

Journal of Pharmaceutical Research International

33(53A): 267-273, 2021; Article no.JPRI.78048

ISSN: 2456-9119

(Past name: British Journal of Pharmaceutical Research, Past ISSN: 2231-2919,

NLM ID: 101631759)

Clinical Presentation of the Epileptic Patients at People Medical College Hospital, Nawabshah, Pakistan

Jeando Khan Daidano ^a, Haresh Kumar Makheja ^b, Moti Ram Bhatia ^c, Waseem Raja Memon ^a, Anwar Ali Jamali ^a, Safdar Ali Pervez Tunio ^a and Arslan Ahmer ^{d*}

Authors' contributions

This work was carried out in collaboration among all authors. Author JKD designed the study, performed the statistical analysis, wrote the protocol and wrote the first draft of the manuscript. Authors HKM, MRB, WRM, AAJ, SAPT and AA managed the analyses of the study and managed the literature searches. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/JPRI/2021/v33i53A33659

Editor(s

(1) Dr. Syed A. A. Rizvi, Nova Southeastern University, USA.

(2) Dr. Takashi Ikeno, National Institute of Mental Health, National Center of Neurology and Psychiatry, Japan.

Reviewers:

(1) P. Vadivukkarasi Ramanadin, Sri Vinoba Bhave Civil Hospital, India.

(2) Ufuoma Bigila Shemishere, Federal University Birnin Kebbi, Nigeria.

(3) Sina Dindarian, Urmia University of Medical Sciences, Iran.

Complete Peer review History, details of the editor(s), Reviewers and additional Reviewers are available here: https://www.sdiarticle5.com/review-history/78048

> Received 29 September 2021 Accepted 03 December 2021 Published 04 December 2021

Original Research Article

ABSTRACT

Objective: study will determine clinical presentation, risk, and type of seizures in epileptic patients. **Methodology:** This observational study was conducted in Medicine department People Medical College Hospital (PMCH) Nawabshah from January 2020 to December 2020. 110 patients were included for study after informed permission of the patient or their relative. Male and female were

*Corresponding author: E-mail: arslan.ahmer@gmail.com;

^a Department of Medicine, Peoples University of Medical & Health Sciences for Women Shaheed Benazirabad, Sindh, Pakistan.

^b Department of Psychiatry, Khairpur Medical College (KMC), Khairpur Mirs, Sindh, Pakistan. ^c Department of Psychiatry, Peoples University of Medical & Health Sciences for Women Shaheed Benazirabad, Sindh, Pakistan.

^d Institute of Pharmaceutical Sciences, Peoples University of Medical & Health Sciences for Women Shaheed Benazirabad, Sindh, Pakistan.

selected. Known epileptic patients were included in the study, patients with false seizures were excluded from the study. Statically analysis was done by software SSPS 22 version.

Results: Age ranged 18 to 60 years. All the patients included in the study epilepsy was noted. The mean age of patients was 37.40 SD 8.71 years. The minimum age was 28 years while maximum 60 years. Pearson Chi-Square Value 105.000^a Asymp. Sig. (2-sided) .000, Likelihood Ratio Value 135.012 Asymp. Sig. (2-sided) .000, Linear-by-Linear Association Value 16.297 Asymp. Sig. (2-sided) .000 which were statistically significant.

Conclusion: Epilepsy is treatable common neurological disease in Pakistan. quality of life can be improved by Education of the patients and their relatives, without socioeconomically burden.

Keywords: Seizure; epilepsy; clinical; medicine.

