Differential Pharmacological Effects on Brain Reactivity and Plasticity in Alzheimer’s Disease

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Differential pharmacological effects on brain reactivity and plasticity in Alzheimer’s disease

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INTRODUCTION

Alzheimer’s Disease (AD) is the most common form of dementia and its treatment one of the most pressing medical issues. Current therapeutic options are limited and underlying neurophysiological mechanisms are still insufficiently understood. Available drugs mainly target cholinergic and glutamatergic pathways.

The degeneration of cholinergic neurons in the basal forebrain is thought to extensively contribute to the cognitive decline characteristic of AD (1). This led to the development of acetylcholinesterase inhibitors (AChEIs), which slow down the breakdown of acetylcholine (ACh). The resulting increase of ACh in affected forebrain regions is thought to improve cognitive functions (2, 3). AChEIs currently in use for patients with mild to moderate AD include donepezil (Aricept®), rivastigmine (Exelon®), and galantamine (Razadyne®), all of which show similar efficacy and side-effect profiles (4, 5).

The N-methyl-D-aspartate (NMDA) receptor antagonist memantine (Namenda™) is traditionally used in moderate to severe AD. At first sight it seems contradictory that blocking NMDA receptors, which play a crucial role in synaptic plasticity and therefore learning and memory, should prove efficacious in AD. Yet, bearing in mind the importance of physiological balance, both hyper- as well as hypoactivation of NMDA receptors leads to dysfunctional effects (6). As NMDA receptors are hyperactive in AD, memantine is used to re-balance glutamatergic pathways and therefore reduce neurotoxic processes.

Clinical effects of both AChEi and memantine have been extensively studied and include temporary and moderate improvements in activities of daily living (ADLs), cognitive, and behavioral functions (4, 7–9), and lead to a reduction of pathological encephalographic rhythms (10).

Several pharmacological studies have shown inconsistent results regarding the benefits of combining cholinesterase inhibitors with memantine. Tariot et al. (11) found beneficial effects of combination therapy, while consecutive studies showed no additional benefit when memantine was added to a regime of cholinesterase inhibitors (2, 12).

Transcranial magnetic stimulation (TMS) is a valuable tool to investigate the neurophysiological effects of drugs. Previous studies have examined various TMS-measures in medicated healthy subjects and in AD patients (13). In healthy subjects, memantine reduces intracortical facilitation (ICF) and enhances short-interval intracortical inhibition (SICI) in the motor cortex (MC) (14), and results in reduced MC plasticity when administered over 8 days.

Acetylcholinesterase inhibitors (AChEIs) are the most commonly prescribed monotherapy medications for Alzheimer’s disease (AD). However, their underlying neurophysiological effects remain largely unknown. We investigated the effects of monotherapy (AChEI) and combination therapy (AChEI and memantine) on brain reactivity and plasticity. Patients treated with monotherapy (AChEI) (N = 7) were compared to patients receiving combination therapy (COM) (N = 9) and a group of age-matched, healthy controls (HCs) (N = 13).

Cortical reactivity and plasticity of the motor cortex were examined using transcranial magnetic stimulation. Cognitive functions were assessed with the cognitive subscale of the Alzheimer Disease Assessment Scale-Cognitive Subscale (ADAS-Cog), activities of daily living (ADLs) with the ADCS-ADL. In addition we assessed the degree of brain atrophy by measuring brain-scalp distances in seven different brain areas. Patient groups differed in resting motor threshold and brain atrophy, with COM showing a lower motor threshold but less atrophy than AChEI. COM showed similar plasticity effects as the HC group, while plasticity was reduced in AChEI. Long-interval intracortical inhibition (LICI) was impaired in both patient groups when compared to HC. ADAS-Cog scores were positively correlated with LICI measures and with brain atrophy, specifically in the left inferior parietal cortex. AD patients treated with mono- or combination-therapy show distinct neurophysiological patterns. Further studies should investigate whether these measures might serve as biomarkers of treatment response and whether they could guide other therapeutic interventions.

Keywords: Alzheimer’s disease, transcranial magnetic stimulation, brain plasticity, pharmacological therapy, acetylcholinesterase inhibitors, memantine

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This effect is likewise observed with the NMDA antagonist dextromethorphan (16). Memantine furthermore abolishes inhibitory effects of continuous theta burst stimulation (cTBS) and facilitatory effects of intermittent theta burst stimulation (iTBS) on brain plasticity (17). AD patients treated with memantine have not yet been investigated with TMS. AChEI given to healthy subjects lead to reduced SICI and increased ICF and have no effect on motor threshold and excitability (18). Contradictory results have been found for the effect of AChEI on MC excitability in AD patients. While one study found improvements in SICI after administration of donepezil (19), another study did not find any reversal in the progressive increase of excitability during 1 year of treatment (20).

