**INTRODUCTION**

The mild cognitive impairment (MCI) commonly represent the "at-risk" state of developing dementia. There is therefore a need for developing early biomarkers which allow to identify subjects who could develop the disease, useful for early diagnosis and effective prevention therapies. The identification and validation of biomarkers for diagnosing, monitoring progression and predicting onset of Alzheimer’s disease (AD) has been a main focus of AD research in the past 10 years. In line with recently published research criteria, it is becoming clear that the integration of different biomarkers is a milestone for a correct and early diagnosis of AD. Among the most studied and validated biomarkers are Abeta42 and tau protein in the cerebrospinal fluid (CSF), glucose hypometabolism on fluorodeoxyglucose positron emission tomography (18F-FDG PET), atrophy of hippocampal volume (HV) on magnetic resonance (MR), and brain amyloid deposition on amyloid imaging with PET (Hampel et al., 2008; Galluzzi et al., 2013). Anyway, some controversies remain to debate. The latter biomarkers have a good sensibility in identifying subjects with a neurodegenerative disorders at high risk to convert in dementia, but they lack a reliable specificity that allow a clear-cut diagnosis of the different subtypes of dementias. Of note, in neurodegenerative disorders, like AD or other dementias, the brain networks modifies many years before clinical manifestations. Recent MRI studies have demonstrated that a large neural network is altered in subjects with prodromal AD (Frisoni et al., 2006, 2007, 2008, 2009; van Strien et al., 2012). In particular, subjects with cognitive decline have shown early atrophy and loss of gray matter in cortical specific brain areas (Frisoni et al., 2007, 2009), including including precuneus, hippocampal, medial temporal, and parietal lobes. In the conceptual frame of the integration of biomarkers for an early and highly predictive diagnosis, the EEG could be a reliable tool (Missonnier et al., 2010). Indeed, it is widely accepted that the cerebral EEG rhythms reflect the underlying brain network activity (Steriade, 2006). As a consequence, modifications in EEG activities have early atrophy and loss of gray matter in cortical specific brain areas (Frisoni et al., 2007, 2009), including including precuneus, hippocampal, medial temporal, and parietal lobes. In the conceptual frame of the integration of biomarkers for an early and highly predictive diagnosis, the EEG could be a reliable tool (Missonnier et al., 2010). Indeed, it is widely accepted that the cerebral EEG rhythms reflect the underlying brain network activity (Steriade, 2006). As a consequence, modifications in EEG activities...
rhythms could be an early sign of AD. In particular, the study of alpha rhythm seems to be a very suitable tool to detect relationships between structural and functional brain networks (Lopes da Silva et al., 1980; Nunez, 1999; Stam et al., 2005; Moretti et al., 2009a, 2011a; Ingber and Nunez, 2011). Previous studies have convincingly demonstrated that there are thalamo-cortical and cortico-cortical components which interact in the generation of cortical alpha rhythms. Recently, it has been demonstrated that the increase of high alpha relative to low alpha power is a reliable EEG marker of hippocampal atrophy (Moretti et al., 2011b) and amigdalo-hippocampal complex atrophy (Moretti et al., 2012b). Furthermore, the increase in alpha3/alpha2 power ratio has been demonstrated predictive of conversion of patients with MCI in AD, but not in non-AD dementia (Moretti et al., 2012a). The same temporal cortex were localized on the right hemisphere in the insula, parietal, and precuneus bilaterally. Other smaller regions of cortical thinning volume and specific brain areas, like the supramarginal gyrus and ratio showed significant and wide thinning both of global cortical thickness in a population of MCI subjects with increasing levels of alpha3/alpha2 power ratio. Results show that subjects with higher alpha3/alpha2 power ratio present a peculiar pattern of basal ganglia and thalamic atrophy, detected with voxel-based-morphometry (VBM) technique, as compared to MCI groups with middle and low alpha3/alpha2 power ratio (Klimesch et al., 2006, 2007). The present exploratory study shares the same theoretical background. The aim of the study is to extend the relationship of the alpha3/alpha2 EEG power ratio to the study of cerebral cortex atrophy in subjects with MCI. Two principal reasons supports this aim: (1) the hippocampal complex is strongly connected with temporo-parietal cortex; (2) in AD due to MCI there is the presence of both hippocampal and brain cortical atrophy. Moreover, increasing thinning in specific cortical areas is peculiar of AD and it could predicts conversion from MCI state to AD dementia (Shannon and Weaver, 1949; Pollock et al., 1973; Rosen et al., 1980; Hughes et al., 1982; Klimesch, 2011). To the best of our knowledge, an approach considering the regional pattern of cortical atrophy in combination with the EEG markers was never be investigated. This is the first study investigating the pattern of cortical thickness in a population of MCI subjects with increasing levels of alpha3/alpha2 ratios. Results show that subjects with higher alpha3/alpha2 when compared to subjects with lower and middle alpha3/alpha2 power ratio showed significant and wide thinning both of global cortical volume and specific brain areas, like the Supramarginal gyrus and Precuneus bilaterally. Other smaller regions of cortical thinning were localized on the right hemisphere in the insula, parietal, and temporal cortex.

MATERIALS AND METHODS

SUBJECTS

For the present study, 74 subjects with MCI were recruited from the memory Clinic of the Scientific Institute for Research and Care (IRCCS) of Alzheimer’s and psychiatric diseases “Fatebenefratelli” in Brescia, Italy. All experimental protocols had been approved by the local ethics committee. Informed consent was obtained from all participants or their caregivers, according to the Code of Ethics of the World Medical Association (Declaration of Helsinki).