1. INTRODUCTION

In developed countries and developing countries epilepsy is most common neurological disorder. Recurrence of seizure is termed as epilepsy, not all convulsions are due to epilepsy [1]. Excessive discharge of neurons in brain alter neurological function termed as a seizure. Seizure occur when there is imbalance between excitation and inhibition in brain [2]. Numerous causes are included in Epilepsy and brain dysfunction [3]. various causes of epilepsy include genetic predisposition, head injury, brain tumors, stroke and drug or alcohol withdrawal. Seizure can be due to hypoglycemia, fever, meningitis and psychogenic. Seizures due to alcohol withdrawal is not epilepsy, epilepsy is generation of seizures cognitive, neurobiological, social psychological consequences of this condition [4]. Epilepsy most common CNS disorder 50 per 100000 new cases per year of the population. Incidence of epilepsy about 1%, refractory epilepsy in 1/3 and epilepsy from childhood 75% [5]. Worldwide 50 million people are affected by this non communicable disease epilepsy. Epilepsy is leading neurological disease in the world. Every year 61.4 per 100000 incidence of epilepsy is reported in population [6]. 5% prevalence rate of epilepsy reported in Iran [7]. All age groups are affected by epilepsy but common in young children and older age group [8]. Frontal lobe epilepsy is second following temporal lobe epilepsy, this type of epilepsy originate from frontal lobe, occur during wake or sleep [9]. 20-30% patients of focal epilepsy are associated with frontal lobe epilepsy [10]. Epilepsy can be misdiagnosed as sleep disorder, non epileptic seizures and psychiatric disorder classified Seizures as generalized. partial(now focal) and epileptic spasm. Limited part of cerebral hemisphere is involved in focal or partial epilepsy. Bilateral distributed neuronal network is involved in generalized seizure.

Initially seizure can be focal later become generalized [12]. Seizure event is common in 10% population [13]. Seizure control is important for Doctors, nursing staff and patients or their relatives with learning disabilities Communication is important for patient with Doctors, regarding drugs side effects and seizures [15]. Patients with learning disabilities are at risk of uncontrolled seizures and increased mortality rate [16]. More than twenty drugs are in use for the successful treatment of epilepsy. These drugs act by preventing neuronal depolarization, blocking calcium or sodium channels, decrease electrical activity of the brain, potassium enhance channel function. neurotransmitter excitation of glutamate is inhibited [17].

2. METHODOLOGY

study was conducted in Medicine department PMCH Nawabshah from January 2020 to December 2020, 105 patients were included for study after informed permission of the patient or their relative. Male and female were selected. Detailed proforma was used for the study, detailed history, clinical examination of the patient, compulsory investigations of the patient, RBS, Urea, creatinine, LFT, Blood CP, Serum Electrolyte, serum Calcium Level, Urine DR, X-ray Chest, EEG, CT Scan Brain and MRI Brain. All patients with true seizures were included for this study, patients with pseudo seizures and seizures due to metabolic disorders were excluded from the study. Statically analysis was done by software SSPS 22 version.

3. RESULTS

The mean age of patients was 37.40 SD 8.71 years. The minimum age was 28 years while maximum 60 years. As shown in Table 1.

Table 1. Descriptive statistics

| | N | Range Minimu | | Maximum | Mean | | Std. Deviation |
|-------------------------|------------|--------------|-----------|-----------|-----------|---------------|-------------------|
| | Statistic | Statistic | Statistic | Statistic | Statistic | Std. Error | Statistic |
| Age in years Valid N | 105 105 | 32.00 | 28.00 | 60.00 | 37.4000 | .85030 | 8.71294 |

The details of different demographic variables occupation, education, Address, addiction SE like age group, gender, marital status, address, Class and Family History are shown in Table 2.

Table 2. The details of different demographical data

| Demographic var | riables | Frequency | Percent (%) | | |
|-----------------|------------------------|-----------|--------------|--|--|
| Age Group | 20-40 Years Young Age | 78 | 74.3 | | |
| | 41-60 Years Middle Age | 27 | 25.7 | | |
| Gender | Male | 69 | 65.7 | | |
| | Female | 36 | 34.3 | | |
| Marital Status | Married | 90 | 85.7 14.3 | | |
| | Un-Married | 15 | | | |
| Occupation | No Occupation | 24 | 22.9 | | |
| · | House Wife | 27 | 25.7 | | |
| | Manual Worker | 44 | 41.9 | | |
| | Office Worker | 10 | 9.5 | | |
| Education | Educated | 65 | 61.9 | | |
| | Un-Educated | 40 | 38.1 | | |
| Address | Rural | 73 | 69.5 | | |
| | Urban | 32 | 30.5 | | |
| Addiction | No | 80 | 76.2 | | |
| | Yes | 25 | 23.8 | | |
| SE Class | Poor Class | 82 | 78.1 | | |
| | Middle Class | 16 | 15.2 | | |
| | Upper Class | 7 | 6.7 | | |
| Family History | No | 86 | 81.9 | | |
| , , | Yes | 19 | 18.1 | | |
| Total | | 105 | 100.0 | | |

