Coexistence of two distinct benign EEG variants in the same subject

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ABSTRACT – Rhythmic temporal theta bursts of drowsiness (RTTD), also known as “psychomotor variant”, and subclinical rhythmic EEG discharge of adults (SREDA) are two EEG patterns of uncertain significance that occur without any correlation with epilepsy. Each of these patterns has been described to occur alone and in the literature there are no previous reports of co-occurrence of the two distinct benign EEG variants in the same patient. We describe the coexistence of RTTD and SREDA in EEG recordings from the same subject. Although the coexistence of two distinct EEG variants in the same patient is a rarity, these patterns are not so infrequently encountered when present alone and should thus be promptly recognised in order to avoid misdiagnosis of epilepsy due to an over-interpretation of normal sharp patterns.

Key words: benign EEG variants, rhythmic temporal theta bursts of drowsiness, subclinical rhythmic EEG discharge of adults (SREDA)

Rhythmic temporal theta bursts of drowsiness (RTTD), also known as “psychomotor variant” or rhythmic midtemporal discharges, and subclinical rhythmic EEG discharge of adults (SREDA) are two EEG patterns of uncertain significance or “benign EEG variants” that occur without any correlation with epilepsy (Fisch, 1999; Mushtaq and Van Cott, 2005; Niedermeyer, 2005; Blume et al., 2011). They are similar to each other in that they resemble epileptiform activity and are therefore likely to be misinterpreted as true epileptiform patterns. As a consequence, it is important to properly recognise these variants in order to avoid misdiagnosis of epilepsy due to errors in EEG interpretation. Each of these patterns has been described to occur alone and in the literature there are no previous reports of co-occurrence of the two distinct benign EEG variants in the same patient. Here, we describe the coexistence of RTTD and SREDA in EEG recordings from the same subject.
Case report and description of EEG recordings

A 43-year-old woman was referred to our epilepsy centre for a diagnostic evaluation. Her medical history was unremarkable and neurological examination was normal. Some months before, she was referred to a neurologist because of some episodes of excessive daytime sleepiness occurring during a period of forced sleep deprivation. A standard and sleep EEG recording were performed and reported to be abnormal due to the presence of epileptiform left fronto-temporal discharges. Brain magnetic resonance imaging was completely normal. Despite the history, a diagnosis of probable epilepsy was made and an antiepileptic treatment with levetiracetam started.

In our EEG laboratory, the patient underwent an EEG recording which showed brief bursts of sharply contoured rhythmic and monomorphic waves at about 4-5 Hz, occurring prevalently in the left mid-anterior temporal region, and sometimes independently over contralateral homologous regions. These discharges lasted only a few seconds (mean: 3-4 seconds) (figure 1A and B), although sometimes in runs of longer duration (figure 1C), and occurred mostly during drowsiness without any variation in morphology, frequency or spatial distribution. They were identified as RTTD because they fulfilled the classic diagnostic criteria for this EEG variant (Westmoreland and Klass, 1990; Fisch, 1999; Mushtaq and Van Cott, 2005; Niedermeyer, 2005; Blume et al., 2011). The same EEG pattern was recorded also in two subsequent standard and two sleep EEG recordings.

During a sleep EEG, in addition to RTTD, a completely different pattern was observed. This pattern consisted of an abrupt onset of monophasic, sharply-contoured delta (2-3 Hz) waveforms, prevalent in amplitude over parietal regions, which gradually evolved into a sustained rhythmic sinusoidal theta pattern (5-6 Hz). Amplitude was maximal from onset and remained constant. The average duration of the discharges was 20 seconds. The onset of this pattern was abrupt without postictal slowing or changes in EEG background activity. During the sleep EEG recording, this pattern occurred three times, twice during drowsiness (figure 2A and B) and once following a K complex with spindle (stage II NREM sleep) (figure 2C). This pattern, which had no evolution in frequency, morphology or distribution, was identified as SREDA since it fulfilled the classic diagnostic criteria for this pattern (Westmoreland and Klass, 1990; Fisch, 1999; Mushtaq and Van Cott et al., 2005; Niedermeyer, 2005; Blume et al., 2011).

Since the subject never had a seizure or any other episodes suggestive of an epileptic nature, and because several EEG recordings never showed true epileptiform abnormalities, the diagnosis of epilepsy was reconsidered and questioned and levetiracetam was discontinued. At the most recent clinical visit, after 15 months follow-up, the woman reported that her daytime sleepiness ameliorated after levetiracetam withdrawal and normalisation of sleep-wake rhythm. The EEG recording showed exactly the same features as before.

Discussion

The co-occurrence of two normal EEG variants in the same subject has not previously been described in the literature. SREDA is a rare EEG pattern with a prevalence of 1 per 2500 recordings (Westmoreland and Klass, 1981), whereas RTTD is a more common EEG variant occurring in 0.1 to 2% of selected normal adults (Maulsby, 1979). Considering these epidemiological data, the coexistence of two distinct pseudoepilepticiform variants may be indeed considered a rarity. In view of the fact that this appears to be the first report of the co-occurrence of these two benign EEG variants, the possibility of a causal association might be considered. Using Laplacian montages, O’Brien et al. (1998) demonstrated that the site of the SREDA activity is maximal in the parietal region or parieto-centrottemporal regions, whereas for non-SREDA discharges, including RTTD, it is maximal in the temporal or fronto-temporal regions. A subsequent source localisation study performed in a single patient (Zumsteg et al., 2006) revealed a posterior hemispheric source localisation maximal in the parietal cortex bilaterally, in large part overlying the anatomical distribution of the vascular watershed areas. On the other hand, a magnetoencephalographic source modelling study (Lin et al., 2003) indicated that the source of RTTD activity is located in the fissural cortex of the posterior inferior temporal region. As a consequence, is it reasonable to hypothesize that in our subject SREDA and RTTD originate from two different localised cortical generators.

The reported case is interesting not only for its EEG features, but also for its methodological implications concerning the reading and interpretation of the EEG, and illustrates the serious problem of over-interpretation of normal EEG patterns resulting in misdiagnoses of seizures. Prior to our investigation, misinterpretation of an otherwise normal EEG variant, despite the fact that the subject never had a seizure, led to a diagnosis of probable epilepsy. Diagnosing epilepsy without a consistent clinical history was indeed a mistake, however, the lack of recognition of a pseudoepilepticiform benign EEG pattern was also a serious mistake which led to a
Figure 1. Examples of rhythmic temporal theta bursts of drowsiness lasting a few seconds (A and B) or occurring as runs of longer duration (C). Speed: 20 sec/page; sensitivity: 7 µV/mm; HFF: 50 Hz; TC: 0.3 sec.
Figure 2. Subclinical rhythmic EEG discharge of adults occurring during drowsiness (A continuing into B) and stage II NREM sleep (C). Note the absence of changes in ECG rhythm. Speed: 30 sec/page; sensitivity: 7 uV/mm; HFF: 50 Hz; TC: 0.3 sec.
Benign EEG variants in the same subject


Brigo F. We should not treat the EEG, but we should read it blind to the patient’s history. Epilepsy Behav 2011a;20:146.


