Combining Transcranial Magnetic Stimulation and Electroencephalography May Contribute to Assess the Severity of Alzheimer’s Disease

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Received 30 December 2010; Revised 3 March 2011; Accepted 13 March 2011

Academic Editor: Fabio Ferrarelli

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Alzheimer’s disease (AD) is the most common form of old age dementia, and mild cognitive impairment (MCI) often precedes AD. In our previous study (Julkunen et al. 2008), we found that the combination of transcranial magnetic stimulation (TMS) and electroencephalography (EEG) was able to find distinct differences in AD and MCI patients as compared to controls. Here, we reanalyzed the small sample data from our previous study with the aim to test the sensitivity of the TMS-EEG characteristics to discriminate control subjects (n = 4) from MCI (n = 5) and AD (n = 5) subjects. Furthermore, we investigated how the TMS-EEG response characteristics related to the scores of the dementia rating scales used to evaluate the severity of cognitive decline in these subjects. We found that the TMS-EEG response P30 amplitude correlated with cognitive decline and showed good specificity and sensitivity in identifying healthy subjects from those with MCI or AD. Given the small sample size, further studies may be needed to confirm the results.

1. Introduction

Alzheimer’s disease (AD) is a neurodegenerative disorder which leads to dementia through a progressive cognitive decline. In Europe, AD affects over 5% of population aged above 70 years [1]. This makes it the most common cause of dementia in old age. It has been postulated that the impairment of the lateral cholinergic pathway originating from the Meynert’s nucleus would characterize AD and contributes to its typical symptom of memory loss [2, 3]. AD-related pathology leads to the degeneration of the large cortical pyramidal neurons [4], and subsequently impairment of functional connectivity takes place [5]. Before the diagnosis of AD can be set, subjects often suffer from impaired episodic memory [6]. The stage characterised by mild memory or other cognitive loss is called mild cognitive impairment (MCI), and it has been proposed as a prodromal state of AD. Thus, subjects with MCI have an increased risk to develop AD [7–9]. Understanding the pathophysiology of MCI would be essential for predicting and possibly in the future preventing the development of AD. It is possible that altered functional connectivity precedes structural changes, and therefore, a sensitive method to detect those early functional changes would be useful in the diagnostics of MCI and AD. Early identification of AD would be desirable, as it could help aiming the current treatment to the appropriate subjects. With the prospects of obtaining treatments that modify the course of AD, accurate identification of subjects who will develop AD is essential.

Earlier it has been shown that the primary motor cortex experiences changes during the development of AD, which also relate to the severity of the disease [10]. Structural
changes in M1 are mild and appear late as compared to other brain areas, and therefore, motor function also appears intact in early AD [11–14]. Several earlier TMS studies have found that AD patients have reduced resting motor threshold (MT) of the primary motor cortex [3, 15–21]. Alagona et al. reported that the resting MT correlates inversely with the disease severity [15]. This implies that the inhibitory control is reduced in AD, which is also supported by reported shortening of cortical silent period [21]. Additionally, previous studies have reported reduction in short-latency afferent inhibition (SAI) in AD [18, 22–24]. SAI has been considered as a marker of central cholinergic activity [25] and is likely of cortical origin [26, 27]. Hence, motor cortex functions, especially intracortical inhibition, suffer during the development of AD. Earlier, Sakuma et al. [23] showed that SAI is not impaired in MCI, suggesting that the cholinergic activity shown to be impaired in AD may still be normal in MCI. Several studies have been conducted to solve this question and supporting as well as contradicting results have been published [28–31]. Hence, the cholinergic changes related to MCI should be interpreted carefully, as the cholinergic regulation in MCI is still unclear. Furthermore, in AD, there is a tendency towards a reduced short-latency intracortical inhibition (SICI), a different form of inhibition evoked by using paired-pulse TMS [3, 18, 19]. SICI has been connected with intracortical GABA A activity [32].

