Epilepsy and phenylketonuria: a case description and EEG-fMRI findings

Summary

Phenylketonuria (PKU) is characterized by phenylalanine accumulation due to phenylalanine hydroxylase deficiency. Up to 50% of PKU patients experience seizures. We evaluated an adult PKU patient who suffered from absences and primarily generalized tonic-clonic seizures in adolescence and early adulthood, associated with features of idiopathic generalized epilepsy (IGE) on EEG. The fMRI pattern observed in our patient during GSWs is similar but not identical to that observed in idiopathic generalized epilepsy, suggesting different neurophysiological mechanisms. This is the first description of BOLD-fMRI patterns in a PKU patient with epilepsy. Similar studies in more patients might help to uncover the pathophysiology of seizures in this disease.

Introduction

Phenylketonuria (PKU) is a metabolic disorder, characterized by accumulation of phenylalanine (Phe) due to phenylalanine hydroxylase (PAH) deficiency. If untreated, it leads to an abnormal phenotype including growth failure, intellectual impairment and seizures. PKU patients may experience different types of seizures, including generalized ones.

Case report

History and clinical features

A 45-year-old Caucasian male was admitted as an outpatient to our epilepsy center at the age of 40. His childhood history showed, from the age of two years, delayed acquisition of walking and language milestones and mild learning difficulties. At the age of 16 years, he experienced a first seizure characterized by eyelid myoclonias and brief psychomotor arrest. In the following months he experienced similar seizures, sometimes triggered by watching television. A year later a generalized tonic-clonic seizure occurred. Carbamazepine induced an increase in seizure frequency, and was thus substituted with ethosuximide and primidone, which gave reasonable seizure control. At the age of 18 years, the patient was diagnosed with PKU in a pediatric neurology unit: his blood and urinary Phe levels on admission were 2516 µmol/l and 297 mmol/l, respectively; PAH gene mutation was confirmed. EEG on admission showed prolonged bursts of 3-4 Hz generalized spike-and-wave discharges (GSWs), which increased during intermittent light stimulation (ILS) (Fig.1A).

A Phe-poor diet was promptly started, primidone was discontinued and the ongoing ethosuximide was asso-
associated with phenobarbital. Within one month cognitive function and seizures had already significantly improved and blood and urinary Phe levels had fallen significantly. Routine EEG had improved too; few GSWs were recorded and were not increased by ILS. Thereafter, the patient experienced absence episodes and sporadic eyelid myoclonias only 2-3 times per year up to the age of 35 years when lamotrigine was associated with phenobarbital and ethosuximide, resulting in complete resolution of absences and eyelid myoclonias up to the present time. EEGs performed annually at our center consistently reveal bursts of 3-4 Hz diffuse sharp waves and GSWs (Fig.1B), not modified by hyperventilation or ILS. A 3T brain MRI scan showed a mild subcortical white matter alteration (Fig.1C).

**EEG-fMRI**

The patient underwent an EEG-fMRI investigation performed using a 3T MR scanner. fMRI data were obtained by a T2*-weighted gradient recalled echo-planar imaging sequence (TR 2500 msec, TE 40 msec, FA 90°, image matrix 128 × 128, in plane field of view 220 × 220 mm²) with 28 interleaved slices (slice thickness 5 mm) parallel to the anterior-posterior commissural plane, repeated over 300 volumes for a total scanning time of 12 min 30 sec. The EEG was recorded with an MR-compatible cap as described elsewhere (Pesaresi et al., 2011). Data were analyzed using the FEAT v5.98 tool of the software-package FSL 4.1.5 (http://www.fmrib.ox.ac.uk/fsl/). The preprocessing of the dataset consisted of spatial smoothing with a 6 mm Gaussian kernel, temporal filtering with a 100 sec high-pass filter, slice-time correction and motion correction. We explored a time window from 10 sec before until 10 sec after the onset of IEDs, proceeding in 2.5 sec steps. For each step we built up a general linear model (GLM) composed of IED regressor convolved with a Gamma hemodynamic response function (HRF). Motion parameters were also added to the GLM as covariates of no interest. In total, 12 GLMs were computed and statistical maps were obtained with a cluster threshold of z=2.3 and applying a p-value =0.05. This study was approved by our institutional ethics committee and the patient gave his written consent to the publication of this case report.