Rivastigmine furthermore leads to improvements of short latency afferent inhibition (SAI), which is associated to long-term treatment response (21). SAI is the most prominent TMS-measure impaired in AD (13, 21–23). It is mediated through GABA-ergic and cholinergic circuits (24), the latter of which plays a dominant role in AD pathology. SAI is assessed by coupling TMS and peripheral nerve stimulation of the contralateral wrist and has a role in AD pathology. SAI is assessed by coupling TMS and peripheral nerve stimulation of the contralateral wrist and has inhibitory interactions with long-interval intracortical inhibition (LICI) (25). To our knowledge, however, no study has investigated LICI in AD patients.

In order to understand the possible mechanisms of action of different pharmacological regimes, we evaluated MC excitability and plasticity, ADLs, cognitive functions, and brain atrophy in two groups of patients receiving either AChEI monotherapy or AChEI combined with memantine. Findings were contrasted against an age-matched healthy control (HC) group on no medications.

**MATERIALS AND METHODS**

**SUBJECTS AND INCLUSION/EXCLUSION CRITERIA**

We investigated 16 patients diagnosed with mild to moderate AD (DSM IV, NINCDS-ADRDA): 7 were stable on AChEI (Table 1; Figure 1), and 9 were treated with combination therapy (AChEI and memantine). Patients were eligible if they had a score of 1 on the Clinical Dementia Rating scale (CDR), and attained between 18 and 24 points on the Mini-Mental State Examination (MMSE). Patients were recruited from other studies investigating AD patients (ClinicalTrials.gov NCT01504958) in comparison to older HCs. Patients were grouped according to psychopharmacological treatment. All experimenters were blinded with regards to pharmacological treatment.

Alzheimer’s disease groups were compared to a group of 13 age-matched HCs with normal neurological and cognitive exams and a score of 0 on the CDR. The local Institutional Review Board approved the study protocol, and all participants or their legally authorized representatives gave written informed consent prior to study onset. The study visits took place at the Berenson-Allen Center for Non-invasive Brain Stimulation and the Harvard-Thorndike Clinical Research Center at Beth Israel Deaconess Medical Center in Boston, MA, USA. Patients with unstable or
chronic medical conditions, major structural or vascular abnormalities on MRI, severe agitation, functional psychiatric disorders, a history of substance abuse or recent withdrawal, epilepsy, other progressive neurological disorders, other diseases causing memory impairment, or intake of drugs listed as a potential hazard for the application of rTMS \(^{(26)}\) were excluded from the study.

**ASSSESSMENTS**

Brain excitability and plasticity, brain-scalp distances, the MMSE, (ADCS-ADL), and the Geriatric Depression Scale (GDS) were assessed by trained research personnel. Certified neuropsychologists and neurologists assessed Alzheimer disease assessment scale-cognitive subscale (ADAS-Cog) and CDR.

**MAPPING, BRAIN REACTIVITY, AND PLASTICITY MEASURES**

The Nexstim system (eXimia NBS 4) was used for neuronavigated single- and paired-pulse stimulation over the left MC. Resting motor threshold (rMT) was defined as the minimum stimulation intensity required to elicit a motor evoked potential (MEP) in the first dorsal interosseous muscle (FDI) of at least 50 µV in 5 out of 10 trials. Active motor threshold (aMT) was defined as the minimal stimulation intensity required to elicit an MEP in 5 out of 10 trials during isometric contraction of the FDI muscle. MEPs were recorded using 30 mm × 22 mm, wet gel surface electrodes (Ambu). The active electrode was placed over the muscle belly and the reference electrode over the proximal interphalangeal joint of the index finger.

Stimulation intensity for single-pulse TMS was set to 120% of rMT. For paired-pulse TMS, intensity was set to 80% rMT for the conditioning pulse and 120% rMT for the test pulse, with an inter-pulse interval of 3 ms to determine SICI and 12 ms to determine ICF. Two pulses at 120% rMT with an inter-pulse interval of 100 ms were used to determine LICI. Paired-pulse measures (each set 50 paired-pulses) were expressed as the ratio of the mean conditioned MEP amplitude to the mean unconditioned MEP amplitude.

To assess the mechanisms of cortical plasticity, neuronavigated intermittent theta burst stimulation (iTBS) was applied using the MagPro X100 (MagVenture). Baseline single-pulse stimulations (90 pulses at 120% rMT with biphasic coil) were followed by iTBS (600 pulses in 2 s trains at 50 Hz repeated every 10 s; stimulation intensity set to 80% of aMT). Sets of 30 single-pulses were delivered at 120% rMT at distinct time periods post iTBS (after 5, 10, 20, 30, 60, and 90 min).

