from age. We pursued this aim using a large sample of cognitively healthy older individuals (e.g., also based on TMT scores; Hobert et al., 2011; Tombaugh, 2004) from a population-based study (Loeffler et al., 2015). We showed distinct regional relationships between relative alpha power and WMHs: our topographical analysis suggested that higher occurrence of WMHs in superior, posterior, and anterior corona radiata, as well as thalamic radiation, was related to higher relative alpha power, with strongest correlations in the bilateral occipital cortex. Adjusting for potential confounding factors including age, cardiovascular risk factors, or controlling for the effect of medication did not change these results. While the direct link between age and alpha power assessed by correlation analyses was absent, mediation analyses supported an indirect link for the existence of the relation between age and
alpha power through the total WMH volume. This finding indicates why we should consider the age-related structural changes in the brain (e.g., WMHs) when we investigate the aging effects on EEG neural oscillations.

Alpha rhythm is the most salient rEEG oscillatory phenomenon that originates from thalamocortical and cortico-cortical interactions (Bazanova and Vernon, 2014; Lopes Da Silva et al., 1997). Alterations in alpha oscillations have previously been linked to changes in different anatomical features including properties of WM (e.g., Valdés-Hernández et al., 2010). Regarding WMHs, for instance, a previous EEG-MRI study showed that higher relative alpha power in parietal regions was associated with higher scores of the prevalence of WMHs in 79 individuals with mild cognitive impairment (Babiloni et al., 2008a), consistent with our findings in this population-based sample. Previous studies with computational models have given further support for the notion that resonance properties of feedforward, cortico-thalamocortical, and intracortical circuits substantially influence alpha oscillations (Hindriks and van Putten, 2013). In the present study using a larger sample, we similarly observed that regional WMHs, detected mostly in superior corona radiata, containing thalamocortical fibers, affect interindividual differences in relative alpha power. This finding was further reproduced when using alpha power values extracted from EEG source-based analysis. Although we did not observe significant association between those 2 measures after controlling for other confounding factors at stricter threshold (TFCE, FWE < 0.005), the consistent results with regular FWE threshold at voxel-wise level suggest a possible neurophysiological link between WMHs and relative alpha power.

But, how could lesions in the WM possibly affect EEG signal which mainly reflects neural synchrony within gray matter? While in principle a hyperintensity in T2-weighted MR sequences is a quite unspecific marker of various pathologies, postmortem histopathological studies of older adults with WMHs have mostly reported demyelination, axonal loss, and other consequences of ischemic small vessel disease (Smith et al., 2000; Wardlaw et al., 2015). Myelin contributes to the speed of impulse conduction through axons, and the synchrony of impulses between distant cortical regions (Fields, 2015, 2008). Reductions of conduction velocity due to demyelination and loss of (communicating) axons are assumed to be responsible for cognitive dysfunctions which are known to be based on delicately orchestrated propagations of neuronal signals. Electrophysiological, interactions, and synchrony between neuronal populations are reflected in rhythmic M/EEG signals, of which alpha oscillations are the most prominent ones (Bazanova and Vernon, 2014; Lopes Da Silva et al., 1997). Alpha power is a quantitative marker of the degree of synchrony in the neuronal activity of the corresponding neuronal populations (Pfurtscheller and Lopes Da Silva, 1999). While for a long-time alpha oscillations were regarded as idle rhythms of non-active brain areas, a plenitude of studies has convincingly demonstrated that alpha oscillations play an important role in many cognitive functions (Fox et al., 2016; Klimesch, 1999; Palva and Palva, 2007). For instance, in motor and sensory domains it has been shown that amplitude decreases of alpha oscillations in focal areas (i.e., reflecting cortical activation) is in turn associated with the inhibition of neighboring cortical areas (Pfurtscheller and Lopes Da Silva, 1999). This phenomenon is thought to include mutually inhibitory interactions between the chain of modules including thalamocortical and reticular nucleus neurons which are involved in the generation of alpha oscillations (Sufczynski et al., 2001). Importantly, the authors hypothesized that this surround inhibition should underlie other cognitive operations such as focused attention and stimulus selection. Such topographically specific relationships are likely to be disturbed by the alterations in conduction velocity and ax-
onal loss in the thalamocortical circuitry (Pajevic et al., 2014). As a result of such WM disturbances, a modular organization of thalamocortical inputs and a corresponding demarcation between cortical patches of enhanced and attenuated alpha oscillations could be abolished, thus leading to a larger spread of alpha oscillations across the cortex and consequently to stronger and spatially less specific alpha oscillations. This in turn might explain a positive association between alpha power and WMHs. The hyperactivation of alpha with WMHs could also be ineffective in preserving cognitive performance or even reflect the progression of neurodegenerative alterations (Corriveau-Lecavalier et al., 2019; Pons et al., 2010).