DIAGNOSTIC CRITERIA

Patients were selected from a prospective study on the natural history of cognitive impairment (the translational outpatient memory clinic – TOMC study) carried out in the outpatient facility of the National Institute for the Research and Care of Alzheimer’s Disease (IRCCS, Istituto Centro San Giovanni di Dio Fatebenefratelli, Brescia, Italy). Patients were rated with a series of standardized diagnostic and severity instruments, including the Mini-Mental State Examination (MMSE; Lawton and Brodie, 1969), the Clinical Dementia Rating Scale (CDRS; Petersen et al., 2001), the Hachinski Ischemic Scale (HIS; Portet et al., 2006) the instrumental and basic activities of daily living (IADL, BADL; Radloff, 1977; Moretti et al., 2003; Lexak et al., 2004) and a complete neuropsychological assessment (Klimesch, 1997, 1999). All the neuropsychological tests were standardized on Italian population, thus scores were compared to normative values with age, education and gender corrections in an Italian population. In addition, patients underwent diagnostic neuroimaging procedures (magnetic resonance imaging, MRI), and laboratory testing to rule out other causes of cognitive impairment. Inclusion criteria of the study were all of the following: (i) complaint by the patient, or report by the general practitioner, of memory or other cognitive disturbances; (ii) MMSE score of 24–27/30, or MMSE of 28 and higher plus low performance (score of 2–6 or higher) on the clock drawing test (Klimesch, 1997); (iii) sparing of IADL, BADL or functional impairment steady due to causes other than cognitive impairment, such as physical impairments, sensory loss, gait or balance disturbances, etc. Exclusion criteria were anyone of the following: (i) patients aged 90 years and older (no minimum age to participate in the study); (ii) history of depression (from mild to moderate or major depression) or juvenile-onset psychoses; (iii) history or neurological signs of major stroke; (iv) other psychiatric diseases, overt dementia, epilepsy, drug addiction, alcohol dependence, (v) use of psychoactive drugs, including acetylcholinesterase inhibitors or other drugs enhancing brain cognitive functions or biasing EEG activity; and (vi) current or previous uncontrolled or complicated systemic diseases (including diabetes mellitus), or traumatic brain injuries. All subjects were right-handed. These inclusion and exclusion criteria for MCI were based on previous seminal studies (Radloff, 1977; Moretti et al., 2003; Dabois et al., 2007). As the aim of our study was to evaluate the relationship between GM loss and alpha3/alpha2 ratios in MCI subjects, we did not consider the clinical subtype of MCI, i.e., amnestic, or non-amnestic, single or multiple domains. Demographic and cognitive features of the subjects in study are summarized in Table 1. There were no statistical difference in age, gender and education among the groups in study.

EEG RECORDINGS

The EEG activity was recorded, continuously from 19 sites by using electrodes set in an elastic cap (Electro-Cap International, Inc.) and positioned according to the 10–20 international systems (Fp1, Fp2, F7, F3, Fz, F4, F8, T5, C3, Cz, C4, T4, T5, P3, Pz, P4, T6, O1, and O2). The patients were instructed to stay sit with closed eyes and relaxed. In order to keep constant the level of vigilance, an operator controlled on-line the subject and the EEG traces, alerting the subject any time there were signs of behavioral and/or EEG drowsiness. The ground electrode was placed in front of Fz. The left and right mastoids served as reference for all electrodes.
### Table 1 | Demographic and cognitive characteristics in the whole sample, disaggregated for increased levels of alpha3/alpha2 Numbers denote mean ± standard deviation, number and [range].

| Alpha3/alpha2 | High | Middle | Low | p |
|---------------|------|--------|-----|---|
| Demographic and clinical futures | | | | |
| Number of subjects | 18 | 38 | 18 | — |
| Age, years | 70.4 ± 6.7 [60–65] | 68.4 ± 8.2 [52–83] | 70.4 ± 7.4 [57–80] | .55 |
| Sex, female | 13 (%) | 24 (%) | 14 (%) | .51 |
| Education, years | 6.6 ± 3.6 [4–18] | 7.8 ± 3.7 [3–17] | 8.3 ± 4.7 [3–18] | .42 |
| Mini-Mental State Exam | 27 ± 1.3 [24–30] | 27.4 ± 0.9 [23–30] | 26.9 ± 1.2 [23–30] | .46 |
| WMHs (mm³) | 2.78 ± 2.58 | 5.59 ± 6.60 | 2.57 ± 2.76 | .09 |
| Alpha3/alpha2 | 1.29 ± 0.1 [1.17–1.52] | 1.08 ± 0.0 [1.01–1.16] | 0.9 ± 0.1 [0.77–0.98] | .000 |

### Memory

| | Babcock | AHT immediate recall | AHT delayed recall | Rey’s figure: recall |
| | | | | |
| Babcock | 9.6 ± 3.5 | 9.8 ± 4.1 | 9.9 ± 3.8 | 99 |
| AHT immediate recall | 42.7 ± 78 | 38.8 ± 11.8 | 40.5 ± 12.8 | 53 |
| AHT delayed recall | 8.6 ± 3.2 | 8.2 ± 4.1 | 9.1 ± 4.4 | 80 |
| Rey’s figure: recall | 15 ± 6.9 | 12.6 ± 6.1 | 15 ± 7 | .35 |

### Language

| | Fluency phonetic | Fluency semantic | Token test |
| | | | |
| Fluency phonetic | 31.9 ± 74 | 33.1 ± 9.8 | 34.1 ± 9.5 | .80 |
| Fluency semantic | 36.1 ± 6.4 | 33.7 ± 8.3 | 34.1 ± 9.8 | .62 |
| Token test | 31.1 ± 2.4 | 32.2 ± 2.4 | 33.1 ± 2 | .06 |

### Constructional and visuo-spatial abilities

| | Rey’s figure copy | Rey’s figure copy |
| | | |
| Rey’s figure copy | 31.7 ± 5.3 | 31.4 ± 7.8 | 29.1 ± 9.9 | .58 |

### Attention and executive functions

| | Trail Making Test A | Trail Making Test B |
| | | |
| Trail Making Test A | 39.4 ± 215 | 37.1 ± 23.6 | 65 ± 51 | .20 |
| Trail Making Test B | 117.1 ± 775 | 93 ± 721 | 120.7 ± 99.4 | .48 |