**Interictal EEG-fMRI findings**

The resulting statistical maps at the level of the subcortical structures (Fig. 2A) revealed deactivation of the left thalamus and the head of the right caudate nucleus from 5 sec before to 5 sec after onset of IEDs. At cortical levels, statistical maps showed early activation of the left perirolandic cortex (10 sec before IEDs), followed by a BOLD signal decrease in cortical regions belonging to the so-called default mode network (DMN), i.e. the precuneus, lateral parietal cortex, frontomesial cortex and prefrontal cortex, from 5 sec before to 2.5 sec after onset of IEDs. A late BOLD signal decrease was detected in the left frontoparietal cortex (7.5 sec after onset of IEDs) (Fig. 2B).

**Discussion**

Phenylketonuria is caused by an inborn error of Phe metabolism (Williams et al., 2008) due to mutation of
the PAH gene located on chromosome 12q23.2 (Lidsky et al., 1985). Because of the enzyme failure, Phe cannot be transformed into tyrosine (Tyr): the accumulation of toxic byproducts of Phe and the deficiency of Tyr and its downstream products (catecholamines, melanin, thyroxine) together cause the PKU phenotype, which includes growth failure, intellectual impairment and seizures. Several factors contribute to the neurotoxicity occurring in PKU. Brain MRI examinations performed in adult PKU patients invariably show white matter abnormalities (Thompson et al., 1990) and post-mortem brain examinations in PKU patients reveal hypo- or demyelination, independently of a Phe-poor diet (Dyer, 1999). In line with these reports, our patient’s brain MRI showed the presence of white matter abnormalities despite the continuation of dietary treatment in adulthood.

There exist no detailed descriptions of the epilepsy/seizure phenotypes in PKU, although generalized tonic-clonic seizures and myoclonic absences have been suggested to occur in up to 25% of treated PKU patients (Brenton and Pietz, 2000). Furthermore, EEG abnormalities are frequently observed even in adult patients under a strict dietary treatment (Gross et al., 1981), a finding that suggests a permanent alteration. The mechanisms linking Phe levels and seizures are far from clear. However: i) high Phe concentrations are associated with a decrease in brain levels of norepinephrine and serotonin (Puglisi-Allegra et al., 2000), which have anticonvulsant effects (Giorgi et al., 2003); ii) in mice bearing the human PAH gene mutation a strict interplay between Phe levels and glutamatergic transmission has been observed: due to an inhibitory effect of Phe upon glutamatergic transmission, these mice might have a latent glutamatergic hyperexcitation linked to chronic compensatory upregulation of glutamatergic receptors caused by high pre-treatment Phe levels (Martynyuk et al., 2007). Interestingly, these mutant mice show increased susceptibility to audiogenic seizures, a model of generalized seizures (Martynyuk et al., 2007).

Interictal epileptiform discharges have been present in our patient, in a varying fashion, since a young age: typical GSWs in adolescence/early adulthood and diffuse sharp waves/GSWs in his forties, with disappearance of ILS-triggered bursts. Our patient’s features fit those of an IGE syndrome, since: i) his EEG constantly showed 3-4 Hz GSWs; ii) he reported a history of both absences and generalized tonic-clonic seizures; iii) during his adolescence he experienced ILS-triggered GSWs, which are often described in IGE, especially juvenile myoclonic epilepsy and childhood-juvenile absence epilepsy (CAE-JAE).

The patient also met at least some of the criteria for a diagnosis of eyelid myoclonia with absences (EMA) (Panayiotopoulos, 2005), since he experienced eyelid myoclonia and brief absences found to be related to EEG GSWs and triggered by ILS; however, he never experienced epileptiform discharge during eye closure – this is considered a hallmark of EMA (Panayiotopoulos, 2005) – nor was such discharge mentioned in previous EEG reports.