Combining TMS with electroencephalography (EEG) offers a direct noninvasive method to study cortical reactivity and connectivity in physiological and pathological conditions [33–38]. Previously, we have shown that TMS-EEG can reveal abnormalities in functional cortical connectivity and reactivity in the AD subjects [39]. Our main finding was that the P30 response of TMS-EEG was significantly reduced in AD as compared to controls and MCI, and that the reduction was localized to the ipsilateral temporoparietal area as well as contralateral frontocentral area, that is, sensorimotor area, connected to M1. In the past, TMS-EEG response, when focused on M1, has been shown to exhibit several distinguishable peaks: N15, P30, N40, P60, and N100 [33, 35, 36, 38, 40–43]. Prior studies have related the early peaks N15 and P30 to the M1 activation. P30 has been suggested to reflect activity around the premotor cortex on the stimulated side, and it has been reported that P30 may increase due to long-term potentiation induced by repetitive TMS [41]. Furthermore, P30 has been suggested to involve pathways between subcortical structures such as thalamic nuclei or basal ganglia and cortex [40]. Also, P30 has been shown to vanish with nonoptimal orientation of the stimulation coil in respect to the cortical structures [40]. Therefore, the use of neuronavigation in combination with TMS allows controlling of the stimulation direction in respect to the subject’s brain anatomy and results in optimized motor responses [44], and likely optimized TMS-EEG responses.

We wanted to investigate subject-specific differences in intracortical connectivity between healthy subjects, and MCI or AD patients. We utilized and reanalyzed our previously published data [39], which indicated that especially the P30 amplitude of the TMS-EEG response could be decreased in AD. We further evaluated the sensitivity of the P30 amplitude changes in discriminating healthy subjects from those exhibiting cognitive impairment (MCI and AD). Furthermore, we tested whether P30 amplitude would directly relate to commonly categorizing scores of dementia rating scales. On the basis of the findings of our previous study, we hypothesized that P30 amplitude would decline as the disease becomes more severe and correlate with the dementia rating scales.

2. Materials and Methods

2.1. Subjects. In the present study, our previously published data was further analyzed. A small size sample including four control subjects (age: 78 ± 3 years, 3 females, 1 male), five MCI subjects (age: 74 ± 8 years, 2 females, 3 males), and five AD subjects (age: 73 ± 8 years, 2 females, 3 males) was recruited for the original study. All subjects were right handed. Each subject gave written informed consent, and the study was approved by the local ethics committee. Categorizing of these subjects to their groups was done based on a standard rating [45] and is explained in more detail in the original paper [39]. Briefly, the MCI subjects fulfilled the following characteristics [7]: (1) subjective memory impairment corroborated by an informant, (2) objective memory impairment, that is, a score of 0.5 in the clinical dementia rating (CDR) scale [45] with at least 0.5 on the memory subscale and a score of 1.5 SD below the average of a normative age-matched sample group in at least one memory test, (3) normal global cognitive function (Mini-Mental-State Examination score (MMSE) of at least 20 [46]), (4) normal activities of daily living, and (5) no dementia according to the NINCDS-ADRDA criteria [11]. The MCI subjects were classified as multidomain amnestic MCI [47]. Diagnosis of AD was made according to the NINCDS-ADRDA criteria for probable AD [11]. All the AD patients were on cholinesterase inhibitors, while other subjects had no medication affecting cognition at the time of measurements.