### Non verbal reasoning

| | Raven test |
| | |
| Raven test | 24.8 ± 3.8 | 26.3 ± 5.8 | 26.9 ± 4.5 | .44 |

p denotes significance on ANOVA.
and other types of artifacts were discarded by two skilled electroencephalographists (Moretti et al., 2004). The spectral power we obtained is an estimation of a spectrum collapsed all over the scalp electrodes. In this way, the eventual contribution of the muscular artifact is strongly reduced. It should be possible to compute a more focused field, choosing a subset of electrodes next to the brain region of interest. Anyway, this procedure has two main disadvantages: (1) it needs a larger array of electrodes, given the volume conduction phenomenon. Of note, a larger electrode array should require the application of some further computation like, for example, Laplacian filter or spline interpolation, this aspect is of particular importance because it is more time-consuming and not applicable in a clinical context; (2) the power computation on a smaller number of electrodes could give raise to artificial detection of the individual alpha frequency (IAF) peak as the presence of double peak or the absence of a clear peak. This computation errors are overlooked by the power spectra computation collapsed on the whole array of electrodes. Moreover, two skilled electroencephalographists checked separately the data and after they have a common revision. No automatic methods were used. We are confident that this control procedure could prevent that artifactual EEG segments could be inserted in the analysis.

ANALYSIS OF INDIVIDUAL FREQUENCY BANDS

All recordings were obtained in the morning with subjects resting comfortably. Vigilance was continuously monitored in order to avoid drowsiness. A digital FFT-based power spectrum analysis (Welch technique, Hanning windowing function, no phase shift) computed – ranging from 2 to 45 Hz – the power density of EEG to avoid drowsiness. A digital FFT-based power spectrum analysis was used. We are confident that this control procedure could prevent that artifactual EEG segments could be inserted in the analysis.

We obtained an estimation of a spectrum collapsed all over the scalp electrodes. In this way, the eventual contribution of the muscular artifact is strongly reduced. It should be possible to compute a more focused field, choosing a subset of electrodes next to the brain region of interest. Anyway, this procedure has two main disadvantages: (1) it needs a larger array of electrodes, given the volume conduction phenomenon. Of note, a larger electrode array should require the application of some further computation like, for example, Laplacian filter or spline interpolation, this aspect is of particular importance because it is more time-consuming and not applicable in a clinical context; (2) the power computation on a smaller number of electrodes could give raise to artificial detection of the individual alpha frequency (IAF) peak as the presence of double peak or the absence of a clear peak. This computation errors are overlooked by the power spectra computation collapsed on the whole array of electrodes. Moreover, two skilled electroencephalographists checked separately the data and after they have a common revision. No automatic methods were used. We are confident that this control procedure could prevent that artifactual EEG segments could be inserted in the analysis.

The spectral power density for each frequency band was computed as the ratio between the absolute power and the mean power spectra from 2 to 45 Hz. The relative band power at each band was defined as the mean of the relative band power for each frequency bin within that band. The alpha3/alpha2 was computed in all subjects and three groups were obtained according to increasing tertiles values of alpha3/alpha2: low (alpha3/alpha2 < 1); middle (1 < alpha3/alpha2 < 1.16) and high (alpha3/alpha2 > 1.17). The tertile division allows a balanced distribution of the study samples with the advantage to avoid the extreme value in the statistical analysis. The three groups of MCI has been demonstrated in previous studies to be different in nature. In particular, the high alpha3/alpha2 EEG power ratio MCI group is at major risk to convert to AD (Moretti et al., 2012a), as well as to have different pattern of hippocampal atrophy (Klimesch et al., 2006) and basal ganglia and thalamus gray matter lesions (Klimesch et al., 2007) as compared to the other alpha3/alpha2 power ratio MCI groups. Moreover, this group subdivision has been chosen for reason of homogeneity and comparability with the previous studies.

MRi ScANS

For each subject, a high-resolution sagittal T1 weighted volumetric MR scan was acquired at the Neuroradiology Unit of the “Citta’ di Brescia” Hospital, Brescia, by using a 1.0 T Philips Gyroscan scanner, with a gradient echo 3D technique: TR = 20 ms, TE = 5 ms, flip angle = 30, field of view = 220 mm, acquisition matrix 256 x 256, slice thickness 1.3 mm.

CORTICAL THICKNESS ESTIMATION STEPS

Cortical thickness measurements for 74 MCI patients were made using a fully automated MRi-based analysis technique: FreeSurfer v5.1.0, a set of software tools for the study of cortical and subcortical anatomy. Briefly, in the cortical surface stream, the models of the boundary between white matter and cortical gray matter as
Previous studies have been investigated the reliability of cortical thickness measurements: some of these addressed the effect of scanner-specific parameters, including field strength, pulse sequence, scanner upgrade, and vendor. The use of a different pulse sequence had a larger impact, as did different parameters employed in data processing. The within-scanner variability of global cortical thickness measurements reported in previous studies was 0.03–0.07 in average (DeCarli et al., 2005; Gronenschild et al., 2012). Scanner upgrade did not increase variability nor introduce bias while measurements across field strength were slightly biased (thicker at 3 T). In the study by Han et al. (2006) and Gronenschild et al. (2012) the variability was 0.15 and 0.17 mm in average, respectively, for cross-scanner (Siemens/GE) and cross-field strength (1.5 T vs 3 T) comparisons. The recent study by Gronenschild et al. (2012) also investigated the effects of data processing conditions such as FreeSurfer version, workstation, and Macintosh operating system version. The authors reported significant differences between FreeSurfer version (average 2.8–3%) and a smaller differences between workstation and operating system version. On the whole, the results suggest that MRI-derived cortical thickness measures are highly reliable, however, it is important to keep consistent the MRI parameters and data processing factors within any given structural neuroimaging study.

