INTRODUCTION

Several disorders, systemic as well as those of the central nervous system, are known for exhibiting seasonal fluctuations with higher incidence or more severe symptoms during winter as compared to summer. This feature is usually regarded as a sign of a disorder being vitamin D dependent as circannual fluctuations in vitamin D levels follow that of sunshine hours and solar UV-B radiation [1]. Systemic diseases most well known for seasonal variations include acute myocardial infarction [2], infectious diseases [3] and cancer [4]. Several neurological disorders such as migraine [5], stroke [6] as well as schizophrenia [7] have also been reported to fluctuate seasonally. Also epilepsy has been shown to exhibit some seasonal features: epileptic births [8-10], febrile seizures [11], onset of epileptic spasms [12] as well as photoparoxysmal discharges [13] have been shown to occur more frequently during winter months. Only one study has assessed the seasonal distribution of the epileptic seizures themselves [14]. This study was based on seizures that were logged at an epilepsy inpatient ward throughout a year and indicated higher seizure frequency during winter as compared to summer months and intermediate frequencies during spring and fall. Here we carried out a study to assess possible seasonal distribution of seizures based on individual seizure diaries from epilepsy patients.

MATERIALS AND METHODS

Seizure diaries

Seizure diaries were obtained from epilepsy patients followed by the epileptologists of the Epilepsy Center in the National Institute of Neuroscience, Budapest. Paper-based seizure diaries were photocopied. A few patients have kept seizure diaries digitally. Patients were asked to assess the reliability of their seizure diary. If the patient assessed his or her diary not to be reliable for any reason we did not include it into the analysis. Patient explanations for non-reliability typically included non-adherence to strictly record seizures or being not aware of some seizures for example those occurring during the night. We also excluded diaries from patients with a history of psychogenic non-epileptic seizures. Those diaries with less than twelve consecutive months were also excluded.

Patients

Seizure diaries included in the study originated from 62 patients. These patients were between the ages of 19 and 61 years (mean±SD: 37.3±11.9) at the time of the study. Age at epilepsy onset was between 0 and 34 years (mean±SD: 11.1±8.7). The study population was heterogeneous with regard to epilepsy syndromes: 59 patients had localization-related epilepsy, two had Lennox-Gastaut syndrome and one had idiopathic generalized epilepsy. All patients were taking antiepileptic drugs. No patient received vitamin D supplementation. The study was approved by the local ethical committee and patients gave written informed consent to use their data for scientific analysis.

Seizure count and statistical analysis

Seizures were counted for each calendar month. If a patient had more than one seizure type the different types were
summed. Epileptic auras were not encountered as seizures. In those patients whose seizure diary spanned more than twelve months the respective months were averaged. Seizure counts were normalized to a 30-day length. In a next step, for handling the substantial difference in seizure numbers across patients, seizure counts were transformed and expressed as percentages in each patient such that the sum of the 12 monthly percentages equals 100%. For statistical analysis these monthly percentages were compared. First repeated measure one-way ANOVA with twelve levels was performed to assess main effect. Then paired t-tests were used to compare months.

RESULTS AND DISCUSSION

Altogether 3011 months and 45,833 seizures originating from 62 patients were included in the analysis. Individual seizure diaries on average contained 49 months (SD: 51, range: 12-276 months). At the group level average seizure counts peaked in January (9.7%) and decreased until August the month representing a nadir in seizure counts (7.4%). Thereafter monthly seizure counts progressively increased until December (9.6%). Repeated measure one-way ANOVA with a Greenhouse-Geisser correction indicated a significant within-subject main effect (F (7.79, 475.17) = 2.22, p=0.026). Post-hoc comparison of monthly values of August with those of the other months revealed significant difference for January (p=0.001), November (p=0.03) and December (p=0.003). The difference between the percentage averages in August and January was 31.1%.

The seizure numbers showed a significant seasonal pattern for epileptic seizures with a peak in January and a nadir in August. This finding is in line with a previous study that found highest seizure numbers during the three winter months and lowest during the three summer months when retrospectively analyzing seizures logged at an epilepsy inpatient ward throughout a calendar year [14]. In that study the seasonal pattern hold true for generalized, complex partial as well as simple partial seizures but not for unclassified (presumably non-epileptic) seizures. Some other epilepsy-related phenomena such as epileptic birth rates [8-10] and photoparoxysmal discharges [13] and interictal epileptic discharges [15] were also shown to exhibit seasonal pattern. Importantly, Danesi, an author of several early papers on seasonal [13] and geographical [16] aspects of epilepsy, was the first to relate these phenomena to differences in the amount of sunshine and vitamin D levels. In a large cohort study of England and Wales sudden unexpected death in epilepsy also slightly increased during the winter months [17].

There is more direct evidence supporting the anticonvulsive effect of vitamin D coming from our recent study [18]. In that study we showed that administration of vitamin D3 results in seizure reduction with a median of 40%. Another study carried out nearly forty years ago [19] indicated a seizure reduction of ~30% following administration of vitamin D2 but no significant seizure reduction in the placebo group. The anticonvulsive effect of vitamin D was also evidenced in animal studies [20-22]. Some other mechanisms such as those related to temperature, light exposure and associated hormonal fluctuations could also be suggested to explain seasonality. Yet we are not aware of evidence supporting a relationship between these environmental factors and epilepsy with a same direction and size as seen in our study. Limitations of the study include self-maintenance of seizure diaries which may allow for inaccuracies. However for the long-term recording of seizure events there is no other choice. Also the large number of total seizure events and months assessed in the present study likely compensated for this possible weakness.

Seasonality is generally overlooked in the clinical practice of epilepsy. Nevertheless the average difference amounting to 31% between January and August is considerable. We suggest that the seasonal effect should be held in mind when assessing the efficacy of antiepileptic drugs over a time period of months. More importantly, seizures exhibiting seasonal pattern, together with other direct and indirect evidence, strongly support a role for vitamin D deficiency in epilepsy.
CONCLUSIONS

Epileptic seizures peaked in January while a nadir was present in August. Such a seasonal variation can be regarded as an indirect evidence for vitamin D deficiency being involved in the pathophysiology of epilepsy.

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REFERENCES