2.2. Measurement System and Protocol. Navigated TMS was used to probe the motor cortex of the subjects (Figure 1), that is, the primary motor cortex (M1) of the subjects was mapped for the representation area of the thenar musculature of both hands. The stimulation system consisted of a Magstim BiStim stimulator (Magstim Ltd., Whitland, UK) and a 70 mm figure-of-eight TMS coil with monophasic pulse form. Stimulation-triggered EEG responses were recorded with 1450 Hz sampling frequency and 16 bit precision using a 60-channel TMS-compatible EEG amplifier (Nexstim Ltd., Helsinki, Finland). Navigation of the TMS system utilized T1-weighted 3D magnetic resonance images (imaged with Siemens Magnetom Avanto, Siemens, Erlangen, Germany). The navigation was conducted using eXimia navigation system (version 2.0, Nexstim Ltd., Helsinki, Finland). Resting state MT at the “hotspot” was determined using a threshold-hunting protocol [48]. For measuring TMS-induced muscle responses, electromyography was recorded (ME6000, Mega Electronics Inc., Kuopio, Finland)
from the opponens pollicis muscle using pregelled disposable Ag-AgCl surface electrodes. TMS-induced EEG responses were recorded from >50 trials elicited with an interstimulus interval of 3–5 s with a stimulation intensity of 110% of the determined MT. For a more thorough system description, the reader is referred to our previous paper [39]. Both hemispheres were separately investigated, and the stimulation order of the hemispheres was randomized.

2.3. Analysis of TMS-EEG. The offline analysis of EEG was performed using Matlab 7.2 (Mathworks Inc., Natick, MA). Zero padding for 10 ms after the TMS pulse was applied to dampen the TMS-induced artefact. Segmented EEG was bandpass filtered to 1–50 Hz. Any segments contaminated by blinks, as observed from vertical electro-oculogram, were removed from the analyses. Also, in some cases, a bad channel signal due to poor contact was replaced with a signal linearly interpolated from the neighbouring good channels. Manual artefact removal was conducted, prior to rereferencing the channel signal due to poor contact was replaced with a signal linearly interpolated from the neighbouring good channels. The o

determination based on the most distinguishable and shortest-latency segment of the TMS-induced artefact. Segmented EEG was performed using Matlab 7.2 (Mathworks Inc., Natick, MA).

2.4. Statistical Analyses. To test how well the P30 amplitude would be able to discriminate the groups from each other, receiver operating characteristic (ROC) curve analysis was conducted. Area under the ROC curve (AUC) was computed to determine how well the groups could be discriminated based on the P30 amplitude. The asymptotic significance for the AUC was computed with the null hypothesis of AUC = 0.5. The optimal cut-off point for the ROC curve was determined as the closest point to the diagonal line connecting points (0, 1) and (1, 0) in the ROC plot.

Differences in P30 amplitude between the groups were analysed applying a mixed linear model, and using group and hemisphere as fixed variables and subject as a random variable. Restricted maximum likelihood estimation was used in the model. Mean effects between the groups were analysed using post-hoc analysis with least significant difference adjustment (LSD). Also, individual mean amplitudes for the P30 (hemispheric values averaged, P30mean) were used in the comparisons. Then, Mann-Whitney test was applied in comparison of the differences between the groups. Correlations between the scores of dementia rating scales and P30 amplitude were conducted using Spearman’s rank correlation (ρ). The tests for correlation significance were two-tailed. The correlated dementia scores were the global score and sum of boxes score of the clinical dementia rating scale (CDR-SOB) as well as MMSE. Statistical tests were conducted using SPSS 17 (SPSS Inc., Chicago, IL). Level of statistical significance was set at P < .05.

3. Results

The resting MTs (average of both hemispheres) for the opponens pollicis muscle of the control, MCI, and AD group were 44 ± 11, 48 ± 12, and 41 ± 4% of the maximum stimulator output, respectively. No significant differences were observed between the groups. The hemispheric data and data for pooled samples were presented in our previous study [39], where we found that the MT of the left hemisphere in MCI subjects (50 ± 13% of the maximum stimulator output) was significantly higher (P < .05) than in AD patients (42 ± 4% of the maximum stimulator output). Additionally, on the right hemisphere, the MT of the controls (40 ± 11% of the maximum stimulator output) was significantly lower (P < .05) than in MCI subjects (48 ± 13% of the maximum stimulator output).