Brain reactivity was defined as the average amplitude of 90 single-pulse MEPs at baseline; brain plasticity was expressed as the ratio of averaged MEP amplitudes obtained at each of the time points after iTBS to the mean baseline MEP amplitude. Safety guidelines were strictly followed \(^{(26)}\).

**BRAIN-SCALP DISTANCE MEASUREMENTS**

As an index of brain atrophy, brain-scalp distances were measured on each individual’s brain MRI (Brainsight) in seven brain regions: left hand MC, right and left dorsolateral prefrontal cortex (DLPFC), right and left inferior parietal cortex (IPL), left superior temporal gyrus (STG), and left inferior frontal gyrus (IFG). Brain regions were determined using conventional Talairach/MNI coordinates \(^{(27-29)}\). For each brain region, three measurements were taken in both the coronal and sagittal view, and the six measurements were averaged for each region \(^{(28)}\).

**DATA ANALYSIS**

Data was analyzed using non-parametric tests (SPSS 19.0 for Mac) with the statistical significance set at \(p < 0.05\). Group comparisons were calculated with the Kruskal–Wallis test, Mann–Whitney \(U\) tests served as post hoc tests. Correlations were calculated using Kendall’s tau. \(p\)-Values are two-tailed. As this study was an exploratory trial, a formal sample size analysis was not possible.

**RESULTS**

**GROUP EFFECTS**

Patient groups did not differ significantly in age, gender, education years, duration of disease or pharmacologic treatment, mean brain-scalp distance, MMSE, ADCS-ADL, GDS, or ADAS-Cog scores (Table 1).

Acetylcholinesterase inhibitor, COM, and HC did not differ significantly regarding age, gender, education years, or...
GDS. However, the three groups differed significantly in rMT \(H(2) = 8.857, p = 0.012\), brain plasticity at T5 \(H(2) = 6.782, p = 0.034\), and in average degree of TBS-induced plasticity over all time points \(H(2) = 6.146, p = 0.046\). Furthermore, a significant group difference was found for LICI \(H(2) = 8.003, p = 0.018\) (Figure 2). Brain-scalp distances were significantly different between groups when averaged over all regions \(H(2) = 8.551, p = 0.014\), but particularly for left IPL \(H(2) = 7.582, p = 0.023\), and left MC \(H(2) = 9.852, p = 0.007\). 

In the cognitive and behavioral assessments, groups differed significantly in ADAS-Cog scores \(H(2) = 21.713, \ p < 0.001\), MMSE \(H(2) = 22.675, p < 0.001\), and ADCS-ADL \(H(2) = 6.316, p = 0.043\) (Figure 3).

**POST HOC GROUP COMPARISONS**

**Motor thresholds, brain reactivity and plasticity, brain atrophy**

Post hoc tests revealed a significantly lower rMT in COM as compared to AChEI \((U = 60.00, z = 3.02, p = 0.001)\), while rMTs of both patient groups and HC were comparable (Figure 2A). HC showed greater plasticity at T5 \((U = 73.00, z = 2.18, p = 0.030)\), T20 \((U = 74.00, z = 2.26, p = 0.024)\), T60 \((U = 71.00, z = 2.02, p = 0.046)\), and at an average of all time points \((U = 78.00, z = 2.58, p = 0.008)\) when compared with AChEI. COM and HC did not differ significantly in TBS-induced brain plasticity (Figure 2B). LICI was significantly reduced in both COM \((U = 20.00, z = -2.24, p = 0.025)\) and AChEI \((U = 12.00, z = -2.40, p = 0.015)\) as compared to HC (Figure 2C).

![Figure 2](image-url) **FIGURE 2 |** TMS-driven measures of corticomotor reactivity and plasticity in Alzheimer’s patients treated with acetylcholinesterase inhibitors (AChEI) or a combination of AChEI and memantine (COM), and a healthy age-matched control group (HC). (A) Resting motor threshold measured from the first dorsal interosseus muscle (mean ± SE). (B) Brain plasticity measures (mean ± SE) over a time-period of 90 min post iTBS. (C) Paired-pulse TMS measures of long-interval intracortical inhibition (LICI).

Acetylcholinesterase inhibitor showed greater atrophy as compared to HC in all three measures: averaged atrophy \((U = 9.00, z = -2.89, p = 0.002)\), left IPL \((U = 15.00, z = -2.42, p = 0.014)\), and MC \((U = 7.00, z = -3.05, p = 0.001)\). However, AChEI also showed a significantly greater atrophy in MC as compared to COM \((U = 51.00, z = 2.06, p = 0.042)\). COM did not differ from HC in atrophy (Figure 3A).