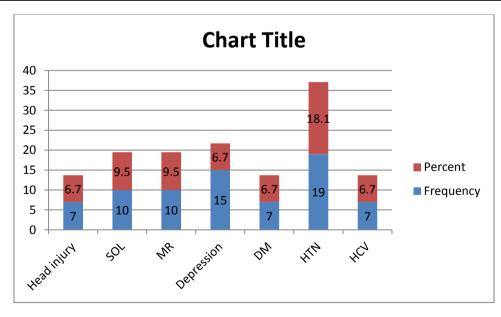


Fig. 1. The comorbidities in epileptic patients like head injury, SOL, MR, depression, DM, HTN and HCV

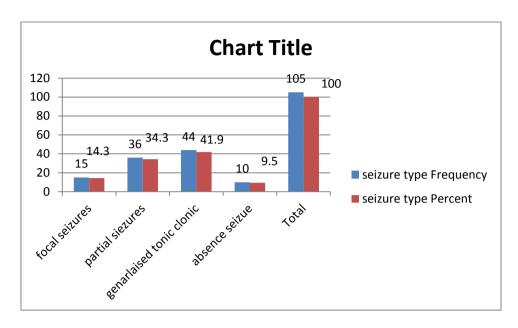


Fig. 2. The different types of seizures noted in epileptic patients

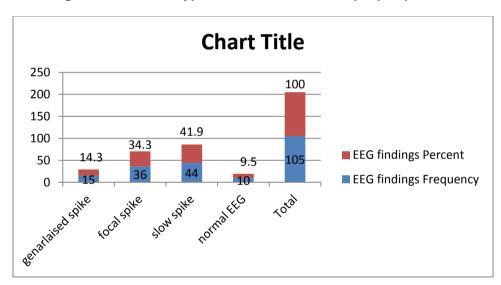


Fig. 3. Different types of EEG findings noted in epileptic patients

Table 3 shows the gender * seizure type Cross tabulation Pearson Chi-Square Value 105.000^a Asymp. Sig. (2-sided) .000, Likelihood Ratio Value 135.012 Asymp. Sig. (2-sided) .000,

Linear-by-Linear Association Value 16.297 Asymp. Sig. (2-sided) .000 which were statistically significant.

Table 3. Statistical results

| Variable | | | Total | | | | |
|----------|--------|------------|----------------|------------------|-------------------|-----------------|--------|
| | | | Focal seizures | Partial seizures | Generalized tonic | Absence seizure | |
| Gender | Male | Count | 15 | 0 | 44 | 10 | 69 |
| | | % of Total | 14.3% | 0.0% | 41.9% | 9.5% | 65.7% |
| | Female | Count | 0 | 36 | 0 | 0 | 36 |
| | | % of Total | 0.0% | 34.3% | 0.0% | 0.0% | 34.3% |
| Total | | Count | 15 | 36 | 44 | 10 | 105 |
| | | % of Total | 14.3% | 34.3% | 41.9% | 9.5% | 100.0% |

| Chi-Square Tests | | | | | |
|---------------------------|----------------------|---------|--|--|--|
| | Value | df | Asymp. Sig. (2-sided) | | |
| Pearson Chi-Square | 105.000 ^a | 3 | .000 | | |
| Likelihood Ratio | 135.012 | 3 | .000 | | |
| Linear-by-Linear | 16.297 | 1 | .000 | | |
| Association | | | | | |
| N of Valid Cases | 105 | | | | |
| a. 1 cells (12.5%) have e | expected count I | ess tha | n 5. The minimum expected count is 3.43. | | |

