CLINICAL PROFILE OF PATIENTS WITH EPILEPSY

Dilli Ram Kafle and Krishna Kumar Oli

Abstract

Epilepsy is a common and diverse disorder with many different causes. Outcomes are varied with 60-70% of newly diagnosed people rapidly entering remission after starting treatment and 20-30% developing a drug-resistant epilepsy with consequent clinical and psychosocial distress. It is a Descriptive Cross-sectional study which was conducted in Tribhuvan University Teaching Hospital from January 2013 to January 2014. A total of 150 patients participated in the study. There was statistically significant association between number of seizures before starting medication and the frequency of seizure after starting medication (p<0.001).

Key words: Neurocysticercosis, Epilepsy

Introduction

Epilepsy is a common and diverse disorder with many different causes. Outcomes are varied with 60—70% of newly diagnosed people rapidly entering remission after starting treatment, and 20—30% developing a drug-resistant epilepsy with consequent clinical and psychosocial distress.¹

About one third of patients with a first unprovoked seizure will have further seizures within five years, and about three quarters of those with two or three unprovoked seizures have further seizures within four years.²

Among 50 million people with epilepsy worldwide, 90% of them are found in developing Countries³ and 90% of these patients are not receiving adequate treatment. They could live normal lives if treated. This huge treatment gap may be due to the limited knowledge, poverty, cultural beliefs, stigma, poor health delivery infrastructure like inadequate supplies of antiepileptic drugs, and shortage of trained health care workers. The prevalence and incidence of epilepsy in Asia is similar to the West but reversible etiologies such as head trauma, infections, stroke, obstetric care are probably more important in Asia.⁴ Epilepsy knows no geographic, social, or racial boundaries and occurs in men and women and affects all ages, but is more frequently diagnosed in infancy, adolescence, and old age. Nepal is one of the poorest countries in the world and it is not uncommon to see huge untreated epilepsy patients in our country. The prevalence rate of epilepsy in Nepal is 7.3 per 1000 population with the treatment gap of over 80%.⁵ People with low socioeconomic status mostly living in the are found be rural areas to more affected.⁶Studies have shown that neurocysticercosis and calcified lesions are the commonest radiological findings.^{7, 8} People suffering from epilepsy in our country do not have good quality of life because of their poor epilepsy control.

The present study was undertaken to study clinical profile of patients with epilepsy.

Methodology

It is a Descriptive Cross-sectional study which was conducted in Tribhuvan University Teaching Hospital from January 2013 to January 2014.

Inclusion Criteria

All the patients attending to neurology outpatient department and those patients

admitted to neurology ward and ICU with recurrent seizure.

Exclusion Criteria

Patients presenting with a history of single seizure ,Patients presenting with a history of multiple seizures within 24 hours without past history of seizure and those who refused to participate in the study.

Statistical Analysis

Data were entered on the computer by using the SPSS Statistical Software (Version 16; SPSS; Chicago, IL) and were analyzed on the same software. Wherever applicable, the data were presented using both the tabular method and descriptive statistics. The strength of associations had been estimated by linear regression analysis and t- test which were used as appropriate. P value of less than 0.05 was considered statistically significant.

Result

The demographic profile of the patients and the clinical characteristics of their seizure are presented in the following table 1.

Table	1.	Demographic	profile	of	the
study pop	ula	tion Baseline D	ata (n=1	50)	

study population Daschile Data (n=150)				
Men	76 (50.7%)			
Women	74 (49.3%)			
Age of patients(Year)	30±15.36			
Seizure type				
(Focal+ secondarily	44			
generalized)	106			
Generalized				
Age at first seizure (Year)	47±17			
Frequency of seizure(Per	80.34±162			
Year)				
Duration of seizure before	2.26±3.5 year			
starting treatment(Year)				
Number of seizures before	24±32.92			
starting treatment				
Family history of epilepsy	14 (9.3%)			
History of status epilepticus	24 (15.7%)			
History of febrile	12 (7.8%)			
	· · ·			

36 (24%)
32 (21.3%)
15 (9.8%)
131 (87.3%)
19 (12.7%)
130 (87 %)
20 (13%)
79 (52.7%)
70 (47.3%)
63 (42%)
87 (58%)

A total of 150 patients were included in the study. There were 76 (50.7%) male and 74(49.3%) female patients included in the study.

