N170 asymmetry as an index of inferior occipital dysfunction in patients with symptomatic occipital lobe epilepsy

Ricardo Lopes a,b,*, Pedro Cabral c, Nuno Canas c, Paula Breia d, John P. Foreid e, Eulália Calado a, Rita Silva a, Alberto Leal a,f

a Department of Pediatric Neurology, Hospital Center of Central Lisbon, Rua Jacinta Marto, 1169-045 Lisbon, Portugal
b Faculty of Psychology, University of Coimbra, Coimbra, Portugal
c Department Pediatric Neurology, Hospital Center of West Lisbon, Lisbon, Portugal
d Department Neurology, Hospital Garcia de Orta, Almada, Portugal
e Department Neurology, Portuguese Oncology Institute, Lisbon, Portugal
f Centre for Psychological Research and Social Intervention (Cis-IUL), Lisbon, Portugal

Article info
Article history:
Accepted 27 May 2010
Available online 20 June 2010

Keywords:
Epilepsy
N170
EEG
Childhood
Occipital lobe

Abstract
Objective: Localizing epileptic foci in posterior brain epilepsy remains a difficult exercise in surgery for epilepsy evaluation. Neither clinical manifestations, neurological, EEG nor neuropsychological evaluations provide strong information about the area of onset, and fast spread of paroxysms often produces mixed features of occipital, temporal and parietal symptoms. We investigated the usefulness of the N170 event-related potential to map epileptic activity in these patients.

Methods: A group of seven patients with symptomatic posterior cortex epilepsy were submitted to a high-resolution EEG (78 electrodes), with recordings of interictal spikes and face-evoked N170. Generators of spikes and N170 were localized by source analysis. Range of normal N170 asymmetry was determined in 30 healthy volunteers.

Results: In 3 out of 7 patients the N170 inter-hemispheric asymmetry was outside control values. Those were the patients whose spike sources were nearest (within 3 cm) to the fusiform gyrus, while foci further away did not affect the N170 ratio.

Conclusions: N170 event-related potential provides useful information about focal cortical dysfunction produced by epileptic foci located in the close neighborhood of the fusiform gyrus, but are unaffected by foci further away.

Significance: The N170 evoked by faces can improve the epileptic foci localization in posterior brain epilepsy.

© 2010 International Federation of Clinical Neurophysiology. Published by Elsevier Ireland Ltd. All rights reserved.

1. Introduction

Epileptic activity in occipital lobe epilepsies tends to propagate to nearby parietal, temporal and also to the distant frontal lobes (Leal et al., 2007, 2008; Williamson et al., 1992). Such phenomenon is responsible for the complex ictal manifestations involving diverse brain areas, making it difficult to obtain a clear picture of the overall dynamics and potentially leading to important errors when surgery for epilepsy is considered. With the exception of elementary visual symptoms, which are relatively rare, the clinical manifestations associated with occipital lobe epilepsy are not very informative about the particular area of onset of the epilepsy within the posterior brain (reviewed in Sveinbjörnsdottir and Duncan (1993)). Complex visual auras, hallucinations, amnesia, version of head and eyes, eye blinking, have a poor localizing value and can be obtained by electrical stimulation in both occipital, parietal and temporal lobes (reviewed in Geller et al. (2000)). Often the tendency for epileptic activity to involve the temporal lobes leads to impaired consciousness and to the inability to report ictal manifestations with localization information. Overall the symptomatology of posterior cortex epilepsy has a poor localizing value, requiring a careful integration of both clinical and neurophysiologic information to formulate good hypothesis about the localization of the epileptic foci.

With the exception of hemianopia, patients with posterior brain epilepsy rarely present with deficits in the neurological examination and in the subjective ability to process daily visual information. In fact classical neurological syndromes of visual agnosia
such as prosopagnosia are more often associated with bilateral lesions (Meadows, 1974) and are rarely found in connection with epilepsy.

