

EEG Deblurring Techniques in a Clinical Context

F. Cincotti¹, C. Babiloni^{2,5}, C. Miniussi⁴, F. Carducci^{2,4,5}, D. Moretti^{2,5}, S. Salinari³,
R. Pascual-Marqui⁷, P. M. Rossini^{4,5,6}, F. Babiloni²

¹IRCCS Fondazione Santa Lucia, Roma, Italy

²Dipartimento di Fisiologia Umana e Farmacologia, Università di Roma "La Sapienza", Roma, Italy

³Dipartimento di Informatica e Sistemistica "Ruberti", Università di Roma "La Sapienza", Roma, Italy

⁴IRCCS "San Giovanni di Dio" Istituto Sacro Cuore di Gesù, Brescia, Italy

⁵FaR and CRCCS Ospedale Fatebenefratelli, Isola Tiberina, Roma, Italy

⁶Cattedra di Neurologia, Campus Biomedico, Rome, Italy

⁷The KEY Institute for Brain-Mind Research, Zurich, Switzerland

Summary

Objectives: EEG scalp potential distributions recorded in humans are affected by low spatial resolution and by the dependence on the electrical reference used. High resolution EEG technologies are available to drastically increase the spatial resolution of the raw EEG. Such technologies include the computation of surface Laplacian (SL) of the recorded potentials, as well as the use of realistic head models to estimate the cortical sources via linear inverse procedure (low resolution brain electromagnetic tomography, LORETA). However, these deblurring procedures are generally used in conjunction with EEG recordings with 64-128 scalp electrodes and with realistic head models obtained via sequential magnetic resonance images (MRIs) of the subjects. Such recording setup it is not often available in the clinical context, due to both the unavailability of these technologies and the scarce compliance of the patients with them. In this study we addressed the use of SL and LORETA deblurring techniques to analyze data from a standard 10-20 system (19 electrodes) in a group of Alzheimer disease (AD) patients.

Methods: EEG data related to unilateral finger movements were gathered from 10 patients affected by AD. SL and LORETA techniques were applied for source estimation of EEG data. The use of MRIs for the construction of head models was avoided by using the quasi-realistic head model of the Brain Imaging Neurology Institute of Montreal.

Results: A similar cortical activity estimated by the SL and LORETA techniques was observed during an identical time period of the acquired EEG data in the examined population.

Conclusions: The results of the present study suggest that both SL and LORETA approaches can be usefully applied in the clinical context, by using quasi-realistic head modeling and a standard 10-20 system as electrode montage (19 electrodes). These results represent a reciprocal cross-validation of the two mathematically independent techniques in a clinical environment.

Keywords

Surface Laplacian, voluntary movements, motor control, cerebral cortex, LORETA

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1. Introduction

It is well known that the EEG potential distributions recorded from the scalp in humans are affected by low spatial resolution and by the dependence on the electrical reference used. Nowadays, modern high resolution EEG technologies are available to drastically increase the spatial resolution of the raw EEG. Such technologies include the computation of the surface Laplacian of the recorded potentials, as well as the use of realistic head models to estimate the cortical sources via a linear inverse procedure. The deblurring techniques are generally used in conjunction with EEG recordings with 64-128 scalp electrodes and realistic head modeling as obtained via sequential magnetic resonance images (MRIs) of the subject's head [1]. However, EEG recordings cannot be performed with more than 19 electrodes (standard 10-20 system) in a clinical context. In addition, it is difficult to acquire and manage MRIs for all patients, in order to build realistic head models. This occurs not only for the unavailability of the technical devices, but also for the emotional problems of some patients. For instance, Alzheimer Disease (AD) patients cannot be subjected to the long procedure of high-density electrodes montage (and relative lowering of electrode impedance). On the other hand, the use of spherical head models instead of the realistic ones decreases the precision of the deblurring techniques. All these considerations were at the basis of our study, aimed at analyzing the performance of two techniques for EEG source estimation such as the surface