Dementia scales for the subject groups were distinctive of the different disease conditions. MMSE for control, MCI, and AD group was 27 ± 4, 25 ± 3, and 22 ± 5, respectively. The global CDR values were 0 for controls, 0.5 for MCI subjects, and 0.5 (n = 4) or 1.0 (n = 1) for AD patients. Corresponding values for the CDR-SOB were 0.0 ± 0.0, 1.9 ± 1.1, and 3.2 ± 2.5, respectively. CDR and CDR-SOB values in controls were significantly lower (P < .001) than in MCI and AD subjects, while MMSE was nonsignificantly higher (P = .055) in controls as compared to AD group. Also, no significant difference was observed between controls and MCI subjects in MMSE (P = .437).

The P30 peak was lower in amplitude in the AD patients than in the controls (Figure 3, Table 1). No significant
Table 1: Group-wise values of P30 amplitude.

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>MCI</th>
<th>AD</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>P30 amplitude (μV)</td>
<td>P30 amplitude (μV)</td>
<td>P30 amplitude (μV)</td>
</tr>
<tr>
<td>Left hemisphere</td>
<td>32.0 ± 6.0</td>
<td>25.6 ± 12.7</td>
<td>17.7 ± 7.1</td>
</tr>
<tr>
<td>Right hemisphere</td>
<td>33.0 ± 14.6</td>
<td>16.3 ± 5.9</td>
<td>11.5 ± 4.9</td>
</tr>
<tr>
<td>P30\text{mean}</td>
<td>32.5 ± 9.8</td>
<td>21.1 ± 8.2</td>
<td>16.0 ± 6.9*</td>
</tr>
</tbody>
</table>

*P < .05 as compared to controls, linear mixed model (pooled values), and Mann-Whitney test (mean or hemispheric values).

Abbreviations:
MCI: Mild cognitive impairment
AD: Alzheimer’s disease
P30\text{mean}: P30 amplitude, mean of P30 amplitudes on both hemispheres.

Figure 2: Grand average curves for TMS-evoked EEG responses as measured from the central electrode (CZ). The mean peak for the P30 has been indicated. However, P30 was analyzed for individuals from the electrode chosen based on the shortest latency and clearest identification on the stimulated hemisphere. The turquoise area represents the 95% confidence interval for the TMS-EEG responses. The vertical black line indicates the moment of stimulation.

Figure 3: Group-wise P30 amplitudes. The individual values are presented as a mean value of P30 amplitude measured from both hemispheres. Black line represents the group-wise mean value when moving from controls to MCI and AD.

If only the discrimination of AD patients from controls was estimated, the AUC increased to 0.950 (P = .027, P30\text{mean}). The optimal cut-off point was 25.4 μV (sensitivity of 0.80 and specificity of 0.75). Discrimination of MCI from AD using ROC curve was found more difficult (AUC = 0.720, P = .251, P30\text{mean}). The optimal cut-off point then was 18.7 μV (sensitivity of 0.60 and specificity of 0.60). However, the more important discrimination of MCI subjects from controls was found stronger although the AUC was 0.850 (P = .086, P30\text{mean}). The optimal cut-off point was 25.5 μV (sensitivity of 0.75 and specificity of 0.80), which was very close to similar as in discriminating AD patients from controls.

Significant correlations were observed between the P30 amplitude and the dementia scales. An inverse correlation was found between the global CDR and P30 amplitude as well as between CDR-SOB and P30 amplitude (Figure 5). As the global CDR is a classification variable, its correlations with P30 amplitude should be interpreted with care.
**Figure 4**: Receiver operating characteristic (ROC) curves for distinguishing (a) controls from MCI and AD, and (b) AD patients from MCI and control subjects based on TMS-EEG P30. Turquoise line indicates the ROC curve for averaged data, while the grey and black lines indicate ROC curves for the right and left hemisphere, respectively. The area under the ROC curve (AUC) has been given separately for the averaged (P30mean) and hemispheric data. The asymptotic significance has been indicated with the null hypothesis of AUC = 0.5 (diagonal line).