Despite the fact that several neuropsychological studies have uncovered an asymmetry in memory for faces (Millner, 1968), these functions have been associated with the temporal lobes and in particular with hippocampus processing (Crane and Millner, 2002). The previous studies demonstrated that the capability to discriminate faces from other objects is not affected in temporal lobe lesions, supporting the thesis that such processing is done in the ventral temporal–occipital areas (Allison et al., 1994; Haxby et al., 1996; Kanwisher et al., 1997; Steeves et al., 2006; Pitcher et al., 2007). The neuropsychological tests presently available have nevertheless been unable to detect reliably abnormalities in face recognition (Duchaine and Weidenfeld, 2003; Duchaine and Nakayama, 2004), making it difficult to produce strong inferences about the localizations of occipital lesions from the obtained results.

The visual analysis of the EEG in occipital lobe epilepsies is poorly localized (Williamson et al., 1992) and the complex dynamics of spike activity requires the use of sophisticated methods of data analysis (Leal et al., 2007, 2008), which have not been used very often in the surgery for epilepsy setting.

In this study, we investigate the possible contribution of the N170, a negative event-related potential peaking around 170 ms, elicited by presentation of complex pictures such as faces (Bentin et al., 1996), to discriminate inferior occipital lobe epileptic foci. The amplitudes of the N170 have been found to be approximately symmetrical (Rossion et al., 2003), despite the well-known differential involvement of both hemispheres in face processing (reviewed in Damásio et al., 1990)). We hypothesize that patients with inferior occipital epileptic foci will show a smaller N170 in the affected hemisphere when compared with the spared one and that the resulting inter-hemispheric asymmetry will provide useful neurophysiologic information about the localization of epileptic foci involving the inferior part of the occipital lobes.

2. Methods and subjects

We selected patients with symptomatic epilepsy of the posterior brain, undergoing evaluation at the program for Surgery of Epilepsy of the Hospital Center of West Lisbon, Portugal. The neurophysiologic studies were conducted as an add-on to the conventional video-EEG monitoring, and were performed before that procedure. Informed consent was obtained from the patients or from their parents. Clinical and neurophysiologic data for the patients is shown in Table 1. All subjects had normal or corrected to normal vision.

The neurophysiologic study consisted in a high-density EEG recording using a cap (EasyCap, Herrsching-Breitbrunn, Germany) with 78 electrodes, including all positions in the 10–10 system plus positions FT11, FT12, TP11, TP12. Particular care was taken to ensure a symmetrical fit of the cap on the head and that the T7/8, P1/2 and O1/2 electrodes were at the conventional 10–20 system positions. Ring electrodes made of sintered AgCl were applied at the cap and connected to two NuAmps 40-channels EEG amplifiers (Neuroscan, Charlotte, USA) chained in such a way as to provide a system with 78 channels. The sampling rate was 1000 Hz and the high- and low-pass filters were 0.5 and 70 Hz.

The N170 was obtained by presentation of neutral emotion black and white faces taken from the Psychological Image Collection at Stirling (PICS) faces database (University of Stirling Psychology Department, Stirling, Scotland – United Kingdom) or 1/2 front view grayscale pictures of cars, using the procedure described by Rossion et al. (2003) (Fig.1a). A collection of 18 different faces (9 males and 9 females) and 18 different cars were presented twice in a random sequence, using the E-Prime 1.2 software package (PST, Pittsburgh, USA). Each picture was shown for 250 ms in a CRT screen, with an inter stimulus interval of 1 s. Four of the previous block sequences were collected for each subject, with an interval of 2 min. In order to maintain attention, subjects were asked to keep a mental recording of the number of female faces.

Thirty healthy volunteers, aged 19–44 years, were included in the study and used as a control group. For epileptic patients an additional sleep recording with an average duration of 30 min was done after the evoked potential acquisition. Intercital spikes with good signal/noise (S/N) ratio were visually identified by an experienced clinical neurophysiologist (AL), and averaged with synchronization by the peak of the spike.

