Cortical and Non-Cortical Myoclonus of Creutzfeldt-Jakob Disease

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Dear Sir,

The characteristic symptoms of Creutzfeldt-Jakob disease (CJD) include rapidly progressive cognitive decline and myoclonus [1]. Myoclonus in CJD patients is supposed to be associated with the appearance of periodic synchronous discharges (PSDs) on routine electroencephalograms (EEGs) [2]. However, a casual relationship between myoclonus and PSDs has not been established. Here, we report the case of a CJD patient with positive myoclonus and analyze the cortical potentials locked to the myoclonic jerks.

A 75-year-old man developed difficulty in targeting and catching objects with his left hand 3 weeks before visiting our neurological clinic. One month after the onset of initial symptoms, he started experiencing semi-rhythmic jerks in the left hand that were sensitive to pinprick stimuli. In the following 2 months, the patient showed significant memory and cognitive impairment, whereas muscle tone, muscle strength, and deep-tendon reflexes were intact. On examination, alien hand phenomenon without intermanual conflict was noted. The left hand tended to wander independently and grope nearby objects. Cerebrospinal fluid samples were negative for protein 14-3-3. The possibility of disorders such as thyroid function disorder, vitamin B12 and folate deficiencies, hepatitis B, vasculitis, and syphilis was excluded by the results of biochemical screening tests. EEGs showed PSDs with the largest amplitude over the central parietal area of the right hemisphere (i.e. C4 and P4; fig. 1a). Magnetic resonance imaging study showed restricted diffusion over the right frontal, parietal, and occipital lobes in diffusion-weighted imaging (DWI) (fig. 1b). 99mTc-TRODAT-1 revealed a bilateral decrease in striatal uptake values (fig. 1c). The patient’s symptoms continuously deteriorated. The patient became bedridden, and showed akinetic mutism and loss of swallowing capabilities at around 2.5 months after the disease onset. According to WHO criteria (1998), probable CJD was diagnosed. The WHO criteria had a diagnostic accuracy of 96.6% in a follow-up study performed in 9.9% of the patients diagnosed with probable CJD on the basis of the relationship between the results of biochemical screening tests. EEGs showed PSDs with the largest amplitude over the central parietal area of the right hemisphere (i.e. C4 and P4; fig. 1a). Magnetic resonance imaging study showed restricted diffusion over the right frontal, parietal, and occipital lobes in diffusion-weighted imaging (DWI) (fig. 1b). 99mTc-TRODAT-1 revealed a bilateral decrease in striatal uptake values (fig. 1c). The patient’s symptoms continuously deteriorated. The patient became bedridden, and showed akinetic mutism and loss of swallowing capabilities at around 2.5 months after the disease onset. According to WHO criteria (1998), probable CJD was diagnosed. The WHO criteria had a diagnostic accuracy of 96.6% in a follow-up study performed in 313 patients [1]. Protein 14-3-3 was absent in 9.9% of the patients diagnosed with probable CJD on the basis of the criteria [1].

Electrophysiological Recordings

The myoclonic jerks were recorded by multi-channel surface electromyography (EMG). Jerk-locked back-averaged EEG recording was performed with three gold-plated electrodes (C3, Cz, and C4) affixed to the scalp. We adopted EMG onset of spontaneous myoclonic jerks from left-hand triceps as the trigger for the EEG back-averaged sampling (sampling rate 2 kHz, filtered band pass 0.05–70 Hz). Artifact-free EEG epochs from 200 ms before to 300 ms after EMG onset were used for averaging. Somatosensory evoked potential (SEP) was also recorded.