At least some aspects of the fMRI-BOLD signal modification we observed in our patient are similar to what has been described during GSWs in IGE patients in other EEG-fMRI studies. In particular, deactivation of
the so-called DMN after onset of GSWs has invariably been described during GSWs in IGE, and particularly in CAE-JAE (Aghakhani et al., 2004; Hamandi et al., 2006; Moeller et al., 2008; Carney et al., 2010; Benuzzi et al., 2012) in which it has been related, theoretically, to the vigilance impairment occurring during prolonged GSWs. Conversely, in many IGE patients, thalamic activation is described at onset of GSWs (Aghakhani et al., 2004; Hamandi et al., 2006; Moeller et al., 2008; Carney et al., 2010; Benuzzi et al., 2012), whereas we detected only an early monolateral thalamic deactivation. Our patient also showed brief unilateral caudate deactivation, similar to that recently reported in a girl with GSWs (Moeller et al., 2010a). Finally, as regards the fMRI findings at cortical level, some studies reported BOLD signal activation in different cortical areas before onset of GSWs (Benuzzi et al., 2012; Moeller et al., 2010b). In a detailed study by Moeller et al. (2010b) in patients with CAE, an early asymmetrical frontal cortical activation was observed in most subjects during absence seizures, similar to what we found in our patient. According to the authors, the difference with other studies describing symmetrical cortical activation in patients with IGE could be related to the different statistical approaches used (individual pattern versus group analysis). In any case, the above findings support the hypothesized role of interplay between the thalamus and the neocortex in the generation of GSWs (Blumenfeld, 2005). Interestingly, a similar left precentral cortex activation during brief diffuse spike-wave discharges has also recently been described (Vaudano et al., 2012) in a patient with reading epilepsy; even though our patient did not show the clinical features of such a localization-related epilepsy, we cannot exclude that, at least in part, these two patients might share some common functional substrates for the generation of IEDs. Finally, it is interesting to note that BOLD activation of the thalamus and DMN deactivation are constantly observed also in EMA patients, during ictal/interictal GSWs (Liu et al., 2008). It is worth mentioning some methodological aspects of our study in which, like other authors (Bagshaw et al., 2004; Jacobs et al., 2009; Moeller et al., 2010a), we investigated multiple time intervals from IED onset. However, our study differed from previous ones in that we did not combine different BOLD maps in order to establish the statistically prevalent BOLD phenomenon, but simply reported all the BOLD changes observed in subsequent time intervals, as we were interested in their temporal evolution. Assuming the existence of dynamic processes that evolve over time, we did not compare alternative hypotheses (i.e. alternative HRFs) and did not perform any correction for the number of HRFs employed as they were included in separate GLMs. It is reasonable to run each analysis with the smallest possible p-value in order to reduce the total false positive rate of the study. The possibility of false positive results should however be borne in mind when interpreting BOLD statistical maps.

In conclusion, some of the cortical BOLD signal changes observed in our patient during GSWs resemble those observed in IGE, and this might be related, at least in part, to the similarity in the interictal EEG patterns between IGE patients and PKU patients with absences. However, the precise meaning of these fMRI features in IGE, particularly with regard to their pathogenetic implications, is still far from being elucidated.

By the same token, the lack of a thalamic activation, which is a frequent fMRI feature in IGE, suggests that: i) our PKU patient is unlikely also to have IGE as an independent disease; ii) almost identical electroclinical phenotypes are not necessarily underlain by the same pathogenesis; iii) the thalamic BOLD activation in IGE may not be as critical for the generation of GSWs as currently believed.

Given the relatively high incidence of PKU among the general population and of seizures among PKU patients, large studies on the characteristics, pathogenesis and possibly fMRI features of the epileptic syndromes occurring in this condition are required to extend our knowledge of this disease, as well as to disclose new mechanisms in seizure generation.

References


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