2.1. Source analysis

In order to transform the information from the 78 channel EEG recording to a ratio expressing the relative contribution of each hemisphere, we performed source analysis with the Source 2 software package (Neuroscan, Charlotte, USA), using a standard realistic finite element model (FEM) of the head, (Fig.1c). Standard electrode coordinates for the cap were downloaded from EasyCap, Inc. (http://www.easycap.de), which were then transformed to the Pre-Auricular-Nasion (PAN) coordinate system. The conductivities were 0.33 S/m for brain and scalp, 0.0042 S/m for the skull (Gonçalves et al., 2003). A regional source dipole was placed at the center of the fusiform gyrus (FG) of each hemisphere (Table2) and the model was fitted to the N170 scalp potential, (Fig.1c) The amplitude of each regional dipole at the peak of the Mean Global Field Power (MGFP) (Lehmann and Skrandies, 1984) of the N170 was used to calculate the left/right (L/R) hemisphere ratio, (Fig.1d).

The brain localizations of the sources of interictal spikes and N170 were studied in patients with the sLORETA method (Pascual-Marqui, 2002; Pascual-Marqui et al., 2002), as implemented in a free downloadable package (http://www.uzh.ch/keyinst/lore-ta.htm), and using the previously described volume conductor model. The solution space was anatomically restricted to the cortex and consisted of 6239 cubic voxel elements with 5 mm side. The instantaneous solution at the spike/N170 peak was used for the mappings shown in Figs. 2 and 3.

Table 1 Clinical and neurophysiological data.

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Age</th>
<th>Sex</th>
<th>Age at onset</th>
<th>MRI scan lesion location</th>
<th>Seizure type</th>
<th>N170 Asymmetry?</th>
</tr>
</thead>
<tbody>
<tr>
<td>ER</td>
<td>43</td>
<td>F</td>
<td>8</td>
<td>Lateral right occipital lobe</td>
<td>Complex partial seizures</td>
<td>No</td>
</tr>
<tr>
<td>MCR</td>
<td>47</td>
<td>F</td>
<td>10</td>
<td>Mesial right occipital lobe</td>
<td>Complex partial seizures</td>
<td>No</td>
</tr>
<tr>
<td>ML</td>
<td>26</td>
<td>F</td>
<td>19</td>
<td>Inferior left temporal lobe</td>
<td>Complex partial seizures</td>
<td>No</td>
</tr>
<tr>
<td>CC</td>
<td>7</td>
<td>F</td>
<td>5</td>
<td>Inferior left temporal lobe</td>
<td>Complex partial seizures</td>
<td>No</td>
</tr>
<tr>
<td>JC</td>
<td>20</td>
<td>M</td>
<td>11</td>
<td>Lateral left occipital lobe</td>
<td>Complex partial seizures</td>
<td>Yes</td>
</tr>
<tr>
<td>MSR</td>
<td>5</td>
<td>M</td>
<td>1</td>
<td>Lateral right temporal lobe</td>
<td>Complex partial seizures</td>
<td>Yes</td>
</tr>
<tr>
<td>GM</td>
<td>9</td>
<td>M</td>
<td>7</td>
<td>No lesion</td>
<td>Complex partial seizures</td>
<td>Yes</td>
</tr>
</tbody>
</table>
Fig. 1. (a) Examples of pictures shown to obtain the N170 potential. (b) Butterfly plot of the 78 channels N170 (above), with the Mean Global Field Power (MGFP) below. (c) Standard realistic volume conductor model, with the electrodes (above), and with regional dipoles at FG coordinates (below). (d) Source activity for the two regional dipoles at N170, right dipole above, left dipole, middle. The activities at the time of MGFP peak (below) were taken to determine the N170 ratio. (e) Ratio between left and right hemisphere N170 for the control group \((N = 30)\), for the presentation of cars (left) and faces (right). The results for faces show a reduced dispersion compared the one for cars.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Lesion (L)</th>
<th>Fusiform gyrus (FG)</th>
<th>Spike source (SS)</th>
<th>DSS-L</th>
<th>DSS-FG</th>
<th>DL-FG</th>
<th>Ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>x</td>
<td>y</td>
<td>z</td>
<td>x</td>
<td>y</td>
<td>z</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ER</td>
<td>28</td>
<td>–66</td>
<td>19</td>
<td>37</td>
<td>–51</td>
<td>–25</td>
<td>20</td>
</tr>
<tr>
<td>MCR</td>
<td>17</td>
<td>–77</td>
<td>5</td>
<td>37</td>
<td>–51</td>
<td>–25</td>
<td>20</td>
</tr>
<tr>
<td>JC</td>
<td>–40</td>
<td>–85</td>
<td>0</td>
<td>–37</td>
<td>–51</td>
<td>–25</td>
<td>–60</td>
</tr>
</tbody>
</table>
3. Results

3.1. Source analysis of interictal spikes

The preliminary visual analysis of the instantaneous scalp potential topographic maps of the average interictal spikes for the epileptic patients revealed spatially consistent patterns from spike onset until the peak for each patient, suggesting stable spatial configurations for the underlying generators. Source analysis was performed at the peak of the spike because this is the point with the best S/N ratio.