Laplacian and low resolution brain electromagnetic tomography (LORETA) [2]. These techniques were applied to the EEG data of a group of AD patients and age-matched controls. EEG data were recorded from 19 scalp electrodes and a quasi-realistic head model was used instead of individual realistic head models. The quasi-realistic head model was realized at the Brain Imaging Neurology Institute of Montreal as an average of 152 individual MRI data sets. The surface Laplacian and LORETA solutions were co-registered with that model. Surface Laplacian and LORETA solutions were also compared to each other.

Convergent results obtained with SL and LORETA techniques suggest that such procedures can improve the efficiency of the spatial details ("deblurring") of the recorded EEG data, also in a clinical context, using the standard 10-20 system (19 scalp electrodes) and a quasi-realistic head model. From a clinical point of view, the computation of SL and LORETA was able to model the sources of movement-related cortical EEG rhythms in mild to moderate AD patients, with respect to an age-matched group of healthy subjects. The results are consistent with the working hypothesis that AD is a brain disease that involves not only the cognitive systems but also neural networks contributing to sensorimotor functions.

2. Methods

Subjects: The AD group consisted of 10 right-handed subjects with an age of $79.4 \pm SD 8.6$

(7 females and 3 males). Only patients with mild to moderate AD were included in the study. The control group consisted of ten right-handed healthy subjects with an age of $72.1 \pm \text{SD } 9.1$ (5 females and 5 males).

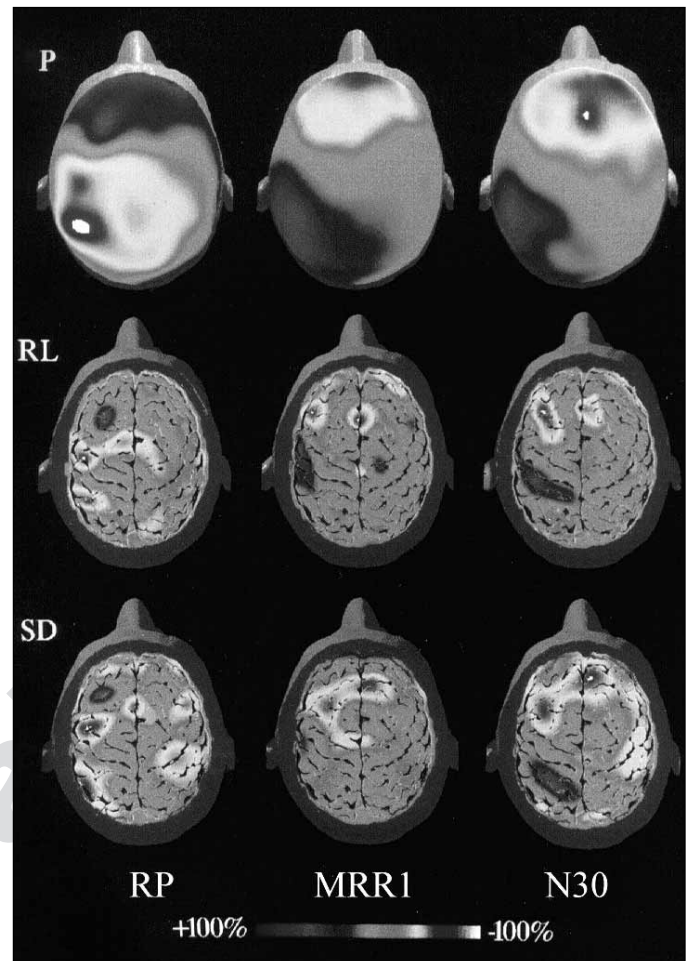
EEG recordings: The motor task consisted of brisk right middle finger extensions, which were triggered by a verbal instruction (i.e. “go”) of the experimenter (8-12 s inter-movement intervals). Nineteen scalp electrodes were positioned according to the international 10-20 system (Fp1, Fp2, F7, F43, Fz, F4, F8, T3, C3, Cz, C4, T4, T5, P3, Pz, P4, T6, O1, O2). EEG data were recorded with 0.3-70 Hz bandpass and linked-earlobes reference. Electrooculogram (0.3-70 Hz bandpass) and surface rectified electromyographic activity of bilateral extensor digitorum muscles (1-70 Hz bandpass) were also collected. For each subject, recorded EEG data were related to about 100 movements.