**ADAS-Cog, MMSE, ADL, GDS**

Alzheimer disease assessment scale-cognitive subscale and MMSE scores were significantly different from HC in both AChEI (ADAS-Cog: \(U = 0.00, z = -3.61, p < 0.001\); MMSE: \(U = 91.00, z = 3.75, p < 0.001\)) and COM (ADAS-Cog: \(U = 0.00, z = -3.91, p < 0.001\); MMSE: \(U = 117.00, z = 4.02, p < 0.001\)) (Figure 3B). COM was significantly more impaired than HC \((U = 78.00, z = 2.60, p = 0.008)\) in ADCS-ADL, while HC and AChEI did not show significant differences (Figure 3C).

**CORRELATIONS (AD AND HC)**

Alzheimer disease assessment scale-cognitive subscale scores were significantly correlated with LICI \((r = 0.352, p = 0.010)\) (Figure 4A) and with mean atrophy over seven brain regions \((r = 0.304, p = 0.021)\), specifically with atrophy of the left IPL \((r = 0.277, p = 0.036)\). Brain-scalp distance in MC was correlated with rMT \((r = 0.287, p = 0.029)\) and aMT \((r = 0.319, p = 0.015)\) and mean brain-scalp distance was correlated with rMT \((r = 0.297, p = 0.024)\). RMT was correlated with aMT \((r = 0.688, p = 0.001)\) and mean brain-scalp distance was correlated with rMT \((r = 0.297, p = 0.024)\).
AChEI patients. COM showed similar plasticity effects as the HC group, while plasticity was reduced in AChEI. This is in line with a previous study (31) showing impaired brain plasticity in AD patients. We found no indication that the reduction in plasticity could be related to the greater atrophy in the left MC measured in AChEI. Future studies need to carefully control for atrophy and adjust stimulation intensity accordingly. However, our results suggest that atrophy does not account for the observed changes in plasticity or the implied medication effects. Disease duration, as expected, was correlated with degree of atrophy. Nonetheless, unexpectedly, we found that plasticity measures positively correlated with duration of disease and time since medication intake. Our data do not allow us to disentangle whether this unexpected finding is related to the duration of the disease or the medication treatment itself. The possibility that medication treatment might reverse an age- and disease-related trend for progressive loss of efficacy of plasticity mechanisms is intriguing. Longitudinal studies would be important to address this question.

DISCUSSION

We investigated the impact of pharmacological monotherapy (AChEI) and combined interventions (AChEI and memantine) on neurophysiological indices of brain reactivity and plasticity in patients with AD and age-matched HCs. Neurophysiological findings were related to cognitive function, disease duration, and measures of brain atrophy.

Patient groups showed differences in rMT and brain atrophy in MC, with COM showing lower motor thresholds but less atrophy than AChEI. Since the current induced by TMS decreases steeply with distance, a greater atrophy would be expected to result in higher MTs. Yet, most studies to date report lower MTs indicating higher motor cortical excitability in AD patients (13), but also opposite effects have been found (30). Differences in MT between COM and AChEI may therefore simply reflect differences in the degree of brain atrophy. Nonetheless, this is somewhat surprising, since greater atrophy would be expected to indicate more progressed disease stage, which might lead to the combination therapy, rather than be found in patients on monotherapy. Disease duration in COM and AChEI was however not significantly different. The addition of memantine to a regime of AChEI may therefore possibly slow down progression of atrophy; a research question that warrants further studies.

Measures of plasticity showed differences between COM and AChEI patients. COM showed similar plasticity effects as the HC group, while plasticity was reduced in AChEI. This is in line with a previous study (31) showing impaired brain plasticity in AD patients. We found no indication that the reduction in plasticity could be related to the greater atrophy in the left
REFERENCES


AUTHOR CONTRIBUTIONS

Anna-Katharine Brem, Natasha J. Atkinson, and Erica E. Seligson prepared the draft report together with Alvaro Pascual-Leone, which was critically reviewed by all authors. Anna-Katharine Brem, Natasha J. Atkinson, and Erica E. Seligson analyzed the data and contributed to data collection.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest. Alvaro Pascual-Leone serves on the scientific advisory boards for Nexstim, Neurotonix, Starlab Neuroscience, Neuroelectrics, and NeuroSync, and is listed as an inventor on several issued and pending patents on the real-time integration of transcranial magnetic stimulation (TMS) with electroencephalography (EEG) and magnetic resonance imaging (MRI).

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