The visual analysis of the orientation of the field of potential over the scalp revealed very distinct patterns for the different patients (Figs. 2 and 3), providing support for the existence of sources with diverse localizations: patients GM, JC and MSR have vertical orientations with phase reversal near the inferior border of the brain, suggesting an origin of the abnormal activity in the inferior occipital cortex; patient ML has a similar field orientation, but shifted to a more anterior localization, suggesting an origin in the inferior cortex of the temporal lobe; patients ER and CC have an horizontal and transverse field orientation with clear maximum over the ipsilateral occipital lobe, suggesting a lateral cortex involvement; patient MR presented a horizontal and anterior-posterior orientation centered over the occipital lobe, compatible with a medial origin.

The sLORETA algorithm produced scores with clear maximum in the areas previously postulated for the different patients,
An analysis of the spatial localization of these sources revealed that they mostly lie in the neighborhood of the structural lesions (average distance of 2.3 cm, Table 2), suggesting that sub-lobar resolution is possible with this method in the posterior brain.

3.2. N170 ratio in patients

A distinct N170 potential was obtained for all healthy volunteers and patients participating in the study, both for faces and cars. The L/R hemisphere ratio of the absolute amplitude of the two regional dipoles generating the N170 potential for the two categories of stimuli revealed a reduced dispersion for faces in the control group (Fig. 1b). This stimulus category was then chosen for comparison with the patient data.

The N170 L/R ratio for the patients revealed values within the control range for four cases (ML, MR, ER and CC) and abnormal (values outside the range of the control group and more than 2 SD away from the mean) results for the other three (GM, JC and MSR) (Table 2). An analysis of the distance from the epileptic spike source to the nearest FG reveals that for the three patients with abnormal L/R ratios it is smaller than for the patients with normal ratios (Table 2). The critical distance separating the two groups of patients seems to be around 3 cm, as is suggested by the crossing
of the statistical threshold of significance of the L/R N170 ratio for distances higher than this value in the plot of Fig. 4. These results suggest that only epileptic foci within the close neighborhood of the FG are able to produce an asymmetry change in the N170 potential for faces.

The distances from the FG to the structural lesions are also superior to 3 cm for the patients with normal L/R ratio (Table 2). This distance is nevertheless a worse predictor of N170 asymmetries as patients JC and MSR have lesions at 4.2 and 3.7 cm away, respectively, from the FG but an abnormal L/R N170 ratio (Fig. 2b and c).

Overall these results suggest that only epileptic sources in the close proximity of the FG were able to significantly change the L/R N170 ratio, in patients with posterior cortex epilepsy.

4. Discussion

The main conclusion of this study is that an abnormal N170 ratio for faces is produced by epileptic sources localized near (<3 cm) the FG, but not by sources further away, and therefore can be used as a physiological marker of an epileptic source localized in the inferior occipital cortex of patients with symptomatic occipital lobe epilepsy.

Interictal epileptic activity in the occipital lobe epilepsies demonstrates a strong tendency to spread to nearby parietal and temporal lobes (Leal et al., 2007), to the contra lateral hemisphere and even to the frontal lobes (Williamson et al., 1992). This fast spread dynamics makes it difficult to localize the epileptogenic cortex not only in the raw EEG traces (Leal et al., 2008), but also using source analysis methods (Van der Meij et al., 1997). Several issues may be responsible for the poor yield of the latte techniques but the reduced spatial sampling over the posterior and inferior scalp, usu- 

al used in clinical studies, seems to be a prominent one (Leal et al., 2008). Careful studies comparing the localization of sources obtained by solving the inverse problem and well-known occipital foci are lacking in the literature, but would represent a major contribution to establish the contribution of source analysis in this clinical setting.