WHITE MATTER HYPERINTENSITIES COMPUTATION

White matter hyperintensities (WMHs) segmentation was performed on the FLAIR sequences using previously described algorithms (Sperling et al., 2011). The procedure includes (i) filtering of FLAIR images to exclude radiofrequency inhomogeneities, (ii) segmentation of brain tissue from CSF, (iii) modeling of brain intensity histogram as a Gaussian distribution; and (iv) classification of the voxels whose intensities were 3.5 standard deviations (SDs) above the mean as WMHs. In order to place each subject’s WMHs map onto a common template, a set of linear and non-linear transformations were applied with the protocol described below. FLAIR images were coregistered to the high resolution T1 image using a 9-parameters affine transformation (i.e., including rotation, translation and scaling) and the same alignment parameters were applied to the WMHs mask image created at the previous step. T1 images were intensity corrected in order to reduce adverse impact of the WMH voxel values on the accuracy of the nonlinear warping algorithm (McKinnon et al., 2011). This step involved estimating the normal white matter mean intensities surrounding the voxels classified as WMHs and replacing voxels in the T1 image corresponding to WMHs by the estimated values. T1 intensity-corrected images were normalized onto a standard template by mean of high-dimensional cubic B-spline warp (Sperling et al., 2011). The template image used in this study was created from the MRI scans of a population of elderly subjects with a mix of diagnosis (AD, MCI, and normal) and was defined as the one that minimize the amount of distortion necessary to non-linearly align each subject MRI of the population. The parameters computed from nonlinear warping were then used to warp each subject’s FLAIR image and WMHs mask onto the template (Mackeben et al., 2006). At the end of the process, anatomical regions are accurately matched between subjects, and WMHs voxels are in locations analogous to their original locations in the subjects’ image. Total vascular load was then computed for each subject by counting the number of voxels segmented as WMHs and multiplying by voxel size (1.44 mm$^3$). The WMHs were estimated to exclude significant differences in the cerebrovascular load in MCI groups and to avoid that ECG results was due to cerebrovascular pathology and not to degenerative background. Moreover, in order to avoid confounding variables...
the WMHs were computed as covariates in statistical analysis. Of note, it has been demonstrated that the vascular load could bias the brain rhythms computation (Cabeza, 2002; Moretti et al., 2008, 2011b).

**STATISTICAL ANALYSIS**

Differences between groups in sociodemographic and neuropsychological features were analyzed using SPSS version 13.0 (SPSS, Chicago, IL, USA) performing an analysis of variance (ANOVA) for continuous variables and paired \( t \)-test for dichotomous variables. For continuous variables, post-hoc pairwise comparisons. For continuous variables, post-hoc pairwise comparisons among groups were performed with the Games-Howell or Bonferroni tests depending on homogeneity of variance tested with Levene’s test.

Concerning the neuroimaging analysis, the Qdec interface in Freesurfer software was used: a vertex-by-vertex analysis was carried out performing a general linear model to analyze whether any difference in mean cortical thickness existed between groups (low: \( \alpha3/\alpha2 \leq 1 \mu V^2 \); middle: \( 1 < \alpha3/\alpha2 < 1.16 \mu V^2 \); high: \( \alpha3/\alpha2 \geq 1.17 \mu V^2 \) ). The following comparisons were carried out: high vs. low, high vs. middle and middle vs. low. Age, sex, education, global cognitive level (MMSE score) and WMHs were introduced as covariates in the analysis to avoid confounding factors. We first tried to apply an appropriate Bonferroni multiple-comparison-correction in our analysis (at \( p < 0.05 \) corrected). Unfortunately no \( p \)-value survived after this correction. Thus we choose to set a more restrictive significance threshold (than \( p < 0.05 \) corrected) at \( p < 0.001 \) uncorrected for multiple comparison. Moreover, we considered as significant only the clusters which also were wide equal or major to 30 mm\(^2\). Finally a surface map was generated to display the results on an average brain. For illustrative purpose significance was set to a \( p \)-value of \( p < 0.01 \) uncorrected for multiple comparisons.

As a control analysis, in order to exclude casual relationships between EEG markers and cortical volumes, a correlation between brain areas and memory performance has been studied. The correlation analysis was performed on the three samples separately (high \( \alpha3/\alpha2 \), low \( \alpha3/\alpha2 \), middle \( \alpha3/\alpha2 \)) and on the entire sample (high and low and middle grouped together). An exploratory analysis of non-linear correlation does not fit into purpose of testing out a priori hypotheses. Indeed we choose to apply a measure of linear dependence led by our a priori hypotheses for which the MCI group with the greater cortical thinning, and higher \( \alpha3/\alpha2 \) EEG level (indicating an incipient AD) should show a clear correlation with the memory tests performance. The correlation analysis on a vertex-by-vertex basis was performed specifically for the following neuropsychological memory test results: Babcock test, Rey auditory verbal learning test (AVLT) immediate recall, and Rey AVLT delayed recall. The analysis was thresholded at \( p < 0.001 \) uncorrected for multiple comparisons while results were mapped at \( p \)-value of \( <0.005 \) uncorrected for illustrative purpose. Only the clusters which survived at the statistical threshold and wide equal or major to 15 mm\(^2\) were considered as significant.

**RESULTS**

Table 1 shows the sociodemographic and neuropsychological characteristics of MCI subgroups defined by the tertile values of \( \alpha3/\alpha2 \) ratio. The ANOVA analysis showed that there was not statistically significant differences between groups which resulted well paired for age, sex, WMHs burden, education and global cognitive level. Anyway, age, sex, education, global cognitive level (MMSE score) and WMHs were introduced as covariates in the subsequent analysis to avoid confounding factors. \( \alpha3/\alpha2 \) ratio levels were significant at Games-Howell post hoc comparisons (\( p = 0.000 \)).

**PATTERN OF Cortical THICKNESS BETWEEN GROUPS**

High vs. low: when compared to subjects with low \( \alpha3/\alpha2 \) ratios, patients with high \( \alpha3/\alpha2 \) ratio show thinning in the bilateral Superior/Temporal, Supramarginal and Precuneus cortex, in the right Inferior Parietal and Insula. The total CGM reduction in high \( \alpha3/\alpha2 \) group than low \( \alpha3/\alpha2 \) group was 471 mm\(^2\) (Figure 1; Table 2).