**Table 2**: Correlation coefficients (Spearman’s ρ) between the P30 amplitude and dementia rating scales.

<table>
<thead>
<tr>
<th></th>
<th>Mini-mental state examination</th>
<th>Clinical dementia rating—global†</th>
<th>Clinical dementia rating—sum of boxes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Left hemisphere</td>
<td>0.456</td>
<td>-0.678**</td>
<td>-0.788***</td>
</tr>
<tr>
<td>Right hemisphere</td>
<td>0.631*</td>
<td>-0.705**</td>
<td>-0.849***</td>
</tr>
<tr>
<td>P30mean</td>
<td>0.537*</td>
<td>-0.698**</td>
<td>-0.808***</td>
</tr>
</tbody>
</table>

*P < .05, **P < .01, ***P < .001.
† As the global CDR is a classification variable, its correlations with P30 amplitude should be interpreted with care.

**4. Discussion**

We have previously shown that TMS-evoked P30 amplitude is reduced in the AD subjects in the temporoparietal area, ipsilateral to the stimulation side as well as in the contralateral frontocentral cortex corresponding to the sensorimotor area [39]. In the present study, we further investigated our previously published data and found that the discrimination of control subjects from MCI and AD subjects may be possible with good sensitivity (Figures 3 and 4). Further, consistently with our hypothesis, we found that there is a significant relation between the commonly used dementia rating scales and the analyzed TMS-EEG response P30 amplitude, when TMS is focused on the M1 with suprathreshold intensity (Figure 5). Our results suggest that the use of TMS-EEG in the evaluation of AD and its initial signs could be feasible as distinct changes occur in the measured responses during the development of AD.

The greatest limitation of this study was that the group sizes were small. In spite of that, the findings of the present study showed clearly significant differences between the groups. In the future, these results should be further verified by other studies with larger group sizes. Nevertheless, it was clear that the discrimination of probable mild AD patients and MCI subjects from control subjects seemed feasible. Furthermore, the present study was able to show a correlation between the P30 amplitude of the TMS-EEG and the dementia rating scales (Figure 5, Table 2). Such relation has not been reported earlier. Therefore, it seems that the P30 peak is indeed related to cognitive decline, or perhaps to the developing motor deficits that the AD patients may exhibit in the advanced stage of the disease [11]. Due to the small sample size, the study may suffer a lack of power to provide reliable answers to its aim, that is, some of the intergroup relations and differences may have been missed. Even with
the small sample size, the found effect size for discriminating controls from mild AD patients based on the P30<sub>mean</sub> was large (Cohen's $d > 0.8$), as was the effect size for the controls and MCI difference (Cohen's $d > 0.8$). A medium size effect was observed between the MCI and AD (Cohen's $d > 0.5$). However, the statistical significance of those comparisons was too weak in the last two cases. This encourages further studies with larger sample sizes.

As opposed to the localized (single-channel) P30 responses reported in the present study, our previous study investigated the global mean field power (GMFP) of the P30 peak [39]. Hence, for comparison, the global mean field power (GMFP) was computed for the P30 response peaks (P30<sub>GMFP</sub>) [49]. We found that the P30<sub>GMFP</sub> correlated strongly with the single-channel P30<sub>mean</sub> ($\rho = 0.810, P < .001$). This correlation is affected by the differences in the P30 spread between the groups, and hence the correlation may not be an ideal indicator for similar behavior. In the discrimination of the different groups, P30<sub>GMFP</sub> was weaker than the single-channel P30<sub>mean</sub>, that is, in the ROC analysis, controls were not discriminated as easily from the MCI and AD (AUC = 0.775, $P = .120$) or the AD group from MCI and control groups (AUC = 0.756, $P = .125$). Furthermore, the P30<sub>GMFP</sub> exhibited some correlations with the dementia scales (P30<sub>GMFP</sub> versus MMSE, $\rho = 0.311, P = .279$; P30<sub>GMFP</sub> versus CDR, $\rho = -0.515, P = .060$; P30<sub>GMFP</sub> versus CDR-SOB, $\rho = -0.755, P = .002$). Therefore, to us it seems that the localized P30 amplitude is more sensitive in observing cognitive decline as compared to global field values. This finding may be influenced by the modified spread of the P30 component in AD, which was observed in our previous study, and which affects the GMFP [39].