4. DISCUSSION

Following factors stigmatization, poverty, attitude, lack of adequate knowledge and coping skills are important in epilepsy care. Majority of the patients and relatives lack the knowledge about precipitating factors and the cause of epilepsy mentioned in earlier studies [18]. Knowledge about epilepsy treatment is important, role of supportive care, information about the disease, precipitants of seizures, role of treatment and adverse effects of drugs [19]. Neuroimaging CT Scan and MRI are helpful for detection of demyelinating disorders, atrophic lesion of brain and structural lesions of brain. Neuroimaging is helpful in patients with partial epilepsy abnormal findings were reported, structural lesion were localized by radiological support in symptomatic epilepsy [20]. Role of Doctors, nurses and health assistant is important in the treatment of epilepsy, communication between patient and health care professional is encouraged [21]. Epilepsy not considered as major public health issue because treatment is readily available and cheap [22]. Epilepsy associated with depression. due to biological and sociological factors. Abnormalities in neurotransmitter 5HT Glutamate leads to depression in epilepsy [23]. Better outcome is associated with surgery in epilepsy, when single region of brain responsible for epilepsy [24]. Mesial temporal sclerosis a structural lesion with intractable seizure, surgery is option [25]. Seizures associated with behavior, memory change, altered responsiveness and posturing in the medial temporal region. Seizures are intractable with co morbidities. Surgical option is considered when two drugs fail. For the pathophysiology of temporal lobe epilepsy investigations are performed, genetic factors are important in temporal lobe epilepsy [26]. Stimulation of nerve where surgical resection is contraindicated, Vagus nerve is safe for stimulation with low complications, like vocal cord paralysis, hematoma and infection [27], Epilepsy associated with other comorbid conditions, these are psychiatric disorders anxiety, depression, learning disabilities, autism, intellectual disability and attention deficit hyperactivity disorder. These comorbid conditions considered to be integral

part of the disease, previously these comorbid conditions considered to be due to side effects of antiepileptic drugs or uncontrolled seizures [28]. Epileptic circuits limbic and hippocampal dysfunction associated with common psychiatric comorbidity depression. Depression is more common in patients with history of epilepsy and epileptic patients develop depression. About 30% patients of epilepsy have depression and 10% patients have bipolar disorder [29].

5. CONCLUSION

Epilepsy common neurological disease in Pakistan, is treatable with cost effective drugs and minimal side effects. Epilepsy is major health problem in our country, long term treatment is needed in majority of the patients. Awareness about seizure, precautionary measures are compulsory. Precautions from fire, water and sudden fall during seizure. Improvement in dietary habits, sleep and early treatment of any infection. Education of the patients and their relatives' quality of life can be improved without socioeconomically burden. Early treatment and education about disease, stress can be reduced with improved quality of life.

CONSENT

As per international standard or university standard, patient's written consent has been collected and preserved by the author(s).

ETHICAL APPROVAL

As per international standard or university standard written ethical approval has been collected and preserved by the author(s).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

 Berg AT, Bercovic SF, Brodie MJ, et al. revised terminology and concepts for

- organization of seizures and epilepsies: report of ILAE Commission on classification and terminology. 2005-2009. Epilepsia. 2010;51: 676-685.
- Stafstorm CE. Pathophysiological mechanism of seizures and epilepsy: A Primer in Epilepsy: Mechanism, models and translational perspectives (ed Rho JM, Sankar R, Stasform CE). 2010;3-19. CRC, Boca Raton, FL.
- Shorvon SD, Andermann F, Gueirrine R (ed) 2011. The causes of Epilepsy. Cambridge University Press, Cambridge.
- 4. Sander JW, The natural history of epilepsy in the era of new antiepileptic drugs and surgical treatment. Epilepsia. 2003; 44(Suppl 1):17-20.
- 5. Hauser W, Hersdorrfer D. Epilepsy: Frequency, causes and consequences. Demos, New York; 1990.
- Fiest KM, Sauro KM, Wiebe S, Patten SB, Kwon C-S, Dykeman J, et al. prevalence and incidence of epilepsy: a systemic review and meta-analysis of International studies. Neurology. 2017;88(3): 296-303.
- 7. Sayehmiri K, Tavan H, Sayhimiri F, Mohammadi I, Carson KV. Prevalence of epilepsy in Iran: a meta-analysis and systemic review. Iran J Child Neurol. 2014;8(4)9.
- 8. m/L/2 Kenneth W, Lindsay, Ian Bone. Neurology and Neurosurgery Illusterated; 4th edition. 2004;90-103.
- Haut S, Frontal lobe epilepsy. Medscope; 2009.
- 10. Doelken MT, Mennecke A, Huppertz HJ. Multimodality approach in cryptogenic epilepsy with focus on morphometric 3 TMRI. J Neuroradiol. 2012;39(2):87-96.
- Mayoclinic staff. Frontal lobe seizure. 2008.
 Available:http://www.mayoclinic.org/diseas es conditions/frontal-lobeseizures/symptoms-causes/dxc-20246880 [Acessed 20 Nov 2016].
- 12. Berg AT, Millichap JJ. The 2010 revised classification of seizures and epilepsy. Contnuum. (Minneapp Minn). 2013;19: 571-597.
- Sander JW. The natural history of epilepsy in the era of new antiepileptic drugs and surgical treatment. Epilepsia. 2003; 44(Suppl1) 17-20.
- 14. Haut S. Frontal lobe epilepsy. Medscope; 2009
- 15. Doelken MT, Mennecke A, Huppertz HJ. Multimodality approach in cryptogenic