Spatial deblurring or source estimation: Artifact-free EEG activity (from 4 s before to 4 s after the onset of electromyographic response, zero time) was used as an input for EEG source estimation with SL and LORETA. EEG segments contaminated by blinking, eye movements, mirror movements, and/or other artifacts were rejected off-line. Artifact-free EEG activity was then considered for the SL estimate by spline function [1]. The single trial analysis was repeated on the SL data to discard the single trials contaminated by computational artifacts. Percentage event-related desynchronization/synchronization (ERD/ERS) of alpha and beta EEG rhythms was computed by the well-known procedure of (Graz) Pfurtscheller’s group. Amplitude gray scale maps of the alpha and beta ERD/ERS peaks were calculated on a 3-D “quasi-realistic” head model by a spline interpolating function, which was constructed based on the quasi-realistic head model of the Brain Imaging Neurologic Institute of Montreal.

LORETA is a well-known technique computing 3-D, discrete, linear solutions for the EEG inverse problem that have been previously tested with an extensive simulation study [2]. In brief, LORETA used a three-shell spherical head model, which was registered to the Talairach brain atlas [3] digitized from magnetic resonance im-

Fig. 1

Cross validation between the cortical source estimation performed with the surface Laplacian (RL row), the linear inverse solution (SD row) from the scalp EEG data (P row). The first column presents the distributions for the pre-movement readiness potential (RP) component of movement-related potentials (MRPs), the second one for the on-going movement evoked potentials (movement-related response 1, MRR1) and the third one for the negative component of median-nerve somatosensory evoked potentials occurring about 30 ms after stimulus delivery. Roughly the same information is conveyed by the two EEG spatial deblurring procedures.



ages at the Brain Imaging Neurologic Institute of Montreal. The LORETA solution space was restricted to the cortical gray matter and hippocampus, as determined by the digitized Probability Brain Atlas. The discretization of the solution space was performed with 2,394 voxels (7 mm resolution), each containing an equivalent current dipole. LORETA solutions were represented as axial, sagittal, and coronal slices ($17 \times 20 \times 25$). These slices were identified based on their distance (mm) from the origin of a 3-D Talairach head coordinate system. In this system, the positive x-axis passed approximately through the right ear (medial-lateral direction), the positive y-axis through the forehead (anterior-posterior direction), and the positive z-axis (representing the superior-inferior direction) through the top of the head.

LORETA provided current density values in each voxel of the source space

from individual EEG power spectrum (digital filtering, narrow bandpass, no phase shift). The EEG bands of interest were alpha (8-12 Hz), beta 1 (13-18 Hz), and beta 2 (19-21 Hz) and the analysis periods were the movement (“movement”) and post-movement (“post-movement”) ones. “Rest” ranged from -4 to -3 s, “movement” from zero time to $+1$ s, and “post-movement” from $+1$ to $+2$ s. Event-related LORETA solutions were computed based on the ratio between dipole power density values at “movement” (or “post-movement”) and rest. Event-related LORETA solutions in AD vs. N subjects were compared by paired t-test, corrected for multiple comparisons.