We used a standard FEM model with standard electrode positions to solve the inverse problem, which can result in a decreased accuracy as compared with individual realistic modeling. Nevertheless a comparison between standard and realistic models for spike data by Fuchs et al. (2002) failed to find systematic differences in localization, and the mean localization differences were in the order of 1 cm. In the same line the effect produced by small inaccuracies in the position of the electrodes seems to have a reduced effect on the final dipole localization error (Khosla et al., 1999; Wang and Gotman, 2001). The use of an electrode cap, ensuring a symmetrical disposition of the electrodes, strict adherence to the 10–20 system rules, minimizing electrode position inaccuracies, and the use of a high spatial sampling, in our view resulted in robust modeling of the intracranial generators. In particular the calculation of the inter-hemispheric ratio of the N170 could be affected by asymmetries either in the placement of the electrodes or by large anatomical head shape differences, neither of which are present in our data. Since we used symmetrical regional dipoles to express the activity of the inferior occipital cortex, the N170 ratio is not expected to be significantly affected by small inaccuracies in the postulated localization.

In our patients, we used anatomical localization of cortical lesions as a surrogate marker for the unknown localization of intracranial epileptic foci. Although this procedure may not provide the ideal localization, in our view, it is reliable enough to provide sub-lobar resolution, allowing us to discriminate among the inferior, lateral and medial occipital cortex subdivisions of each hemisphere. The exception seems to be patients JC and MSR, which have lesions in the lateral occipital lobe, but topography of the epileptic spikes with a vertical orientation and phase reversal near the scalp projection of the lateral and inferior occipital cortex, suggesting an origin in the inferior occipital cortex. The latte localization agrees well with the localization of the source recovered by the sLORETA algorithm (Figs. 2b and c).

Overall the spatial distance between the calculated sources and the associated lesions for the five patients is on average 2.3 cm, which is very close to the resolution power of source analysis methods using realistic models for point sources (Fuchs et al., 2001). This result supports the suggestion that the methodology may provide sub-lobar resolution for the epileptic foci in occipital lobe epilepsy.

The N170 potential for faces has demonstrated slightly higher amplitude in the right hemisphere as compared with the left one, but in most studies the difference did not reach statistical significance (Bentin et al., 1996). In line with these results in our study the control group showed a L/R ratio slightly bellow 1, but with no significant asymmetry (Fig 1e).

None of our patients complained of difficulties in subjective visual recognition of faces, even those with an abnormal N170 ratio, a fact that most likely relates to the independent capability of each hemisphere to perform this function (Levy et al., 1972).

Three patients presented an age in the pediatric range, making the comparison with the control group of adults less reliable than for all the other patients. The two patients with asymmetrical N170 (MSR and GM) both presented very small distances from the spike source to FG (1.6 and 0.6 cm), while the one with normal ratio (CC) showed a larger distance (4.7 cm); Previous studies in the normal developmental aspects of the N170 throughout childhood, demonstrating preserved symmetry (Taylor et al., 1999), suggest that an abnormal left hemisphere function it is the most likely explanation for the inter-hemisphere N170 difference.

Rosburg et al. (2010) studied the effect on the vertex positive potential (VPP) of face inversion in a heterogeneous sample of epileptic patients undergoing surgery for epilepsy. Lateralization effects were not searched for, and no correlation of the findings with the particular type of epilepsy was performed. Because the VPP merges contributions from both hemispheres, no independent evaluation of those effects in the published material is possible.
Despite the small patient sample, the overall results of our study suggest that the N170 for faces can be a useful marker of inferior occipital lobe dysfunction, providing additional information on the localization of the epileptic focus when the L/R inter-hemisphere ratio is significantly altered. The association of this neurophysiologic parameter with the conventional neuropsychological tests for face processing could provide more robust inferences on occipital lobe dysfunction in patients with focal epilepsy of the posterior brain.

Acknowledgements

The authors are grateful to Heloisa Silva and Daniela Dias for their technical support.

Ricardo Lopes has been supported by the Grant SFRH/BD/65617/2009 from the Portuguese Foundation for Science and Technology (FCT).

References