- epilepsy with focus on morphometric3TMRI. J Neuroradiol. 2012; 39(2): 87-96.
- 16. Selemon L, Goldman-Rakic P. Longitudinal topography and interdigitation of corticospinal projections in the rhesus monkey. J Neurosci. 1985;5:776-94.
- Bui A, Kim H, Moroso M, Soltez I. Microcircuits in epilepsy: Heterogeneity and hub cells in network synchronization. Cold Spring Harb Perspect Med. 2015;10. 1101/cshperspect a022855.
- Ogunrin OA, Adeyekun A, Adudu P. Etiologies of epilepsy and health-seeking itinerary of patients with epilepsy in a resource poor setting: analysis of 243 Nigerian Africans. Seizure. 2013;22(7): 572-6.
- Nwani PO, Arinzichi EO, Asomugha AL, Enwereji KO, Nwosu MC, Ogunniyi AO. Illness concept among people with epilepsy and their caregivers and preferred treatment methods in a suburban community in South East Nigeria. West Afr J Med. 2013;32(1): 26-30.
- 20. manz. Lakh.15 Obajmi MO, Fotunde OJ, Oganseyinde. Computed Tomography in child hood seizure disorder in lobdor West Africa. J Med. 2004;232: 16772.
- 21. Rodgers J, Namaganda S. Making information easier for people with learning disalities. Br J Learn Disabil. 2005;33(2): 52-8.
- 22. Bertolote JM. Epilepsy as a public health problem. Role of the World Health Orgnization and the cooperation between WHO and non Governmental organizations. Top Geogr Med. 1994;46 (3Suppl): S 28-30.
- 23. Lothe A, Didilot A, Hammers A, et al. Comobidity between temporal lobe epilepsy and depression. a [18F] MPPF PET Study Brain. 2008;131:2765-2782.
- 24. De Tisi J, Bell GS, Peacock JL, et al. The long term outcome of adult epilepsy surgery, pattern of seizure remission, a relapse: a cohort study Lancet. 2011;378: 1388-1395.
- 25. Thom A, Mathern JW, Cross JH, Bertram EH. Mesial temporal lobe epilepsy: How do we improve surgical outcome? Ann Neurol. 2010;68:424-434.
- 26. Liu Z, Mikati M, Holmes GL. Mesial temporal sclerosis: Pathogenesis and significance. Peditr Neurol. 1995;12:5-16.
- 27. Revesez D, Rydenhag B, Ben-Menacham E. Complications and safety of vagus

- nerve stimulation: 25 years experience at a single center. J Neurosurg Pediatr. 2016; 18: 97-104.
- 28. Brookes-Kayal AR, Bath KG, Berg AT, Galanopoulou AS, Holemes GL, Gensen FE, Kanner AM, O' Brien TJ, Whittemore VH, Vinawer MR, et al. Issues related to
- symptomatic and disease-modifying treatments affecting cognitive and neuropsychiatric comobidities of epilepsy. Epilepsia. 2013;54: 44-60.
- 29. Kanner AM. The treatment of depressive disorders in epilepsy: What all Neurologists should know. Epilepsia. 2013;54:3-12.

© 2021 Daidano et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
https://www.sdiarticle5.com/review-history/78048