In both techniques, individual alpha or beta frequencies showing maximal differences between the “rest” and other periods of interest were chosen for the ERD/ERS computations. This provided individual (i)

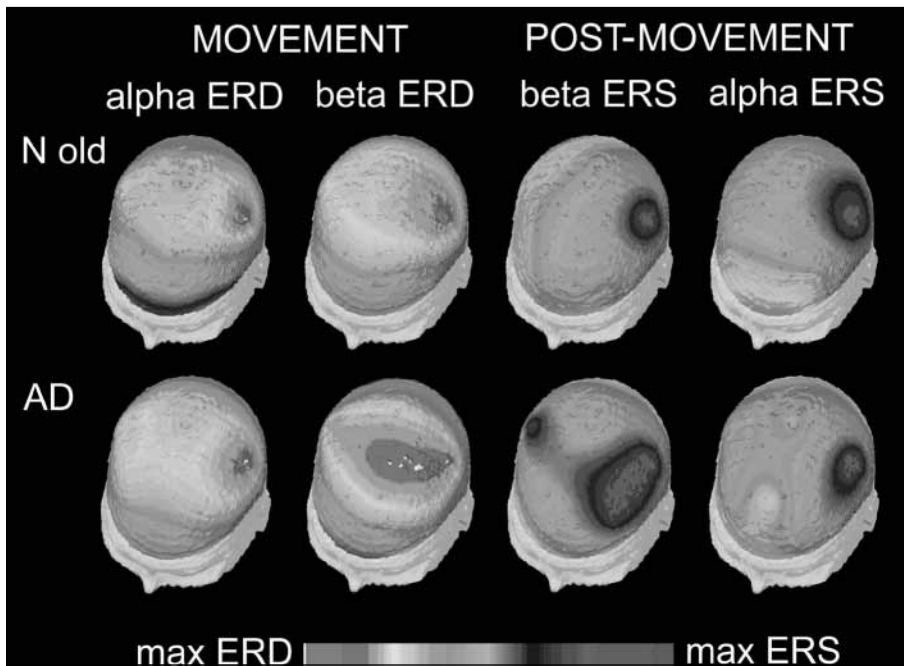


Fig. 2 Three-dimensional maps of across-subjects alpha and beta event-related desynchronization/synchronization (ERD/ERS) peaks in Alzheimer disease (AD) and normal subjects. These ERD/ERS peaks were computed during the movement and post-movement periods. Gray scale: maximum ERD and ERS are coded in light and dark gray, respectively. Note that AD data showed a paradoxical ipsilateral preponderance of post-movement ERS distributions. Of note, ERD/ERS was computed by the well known procedure of (Graz) Pfurtscheller's group.

“movement”/alpha reactivity band, (ii) “movement”/beta reactivity band, (iii) “post-movement”/beta reactivity band, for ERD/ERS computations.

Cross-validation between SL and linear inverse solutions. The comparisons between SL and linear inverse (mathematics at the basis of LORETA) solutions were accomplished by also using standard median-nerve somatosensory evoked potentials (SEPs) and averaged movement-related potentials (MRPs) recorded in normal subjects. This was because the neural sources of SEPs and MRPs are well known in literature by intra-cortical recordings.

3. Results

Figure 1 presents the results obtained by the two spatial deblurring/source estimation techniques on a common dataset of SEPs and MRPs recorded in one healthy subject. The SL distributions computed from SEPs and MRPs were projected radially onto the cortical reconstruction of a realistic head model for a better comparison with those

computed with the linear inverse technique. Each column in Figure 1 presents the raw scalp potentials, the SL solution, and the linear inverse solution for components of MRPs and SEPs. In particular, the first column presents the distributions for the pre-movement readiness potential (RP) component of MRPs, the second one for the on-going movement evoked potential (movement-related response 1, MRR1) and the third one for the negative component occurring after 30 ms of stimulus delivery for the SEPs (N30). It is well known that these components are generated in primary sensorimotor and supplementary motor cortical areas. It is interesting to note that the cortical potentials estimated with the two procedures are really similar, demonstrating the activation of the appropriate cortical areas. This result encouraged us to apply these deblurring tools to the EEG recordings in both normal and AD patients.