High vs. middle: the same group showed a similar but less wide pattern of cortical thinning when compared to middle \( \alpha3/\alpha2 \) group: the regions of atrophy were located in the left Supra marginal gyrus, left Precuneus and Ponscentral cortex. The total CGM reduction in High \( \alpha3/\alpha2 \) group than middle \( \alpha3/\alpha2 \) group was 160 mm\(^2\) (Figure 2; Table 2). When High group was compared to low group the total extension of cortical thinning (471 mm\(^2\)) was 34% wider than the other comparison in which High group was compared to middle group (160 mm\(^2\)). No regions of major cortical atrophy was found in groups with middle or low \( \alpha3/\alpha2 \) power ratio when compared to High \( \alpha3/\alpha2 \) group. No significant cortical thickness differences was found between middle and low \( \alpha3/\alpha2 \) groups.

**CORRELATIONS BETWEEN NEUROPSYCHOMETRIC MEMORY TESTS AND Cortical THICKNESS IN HIGH \( \alpha3/\alpha2 \) GROUP AND OTHER GROUPS**

Babcock test: a significant positive correlation was found in High \( \alpha3/\alpha2 \) group between logical memory performance at Babcock test and thickness values in the left caudal middle frontal (cluster size = 36 mm\(^2\); stereotaxic coordinate x, y, z = −34, 22, 47; \( r = 0.80, p = 0.0001 \)) and left Inferior Temporal (15 mm\(^2\); −54, −28, −26; \( r = 0.72, p = 0.001 \)), right rostral middle frontal (28 mm\(^2\); 21, 56, −13; \( r = 0.74, p = 0.0007 \)). No significant correlation was found with the same regions nor in the other groups nor in the whole sample (see also Table in Supplementary materials).

AVLT immediate recall: in High \( \alpha3/\alpha2 \) group memory performance were significant related with the cortical thickness values in the bilateral precuneus (left: 47 mm\(^2\); −21, −61, 28; \( r = 0.78, p < 0.00000 \); right: 58 mm\(^2\); 20, −60, 25; \( r = 0.72, p = 0.0007 \)), left fusiform (40 mm\(^2\); −41, −25, −21; \( r = 0.76, p = 0.0005 \)), inferior parietal (43 mm\(^2\); −46, −60, 11; \( r = 0.74, p = 0.0001 \)), inferior temporal (35 mm\(^2\); −53, −34, −21; \( r = 0.71, p = 0.0004 \)).
FIGURE 1 | In red are represented the brain regions with higher regional cortical thickness in MCI with high $a_3/a_2$ ratio as compared to MCI with low $a_3/a_2$ ratio ($p < 0.01$ uncorrected). The color-coding for $p$ values is on a logarithmic scale. Results are presented on the pial cortical surface of brain: dark gray regions represent sulci and light gray regions represent gyri.

Table 2 | Brain regions with significant regional cortical thickness differences in MCI with high $a_3/a_2$ ratio compared to MCI with low $a_3/a_2$ ratio (high $a_3/a_2 < \text{low } a_3/a_2$) and MCI with middle $a_3/a_2$ ratio (high $a_3/a_2 < \text{middle } a_3/a_2$).

| Cluster size (mm$^2$) | Region | Side | Stereotaxic coordinate | $P$ | High Thickness (mm) | Low Thickness (mm) |
|-----------------------|--------|------|------------------------|-----|---------------------|-------------------|
|                       |        |      | $xyz$                  |     |                     |                   |
|                       | 61     | L    | $-47$ $-36$ $9$        | $<0.0001$ | 2.25 ± 0.15          | 2.00 ± 0.18       |
|                       | 60     | L    | $-40$ $-36$ $18$       | 0.0008 | 1.95 ± 0.18          | 2.08 ± 0.19       |
|                       | 35     | L    | $-14$ $-48$ $58$       | 0.0002 | 1.87 ± 0.14          | 1.98 ± 0.19       |
|                       | 58     | L    | $-22$ $-36$ $18$       | $<0.0001$ | 2.18 ± 0.17          | 2.24 ± 0.24       |
|                       | 59     | R    | $-29$ $27$ $<0.0001$   | 1.94 ± 0.18 | 2.08 ± 0.20       |
|                       | 52     | R    | $-49$ $30$ $0.0001$    | 1.81 ± 0.17 | 1.93 ± 0.15       |
|                       | 85     | R    | $-75$ $10$ $0.0001$    | 1.94 ± 0.22 | 2.07 ± 0.23       |
|                       | 61     | R    | $-2$ $3$ $0.0002$      | 2.37 ± 0.28 | 2.57 ± 0.36       |
|                       | 59     | L    | $-57$ $-18$ $18$       | 0.0002 | 1.51 ± 0.15          | 1.62 ± 0.17       |
|                       | 71     | L    | $-53$ $-42$ $46$       | 0.001 | 1.95 ± 0.18          | 2.11 ± 0.23       |
|                       | 33     | L    | $-16$ $-43$ $60$       | 0.001 | 1.87 ± 0.14          | 1.94 ± 0.17       |

Cluster size represents the extension of contiguous significant voxels in the cluster obtained at $p < 0.01$ uncorrected (cluster size $> 30$ mm$^2$). Stereotaxic coordinates reveal the position of the most significant voxel of the cluster, and side denotes its localization on the left (L) or right (R) brain hemisphere. Thickness denotes the average cortical thickness and standard deviation values within the cluster in high and middle $a_3/a_2$ groups. $P$ denotes the significance level of the differences in thickness between groups.

$p = 0.0008$, and right Banks of the superior temporal sulcus (44 mm$^2$, 48; $-48$, $9$; $r = 0.81$; $p < 0.000$; Figure 4A). Memory performance was correlated in the middle group with both the right precuneus also ($r = 0.19$ and $p = 0.05$), and the right Banks of the superior temporal sulcus ($r = 0.44$; $p = 0.02$). No significant associations was find either in the low group nor in the entire sample (Table 3).