Results

Surface EMG recording revealed semi-rhythmic bursts for the recorded muscles at a duration of around 60 ms (fig. 1d). On the basis of the relationship between the left-hand triceps jerks and the EEG signals, four distinct types (A–D) of conditions were characterized (fig. 1e). Type-A jerks had a mean duration of 60.35 ± 10.11 ms and mean amplitude of 179.13 ± 64.12 μV. The mean latency of the jerk onset to the maximal EEG negativity was 24.22 ± 11.07 ms. In comparison to type-A jerks, type-B jerks had a smaller amplitude (60.86 ± 40.43 μV) and similar duration (56.85 ± 9.01 ms). No significant EEG potentials were detected by locking the onset of type-C jerks (mean duration 32.45 ± 3.69 ms, mean amplitude 24.3 ± 3.40 μV). Some PSD were not jerk-locked (type D). No giant SEP was found.
Fig. 1. a Routine EEG study shows PSDs with predominance over the C4 and P4 electrodes. b Increased signal intensity in DWI at right frontal, parietal, and occipital cortical areas. c 99mTc-TRODAT-1 SPECT analysis of the brain shows decreased uptake in right striatum (1.12) and left striatum (1.07). The specific uptake is obtained by subtracting the mean counts per pixel in the occipital cortex (OC) and the mean counts per pixel in the whole striatum (ST) and dividing the result by the mean counts per pixel in the background (equations: (ST – OC)/OC; referential normal values in striatum: 1.69–2.15). d Surface EMG recorded from the left-hand triceps muscle of the patient. The signals show a semi-rhythmic pattern of the EMG bursts. e Four types of the averaged EMG and EEG recorded from the patient. Type A and B myoclonic jerks are locked to the cortical potentials, most obviously at C4, and are shown in the first and second rows, respectively. The third row demonstrates the absence of time-locked cortical potentials for some of the myoclonic jerks. The fourth row shows the cortical potentials without time-locked myoclonic jerks. The numbers of the averaged EEG epochs are 193, 43, 129 and 114 for types A, B, C, and D, respectively.
PSD is a common and characteristic EEG finding in CJD patients [3]. An abnormal subcortical pacemaker, which is probably synchronized with the cortical potentials, was supposed to play a pivotal role in the PSD generation [4]. Impairment of normal inhibitory function at the cortical level may further amplify the influence of this pathological pacemaker [4]. Myoclonus is another important symptom for the diagnosis of CJD and often diffuse, generalized and relatively rhythmic in CJD patients. In asymmetric cases, the jerks are usually accompanied by dystonia or alien hand syndrome [5]. Both positive and negative myoclonus have been reported in CJD. Some cases did not show a giant SEP or enhanced C reflex [6]. To which extent the observed PSD in CJD patients is relevant to the myoclonus remains unclear. Shibasaki et al. [7] employed a jerk-locked SEP technique and reported that cortical excitability was suppressed between periodic myoclonic jerks, and this suppression was associated with PSD in a CJD patient. Our findings for the current CJD patient provide further insight into this phenomenon and illustrate that the cortical potentials may or may not lock to the myoclonic jerks and vice versa. One of the possible explanations for this finding is that the generators of the myoclonic jerks are located at different cortical and subcortical levels. The fact that some jerks were not associated with any detectable cortical potentials may indicate that their generator was located very far away from the cortex, i.e. subcortically located, and such jerks were unable to be recorded during the current scalp EEG recording; alternatively, these jerks were randomly evoked and not locked to any specific cortical potentials in a fixed time period. In a typical subcortical myoclonus, i.e. reticular reflex or hyperekplexia, the jerks may involve all four limbs and the muscles innervated by the cranial nerve. The jerk duration may range from 50 to 100 ms [8], which is longer than the duration of the type-C jerks observed in the present study. In contrast, the appearance of robust cortical potentials or PSDs was not always restricted to any myoclonic jerks. These findings support the notion that the generalized PSD may have different influences on the motor neuron excitabilities at cortical and subcortical levels. The subcortical involvement in the current patient was also evidenced by the bilaterally decreased striatal uptake (fig. 1c), although this phenomenon may be irrelevant to the generation of jerks or PSDs.

The direct evidence that confirmed the cortical origin of type-A and -B jerks was the presence of a jerk-locked EEG event. The latency between the onset of jerks and the peak of the cortical potentials was ~24 ms, which is close to the physiological range of the SEP. In the cases of jerk-locked EEG events, we did not find giant SEPs in the scalp recordings. Giant SEPs were classically found in patients with cortical myoclonus [9–11]. However, giant SEP was not seen in all patients with cortical myoclonus. This finding implied that either these patients have no abnormality in the cortical processing of sensory inputs, or the dysfunction is not sufficient to be detected by classical SEP studies [12]. For example, there were usually no giant SEPs in corticobasal degeneration patients with cortical myoclonus. This may result from cortical lesions with functional loss in the corresponding area [13, 14] and this could also be the reason in the current CJD patient who showed a lateralizing cortical involvement in DWI (fig. 1b).

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References