3.1 Application to Normal and AD Patients of SL Computation

The topography of the alpha and beta ERD/ERS peak obtained with the SL technique is illustrated in the gray scale maps of Figure 2. The “movement” alpha and beta ERD distribution of all groups was characterized by maximum response in the centroparietal areas of both sides (contralaterally preponderant). In addition, there was a marked “movement” beta ERD around scalp vertex in the control group with respect to the AD data. During the “post-movement”, the topographical ERS difference between the two groups appeared to be more pronounced. The “post-movement” beta ERS distributions showed a strong preponderant response on the contralateral central area in the control group. In contrast, AD group data showed a paradoxical emerging beta ERS maximum in the ipsilateral centroparietal area. Furthermore, the contralateral beta ERS maximum tended to be localized more anteriorly and to be distributed more largely in the AD than in the normal group.

3.2 Application to Normal and AD Patients of LORETA Computations

Figure 3 plots 3-D statistical maps of movement-related alpha and beta band power density changes computed by LORETA in AD patients vs. normal subjects. The maps were registered to Talairach space and digitized Probability Brain Atlas. For the “movement”, statistical maps showed significantly lower desynchronization of alpha and beta oscillations (i.e. reduction of the spectral power density during the event compared to baseline period) in the AD than in the control subjects (dark gray region) within contralateral inferior frontal gyrus (alpha $t = 3.41$, $p < 0.0016$), bilateral superior frontal gyrus (beta 1 $t = 3.93$, $p < 0.0005$), and supplementary motor area/cingulate gyrus (beta 1 $t = 3.65$, $p < 0.0009$ /beta 2 $t = 2.93$, $p < 0.0045$). In contrast, a lower desynchronization of alpha and beta oscillations in the normal group than in the AD one (light gray region) was computed but not showed here in contralateral (i.e.

left), middle temporal gyrus (beta 2 $t = 4.23$, $p < 0.0002$), and parietal lobe (alpha $t = 2.14$, $p < 0.02$). For the “post-movement”, not shown in the figure, statistical maps indicated a statistical trend for higher synchronization or “rebound” of alpha and beta oscillations in the AD than in the normal subjects within ipsilateral superior frontal gyrus (alpha $t = 1.97$, $p < 0.0322$). On the contrary, a statistical trend for higher synchronization of alpha and beta oscillations in the normals than AD subjects was observed in contralateral inferior parietal lobule (alpha $t = 2.40$, $p < 0.0137$).

4. Discussion

Results of the present study suggest that the use of deblurring tools like SL and LORETA improved the spatial details of the raw recorded EEG scalp potential distributions. In addition, similar cortical distributions were retrieved by processing SEPs and MRPS with the two procedures. In precedence, the agreement between linear inverse estimation and SL techniques was also assessed by using data simulations within an unrealistic spherical head model [4]. Although in the present study a formal correlation coefficient between the two cortical distributions was not computed, it seemed reasonable from the observation of Figure 1 to assert a general agreement between the two EEG spatial deblurring/source estimation methods. The application of EEG spatial deblurring/source estimation tools in a clinical context produced interesting results. These results showed that the activation of contralateral rolandic cortex and motor performance are preserved in mild to moderate AD. Moreover, SL (Fig. 2) and LORETA (Fig. 3) procedures demonstrated an abnormal recruitment of other functionally-related nonprimary frontal and ipsilateral sensorimotor areas in AD patients with respect to normal subjects. This would support the hypothesis that AD is a brain disease globally affecting also neuronal connectivity involved in the processing of sensorimotor information (despite no overt movement disorder).