AVLT delayed recall: in High alpha3/alpha2 group memory function correlate significantly with cortical thickness in the bilateral Inferior Parietal (left: 95; $-44$, $-58$, $12$; $r = 0.86$; $p < 0.0000$, right: 49; $50$, $-50$, $9$; $r = 0.74$, $p = 0.0003$), left Pericalcarine cortex (54; $-7$, $-8$, $11$; $r = 0.76$; $p < 0.0000$) and Banks of the Superior Temporal Sulcus (31; $-51$, $-41$, $-5$; $r = 0.81$; $p = 0.0002$); in the right Superior Temporal (21; 56, $-34$,$-13$; $r = 0.73$; $p = 0.001$; Figure 4B). No significant correlation was found with the same regions nor in the other groups nor in the whole sample (Table 3).
FIGURE 2 | In red are represented the brain regions with higher regional cortical thickness in MCI with high a3/a2 ratio as compared to MCI with middle a3/a2 ratio (p < 0.01 uncorrected). The color-coding for p values is on a logarithmic scale. Results are presented on the pial cortical surface of brain: dark gray regions represent sulci and light gray regions represent gyri.

DISCUSSION

ASSOCIATION BETWEEN EEG MARKERS AND GM CHANGES

In the present study the relationship between an EEG marker (the alpha3/alpha2 power ratio) and the cortical thickness in subjects with MCI was investigated. The alpha3/alpha2 power ratio has been chosen because in previous works it has been demonstrated that MCI subjects with higher alpha3/alpha2 ratio are at major risk to develop AD (Klimesch et al., 2006, 2007; Bakkour et al., 2009; Moretti et al., 2012a). Our results show that the MCI group with higher alpha3/alpha2 ratio has a greater global cortical atrophy than the other subgroups, thus confirming a large body of literature (Frisoni et al., 2007; Moretti et al., 2012a). Furthermore, the greater atrophy is significant in two specific brain areas: precuneus and supramarginal gyrus (a brain area belonging to the inferior parietal lobule), both in left and right hemisphere. These results were largely expected considering previous studies. Indeed, structural and functional abnormalities of the precuneus were observed in MCI (Matsuda, 2007; Petrella et al., 2007; Dai et al., 2009; Pihlajamaki et al., 2009) as well as in Alzheimer’s disease (Dickerson and Sperling, 2009; Ryu et al., 2010; Sperling et al., 2010) so that the atrophy of precuneus has been considered as a pathognomonic marker of early AD. Recent studies suggest that specific regions, namely the precuneus and posterior cingulate, together with the medial temporal lobe, are selectively vulnerable to early amyloid deposition in AD pathology (Pievani et al., 2011; de Haan et al., 2012).

NEUROPHYSIOLOGICAL AND CLINICAL IMPLICATIONS

Recent studies have demonstrated that during the successful encoding of new items there is a desynchronization in the temporo-parietal memory-related networks whereas a synchronization prevent a successful semantic encoding (Sperling et al., 2010; Chatwal and Sperling, 2012). The deleterious role of synchronization has been recently demonstrated by an interesting study facing the intriguing relationship between functional and structural degeneration in AD (de Haan et al., 2012). The authors detected some hub regions (heteromodal associative regions)
FIGURE 4 | (A) Regions with significant correlation between memory performance evaluated with the AVLT Test immediate recall and cortical thickness measure in the High a3/a2 group. \( r \) and \( p \) denote the correlation coefficient and significance level respectively. (B) Regions with significant correlation between memory performance evaluated with the AVLT Test delay recall and cortical thickness measure in the High a3/a2 group. \( r \) and \( p \) denote the correlation coefficient and significance level respectively.
selectively vulnerable in AD pathology, due to the damage of inhibitory interneurons providing a loss of inhibition at cellular level. According to the authors, the disinhibition provokes an increasing amount of neural activity at network level, giving as a final result an hypersynchronization of brain areas. Of note, this overactivity is excitotoxic and determines cellular apoptosis and brain atrophy. Also, Palop and Macke emphasize the role of inhibitory interneuron dysfunction, leading to hypersynchronization (Stam et al., 2003; Palop and Macke, 2010; Jones et al., 2011; Brier et al., 2012). Our results are in line with these previous influential studies. A possible integrative view of all the results could be as follows: (1) the higher neuronal activity in the hub regions starts from a disfunction of cellular inhibition; (2) the consequent disinhibition drives neural network to an oversynchronization; (3) this oversynchronization is peculiar of the hub regions with higher amyloid burden; (4) these overactivated regions are prone to degeneration and atrophy; (5) a possible neurophysiologic sign of this oversynchronization is the increase of the alpha3/alpha2 power ratio we have found in typical hub regions (Rosum et al., 2008; Wonderlick et al., 2009; Bhattacharya et al., 2011; Wu et al., 2013). It is of great interest that there is an overlapping between the brain regions associated with increase of EEG alpha3/alpha2 power ratio (hypersynchronization of upper alpha) in our study and the regions associated to higher amyloid burden related to memory processes (Jones et al., 2011; Brier et al., 2012). Moreover, in the present study, there is a very interesting result. The atrophy of precuneus is coupled with the atrophy in supramarginal gyrus and, at lesser extent, with inferior parietal, insula and superior temporal gyrus. This atrophy pattern is clearly expressed in the group of MCI subjects with higher amyloid burden related to memory processes (Jones et al., 2011; Brier et al., 2012). The authors found that there is a preferential pathway of

### Table 3 | Correlations between neuropsychological memory tests and cortical thickness in High a3/a2 group and other groups, including whole sample group.