The CDR is a standard assessment tool that yields global and CDR-SOB scores. The global CDR score is often used to stage dementia severity [45]. However, the CDR-SOB score is a more detailed general index and is more sensitive in assessing mild dementia [50–52]. Based on the CDR-SOB, the AD patients in this study had very mild or mild dementia [52]. In agreement with our hypothesis, we found significant correlations between the P30 amplitude and the dementia scales (Table 2). As the CDR-SOB is a detailed and one of the most used dementia scales, it also related best to the P30 amplitude as was seen from their strong correlation (Figure 5). Also, the other applied dementia scales correlated with P30. Therefore, it seems that P30 indeed relates to the severity of the AD even in a mild stage of the disease as the patients were in the present study. Since our recent study showed some effect of cognitive decline on the N100 [39], we analyzed the N100 response from the TMS-EEG at the vertex for comparison with the P30 amplitude. We found that none of the dementia scales correlated significantly with the N100 amplitude (N100 versus MMSE, $\rho = -0.029$; N100 versus CDR, $\rho = 0.333$; N100 versus CDR-SOB, $\rho = 0.177$). Therefore, P30 appears more specific than N100 in identifying cognitive decline with TMS-EEG. The reason for this may be that TMS-induced N100 response also includes an auditory component, which does not affect P30 [42].

Currently, the origin of P30 is not precisely described. It has been suggested to originate from the ipsilateral sensorimotor/premotor cortex border [41] or the ipsilateral supplementary motor area [53]. As discussed by Mäki and Ilmoniemi, the P30 may not reflect activation directly at the location activated by the stimulus. However, it may still reflect the degree of excitation in M1 [33]. Bonato et al. showed that P30 vanishes if the activation of M1 is induced with TMS coil oriented nonoptimally, which supports the idea that P30 is descriptive of cortical activation related to M1 excitation [40]. Considering the findings of the present study, the decrease in P30 in AD as compared to healthy controls may reflect impaired cortical activation in response to M1 activation. However, if we consider the earlier findings relating MEPs with P30, the P30 measured in the present study would not directly relate to motor activation, as the earlier reports have indicated a positive correlation between the two [33, 40]. In our earlier study [39], we showed that the correlation between the induced MEPs and P30 exhibits a nonsignificant negative trend when correlated over different patient groups ($\rho = -0.224, P = .441$). When comparing the mean values of MEP amplitudes, the controls exhibit the lowest mean MEP amplitude of 1.0 mV while showing the highest P30 amplitude. Instead, the highest mean MEP amplitude of 2.9 mV was in the AD group with the lowest P30 amplitude (Figure 3). This suggests that P30 is not only related to the excitation of M1, but also other mechanisms may influence it when AD progresses. Ferreri et al. [38] suggested that P30 may be connected to GABA<sub>A</sub> receptors, provided that P30 modulation is related to fast inhibitory

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**Figure 5**: Scatter plot indicating the relation between the clinical dementia rating sum of boxes and P30 amplitude (average of each subject's left and right hemisphere measurement, P30<sub>mean</sub>). The thin curved lines represent the 95% confidence intervals for the curve fit.
5. Conclusions

We found differences in TMS-induced P30-component amplitude between the controls and AD patients, indicating impaired/altered connectivity in AD. In addition, we found that the cognitive decline correlated with the P30 amplitude. Further investigations with larger sample sizes are needed to support our conclusion that TMS-EEG could be a potential noninvasive biomarker for identifying MCI and AD subjects and separating those from healthy population, and for identifying connectivity changes occurring during the development of AD.

Conflict of Interests

Professor J. Karhu works part time as Chief Medical Officer in Nexstim Ltd., the manufacturer of navigated brain stimulation instruments.

Acknowledgment

This study was supported by EVO (5772739) Grant from Kuopio University Hospital.

References


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