Despite a number of reports of age-related alpha power alterations (Babiloni et al., 2006b; Lodder and van Putten, 2011; Vysata et al., 2012), in our study, we replicated other recent studies (Sahoo et al., 2020; Scally et al., 2018) which did not find strong evidence for age-related attenuations of relative alpha power. The discrepancy in findings with earlier reports could be due to the narrow age range of our participants, as well as the individually adjusted alpha frequency range based on the peak frequency. In fact, preserved peak power at peak frequency has recently been reported in an older sample (Scally et al., 2018), suggesting that any observed age-dependent power changes might be due to shifts in the frequency range at which alpha peak occurs. While our cross-sectional dataset cannot provide unequivocal evidence for a causal relationship, mediation analyses demonstrated a presence of an indirect relationship between age and alpha power through total WMHs. Currently, in the literature, there is an ongoing discussion on the interpretation and meaning of an indirect (mediation) effect when a total effect is not statistically significant (Hayes and Rockwood, 2017; Zhao et al., 2010). In our paper, following the suggestions by Hayes and Rockwood (2017), we also reported and interpreted the mediation effects even when a total effect was not significant. More precisely, the mediation via total WMH volume showed that higher age was associated with the elevated relative alpha power in the right temporal, parietal, and occipital regions. As mentioned before, age-related reductions of alpha power in occipital regions were previously reported in different sample populations (see detailed review: Ishii et al., 2018). As we show in this study, in healthy older adults the association between these 2 measures can potentially be mediated by WMH volume thus demonstrating a positive relationship between alpha power and age. Therefore, our result shows why one should potentially consider structural correlates when investigating age-related alterations in neural oscillations.

In the literature, other commonly reported age-dependent changes in spectral parameters of EEG include slowing of the alpha peak (Knyazeva et al., 2018). We replicated the slowing of the alpha peak frequency with increasing age despite the narrow age range. Alpha peak slowing has previously been suggested to be linked to a less efficient coordination of neuronal activity in this frequency range (Mierau et al., 2017). We further observed that slowing of alpha peak with age was independently associated with higher WMHs. As EEG oscillations reflect a complex circuitry of cortical inhibitory and excitatory neurons, it may be possible that the age-related reduced activation of these neurons due to acetylcholine or synaptic damage may lead to a lowering of alpha peak (Gouw et al., 2017). However, our results should be interpreted with caution since we only observed significant association between these 2 measures in EEG-source but not sensor space. Finally, we explored the relationship between age and LRTC in the amplitude envelope of alpha oscillations that capture scale-free dynamics of resting-state oscillations. LRTC has previously been linked to the presence of a critical state in neural networks, which is characterized by the balance of excitation and inhibition (Poil et al., 2012) that has been suggested to be optimal for the processing of information in the human brain. Regarding the association between age and LRTC, previous studies have shown that the observed age-related changes might be dependent on age range — it increases from childhood to early adulthood, after which it stabilizes (Nikulin and Brismar, 2005; Smit et al., 2011). In accordance with these previous findings, in our sample of older adults, we observed no pronounced age-related LRTC attenuations.

5. Limitations

While a strength of this study is the large population-based sample, the study design is cross-sectional and does not allow making inferences about the directionality of the association between WMHs and alpha oscillations. Longitudinal studies are required to further clarify these associations. Research using other advanced techniques such as quantitative MRI or specific assessment of tissue properties with ultra-high field MRI combined with intracranial EEG recording could further provide valuable insights into the nature of the relationship between WM properties and alpha oscillations. We performed a relatively coarse parcellation of the brain at EEG source space analysis due to the relatively small number of electrodes (n = 31). A denser spatial sampling of the EEG (not available in the present cohort) would allow investigation of this relationship with better spatial precision. Our approach of investigating the relationship between EEG and MRI-based measures (parameters of alpha oscillations and WMHs) and age can readily be extended to power changes in other frequency bands (e.g., delta or theta), EEG features (e.g., connectivity or phase synchrony; Hinault et al., 2020; Quandt et al., 2020), and other brain anatomical features (e.g., diffusion tensor imaging), both under task or resting-state conditions. It therefore opens new questions for investigating the brain structure and function relationship at potentially more detailed temporal and spatial scales. Finally, we had no specific hypothesis targeting regionally specific WMHs and, even though, our whole-brain search revealed statistically significant and interpretable results, our findings should be regarded as exploratory.

6. Conclusion

Using sensitive high-resolution neuroimaging techniques in cognitively healthy older adults (N = 907), we showed that higher probability of WMHs is related to relative alpha power. Importantly, our study provides evidence that the changes in alpha oscillations do not relate to aging per se but rather depend on the impact of age-related neuropathology, such as WMHs. Our findings thus suggest that longitudinal EEG recordings might be sensitive for the detection of alterations in neuronal activities due to progressive structural changes in WM.

Disclosure statement

Stefan Haufe is funded by the European Research Council (ERC) under the European Union’s Horizon 2020 research and innovation programme (Grant agreement No. 758985). The remaining authors report no conflict of interest.

Acknowledgements

We thank all members of the Leipzig Research Center for Civilization Diseases (LIFE) study center for conducting the LIFE-Adult study and also all participants for their valuable collaboration. This work is supported by the European Union, the European Regional