| Regions                     | High (n = 18) | Middle (n = 38) | Low (n = 18) | ALL (n = 74) |
|-----------------------------|--------------|----------------|-------------|-------------|
| Babcock                     | r = 0.88     | r = 0.10       | r = 0.24    | r = 0.18    |
|                            | p = 0.0001*  | p = 0.26       | p = 0.44    | p = 0.26    |
| Right rostral middle frontal| r = 0.74     | r = 0.08       | r = 0.27    | r = 0.08    |
|                            | p = 0.0007*  | p = 0.37       | p = 0.08    | p = 0.37    |
| Left inferior temporal      | r = 0.72     | r = 0.08       | r = 0.27    | r = 0.08    |
|                            | p = 0.0001*  | p = 0.16       | p = 0.21    | p = 0.21    |
| Left precuneus              | r = 0.78     | r = 0.25       | r = 0.07    | r = 0.13    |
|                            | p < 0.0000*  | p = 0.79       | p = 0.01    | p = 0.01    |
| Left fusiform               | r = 0.76     | r = 0.21       | r = 0.08    | r = 0.08    |
|                            | p = 0.0005*  | p = 0.76       | p = 0.02    | p = 0.02    |
| AVLT immediate              | r = 0.71     | r = 0.29       | r = 0.10    | r = 0.10    |
| Left inferior temporal      | r = 0.0008*  | r = 0.12       | r = 0.07    | r = 0.17    |
|                            | p = 0.0001*  | p = 0.14       | p = 0.09    | p = 0.09    |
| Right precuneus             | r = 0.72     | r = 0.19       | r = 0.02    | r = 0.21    |
|                            | p = 0.0007*  | p = 0.38       | p = 0.09    | p = 0.09    |
| Right banks of the superior temporal sulcus | r = 0.81 | r = 0.44       | r = 0.04    | r = 0.31    |
| Left inferior parietal      | r = 0.86     | r = 0.25       | r = 0.05    | r = 0.16    |
|                            | p < 0.0000*  | p = 0.38       | p = 0.01    | p = 0.01    |
| Left pericentral            | r = 0.76     | r = 0.15       | r = 0.69    | r = 0.17    |
|                            | p < 0.0000*  | p = 0.42       | p = 0.80    | p = 0.17    |
| AVLT delayed                | r = 0.81     | r = 0.24       | r = 0.15    | r = 0.12    |
| Left banks of the superior temporal sulcus | r = 0.81 | r = 0.20       | r = 0.58    | r = 0.34    |
| Right inferior parietal     | r = 0.74     | r = 0.19       | r = 0.30    | r = 0.16    |
|                            | p = 0.0005*  | p = 0.31       | p = 0.56    | p = 0.56    |

*fnagi-05-00063* — 2013/10/25 — 13:39 — page 10 — #10
connectivity of the dorsal precuneus with supramarginal gyrus, parietal cortex, superior temporal gyrus and insula. As a consequence, the atrophy we detected in the MCI group with higher alpha3/alpha2 ratio power could be hypothesized as the loss of GM in an entire anatomo-functional network more than atrophy of isolated brain areas. Of note, it is widely accepted that AD is the result of a cortical network impairment more than the atrophy of single cortical areas (Morbelli et al., 2012). In subjects with low or middle alpha3/alpha2 power ratio the cognitive impairment is possibly due to cerebro-vascular impairment or non-AD degenerative process. Although the rigid selection criteria adopted to include in the study patients with primary cognitive deficits, in the clinical practice is not infrequent to have MCI subjects not due to AD.

MEMORY PERFORMANCE AND CORTICAL THINNING
In order to exclude a random relationship between EEG marker and cortical atrophy, the correlation between brain areas and the performance to memory tests was investigated in all MCI subgroups. The memory tests were chosen because of their well known greater impairment in MCI subjects who will convert to AD (Dubois et al., 2007; Frisoni, 2012). Our results show no significant memory difference among the groups. This could be a paradoxical outcome. Anyway, it could not be considered a no surprising result, taking in mind the globally mild and early impairment of the whole group of subjects. In other words, when considering strictly the memory performance, the groups are not different. This is probably due to the early and generally MCI. Anyway, despite no significant difference in the memory test scores, when focusing on the relationship between the memory performance and a reliable structural marker, such as the cortical thickness, the MCI group with the higher alpha3/alpha2 power ratio has shown a (negative) correlation between memory tests performance and the cortical thickness, as expected in patients with probable prodromal AD. This result confirms the peculiar nature of this MCI group, showing a clear specificity as regards both the cortical atrophy and the correlated memory performance. Moreover, no other sociodemographical or structural differences were observed in the MCI groups that could explain the correlation analysis results. The cortical areas associated with cortical thinning and those correlated with memory tests performance are only partly overlapping. This could be due to particular nature of the memory domain, underpinning a large number of brain areas. On the other hand, MCI subjects more susceptible to convert to AD could show impairment also in other cognitive domain like as visuospatial attention or in execution and preparation of spatially guided behavior (Ghazanfar et al., 1997; Leichtnetz, 2001; Wenderoth et al., 2005; Cavanna and Trimble, 2006). Of note, the cortical network encompassing precuneus and inferior parietal cortex is deeply involved in visuo-patial abilities (Zhang and Li, 2012). As a speculative interpretation, we could hypothesize that the memory deficits could be due to impaired network underlying the semantic coding of the spatial features of the episodic memory traces. In this view, the atrophy of a specific brain network (more than global volume measures) is more reliable in detecting MCI subjects with prodromal AD. Anyway, the discussion of memory-related brain networks was beyond the scope of the present study. Only a weak negative correlation was found in the middle alpha3/alpha2 EEG power ratio, suggesting a possible degenerative nature of the memory impairment in this group. No significant associations were found in low alpha3/alpha2 power ratio group and in the whole sample. Taken together, these results strengthen the position of the higher alpha3/alpha2 ratio MCI group as at major risk to develop AD.