Table	2:	Age	distribution	of	study
population	on (y	years)			

population (Jearb)	
Age distribution of study	Number
population(years)	
≤20	40(26.7%)
21-40	80(53.3%)
41-60	21(14%)
>60	9(6 %)

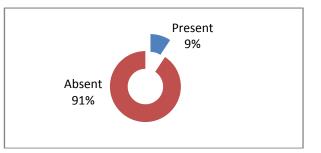


Fig 1. Family history of seizure

14 (9.3%) patients reported having a family history of seizure while 136 (89.7%) patients did not have a family history of epilepsy.

Precipitants	Seizure type	Number of patients
Sleeep	GTCS	6
deprivation	Myoclonic	4
	seizure	4
	Absence	
	seizure	
Alcohol	GTCS	6
intake	Partial seizure	6
Emotional stress	Tonic seizure	2
Fatigue	Atonic seizure	1
Hunger	Myoclonic	1
-	seizure	

Table 3: Precipitants of seizure

30(20%) patients reported having one or more precipitants for their seizure. The precipitants in decreasing order were sleep deprivation, alcohol intake, emotional stress, fatigue and hunger. The presence of precipitants was significantly associated with seizure frequency (p=0.004)

The mean duration of seizure before treatment in years was 2.26 ± 3.5 . The mean number of seizures before treatment was 24.11 ± 32.92 . Mean frequency of seizures after treatment per year was 80.34 ± 162 .

Discussion

In our study, 30(20%) patients reported having one or more precipitants for their seizure. The precipitants in decreasing order were sleep deprivation, Alcohol intake, emotional stress, fatigue and hunger. The presence of precipitant was significantly associated with seizure frequency (p=0.004). E Balamurugan at el also found missing medication, sleep deprivation, fatigue, and emotional stress to be important triggers for seizure onset.¹ 131(87.3%) patients were on monotherapy. 17(11.3%) patients were taking 2 drugs while 2(1.3%) patients were taking 3 drugs. A study conducted in India showed 75.5% of patients maintained on monotherapy while 24.5% percent on polytherapy. ⁹

While 130(87%) patients reported taking their medications regularly, only 20(13%) patients were taking medicine irregularly. Patients who were noncompliant to medication had significantly higher seizure frequency than those patients who were compliant to their medication. (P<0.001)3(2%) patients reported rash with carbamazepine. 2(1.3%) patients developed stevens Johnsen syndrome with phenytoin. 1(0.67%) patient developed hepatitis while on valproic acid. The overall major adverse effect leading to change in current medication was observed in 6 (4%) of patients.

14(9.3%) patients reported having a family history of epilepsy in first degree relative. Family history of epilepsy in Nepalese epileptic patients were found to be higher than that found in India (8.4%) as shown in the study done by Joseph et al.² 116(77.3%)patients had seizure onset before age 30 in our study. In India Joseph et al.² found 68.9% of patient to be having seizure onset before age 25.

History of febrile convulsion was given by 12(7.8%) patients. History of febrile convulsion was statistically associated with seizure frequency (p=0.041).

There was no statistically significant correlation between the duration of epilepsy before starting medication and the frequency of seizure (p=0.9) in our study.

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131(87.3%) patients were on monotherapy. 17(11.3%) patients were taking 2 drugs while 2(1.3%) patients were taking 3 drugs. While 130(87%) patients reported taking their medications regularly, only 20(13%) patients were taking medicine irregularly. Patients who were noncompliant to medication had significantly higher seizure frequency than those patients who were compliant to their medication. (P<0.001)

Nicolas Carpentier et al ¹¹ 11 found in their study that the the rate of nonadherence was 40.9%.Most of the patients in our study were prescribed phenytoin or carbamazepine or valproic acid for their seizures. Some patients were maintained on lamotrigine, oxcarbamazepine or levetiracetam.

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