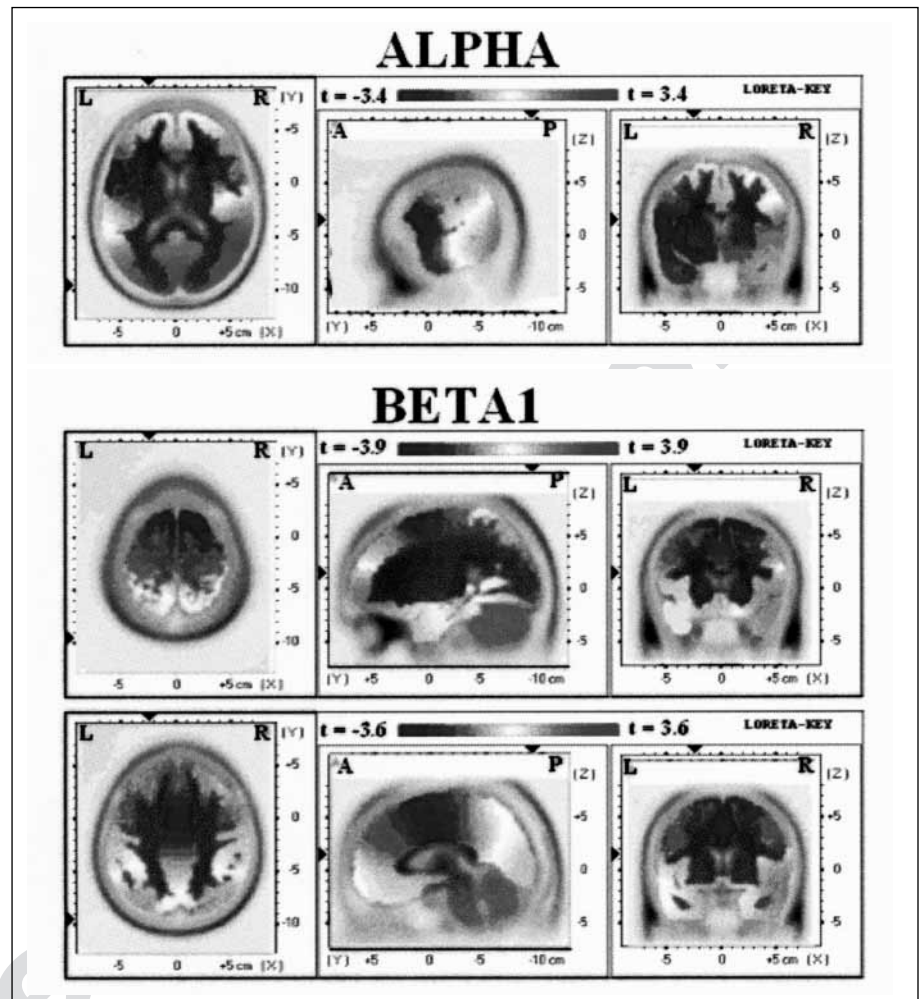


Fig. 3 Three-dimensional statistical maps of movement-related alpha and beta band power density changes computed by low resolution brain electromagnetic tomography (LORETA) in AD patients vs. normal age-matched subjects. Brain slices (axial, sagittal, coronal) illustrating LORETA solutions were identified based on their distance (mm) from the origin of the Talairach head coordinate system. Statistical comparisons were performed with a t-test. T values were represented by a gray percent scale. Negative t values (dark gray) mean lower power density in AD than normal age-matched subjects and viceversa for positive t values (light gray).

5. Conclusion

The results of the present study suggest that both SL and LORETA approaches can be usefully applied in the clinical context, by using quasi-realistic head modeling and the standard 10-20 system as electrode montage (19 electrodes). These results represent a reciprocal cross-validation of the two mathematically independent techniques in a clinical environment.

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Correspondence to:

Dr. Fabio Babiloni
Dipartimento di Fisiologia Umana e Farmacologia
Università di Roma “La Sapienza”
P.le A. Moro 5, 00185 Roma, Italy
E-mail: Fabio.Babiloni@uniroma1.it