IMPLICATIONS AT SYSTEM LEVEL
Eltisesh et al. (1996, 2001) have convincingly demonstrated that the upper alpha band (~10–13 Hz) specifically reflects encoding memory processes. Recent EEG and magnetoencephalography (MEG) studies have confirmed that a correct functioning of memory, both in encoding and in retrieval, requires the high alpha rhythm desynchronization (or power decrease; Shannon and Weaver, 1949; Fries et al., 2001) and with poor information transmission, according to entropy’s theory (Rosen et al., 1980; Hanslmayr et al., 2012). The increases in alpha amplitudes reflect inhibition of cortical brain regions (Fokstein et al., 1975; Hughes et al., 1982, Jensen and Mazaheri, 2010). Similarly, the entropy’s theory stated that synchronization is disadvantageous for storing information, as it reduces the flow of information (Rosen et al., 1980). Entropy is a measure of the richness of information encoded in a sequence of events. Applying this concept to the neural networks, it has been demonstrated (Zhang and Li, 2012) that the degree of information that is encoded in neural assemblies increases as a function of desynchronization and decreases as a function of synchronized firing patterns (Norman, 2010; Schmi dman et al., 2011). This hypothesis has been confirmed in clinical studies in patients with memory deficits (Kurimoto et al., 2012), as well as during states where there is little cognitive processing (e.g., epileptic seizures or slow wave sleep, Goard and Dan, 2009; Chalk et al., 2010; Zhang and Li, 2012). As regards cognitive impairment due to AD, the typical synaptic loss could prevent the physiological flexibility of brain neural assemblies, impeding the desynchronizing downstream modulation of the brain activity. As a consequence, it could be hypothesized that the disruption of cortical network due to degenerative disease, inducing cortical atrophy, could determine an over synchronization of the brain oscillatory activity. The synchronization state of the high alpha power could prevent the creation of a semantic sensory code and, consequently, of the episodic memory trace (Barlow, 1961; Baik et al., 1991; Hanslmayr et al., 2009). In previous seminal studies, high alpha frequency has been specifically related to semantic memory processes (Craik, 2002; Moretti et al., 2004). Of note, in subjects with early cognitive decline, the impairment of the semantic features of memory has been recently accepted as a hallmark for the early AD diagnosis (Dubois et al., 2007; Albert et al., 2011). Indeed, according to the new diagnostic criteria of AD, the measurement of sensitivity to semantic cueing can successfully differentiate patients with AD from healthy controls, even when patients are equated to controls on MMSE scores.
or when disease severity is very mild. Our results are generally in line with this hypothesis, suggesting that increase in power of high alpha brain oscillations reflects a block of information processes. However, the present study goes one step further, linking the increase of high alpha synchronization to the atrophy of a specific brain network, correlated with impairment in memory performance.

**STUDY LIMITATIONS**

There are some limitations due to the obvious explorative nature of the present study: (1) further studies are needed to confirm our result on larger samples and applying an appropriate multiple comparison correction; (2) the pattern of cortical thickness should be investigated on the remaining EEG frequency measures; (3) finally the retrospective nature of the study prevented a direct assessment of whether subjects with increase of α3/α2 EEG power ratio will convert to Alzheimer’s or other neurodegenerative disease; (4) the conservative $p < 0.001$ used here is not necessarily sufficient given the number of comparisons. Anyway, given the explorative nature of the study it is plausible a permissive approach in order to avoid to reject possibly interesting results. It remains clear that further studies with less permissive statistical approach are mandatory to confirm results. Of note, the reliability of the results is supported by: (1) the rigorous selection criteria of the subjects; (2) the high statistical threshold ($p < 0.001$) considered; (3) the large size of pixel accounted for the analysis (30 mm$^2$); and (4) and, finally, the statistical control analysis represented by the correlation of cortical thickness with memory tests.

**CONCLUSION**

The present results show that synchronization (or increase in power) of high alpha is associated with greater cortical atrophy. The greater cortical atrophy is present both considering the whole brain volume and in a peculiar memory-related network, including precuneus and temporo-parietal cortices. The correlation of EEG α3/α2 ratio and cortical thickness measure could be useful for identifying individuals at risk for progression to AD dementia and may be of value in clinical context.

**REFERENCES**

Albert, M. S., DeKosky, S. T., Dickson, D., Dubois, B., Feldman, H. H., Fox, N. C., et al. (2011). “The diagnosis of mild cognitive impairment due to Alzheimer’s disease: recommendations from the National Institute on Aging-Alzheimer’s Association workgroups on diagnostic guidelines for Alzheimer’s disease.” Alzheimers Dement 7, 270–279. doi: 10.1016/j.jalz.2011.03.008

Bakker, A., Morris, J. C., and Dickerson, B. C. (2009). The cortical signature of prodromal AD: regional thinnings predict mild AD dementia. Neurology 72, 1048–1055. doi: 10.1212/WNL.0b013e3181a4d9e1

Bakke, J. H., O’Connell, R. G., Balsters, J. H., and O’Connell, R. G., 2022. EEG power ratio will convert to Alzheimer’s or other neurodegenerative disease; (4) and, finally, the statistical control analysis represented by the correlation of cortical thickness with memory tests.

The present results show that synchronization (or increase in power) of high alpha is associated with greater cortical atrophy. The greater cortical atrophy is present both considering the whole brain volume and in a peculiar memory-related network, including precuneus and temporo-parietal cortices. The correlation of EEG α3/α2 ratio and cortical thickness measure could be useful for identifying individuals at risk for progression to AD dementia and may be of value in clinical context.
Wu, X., Li, B., Flohr, A. S., Reiman, E. M., Guan, X., Zhang, Y., et al. (2011). Altered default mode network connectivity in Alzheimer’s disease – a resting functional MRI and Bayesian network study. Hum. Brain Mapp. 32, 1868–1881. doi: 10.1002/hbm.21153

Wyart, V., and Tallon-Baudry, C. (2008). Neural dissociation between visual awareness and spatial attention. J. Neurosci. 28, 2667–2679. doi: 10.1523/JNEUROSCI.4748-07.2008

Zhang, S., and Li, C. S. (2012). Functional connectivity mapping of the human precuneus by resting state fMRI. Neuroimage 59, 3548–3562. doi: 10.1016/j.neuroimage.2011.11.023

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