

Original Research Article

COMPARATIVE STUDY OF NEUROBEHAVIORAL PROFILE AND SLEEP DISORDER IN EPILEPTIC CHILDREN

Kapil Bainade¹, Pallavi Gahlowt², Veeranna Kotrashetti³, Arnab Bhattacharjee⁴

 Received
 : 21/06/2025

 Received in revised form
 : 04/08/2025

 Accepted
 : 25/08/2025

Corresponding Author:

Dr. Arnab Bhattacharjee,

Junior Resident, Department of Paediatrics, Dr D. Y. Patil Medical College, Nerul Navi Mumbai, India. Email: arnabbhattacharjee2407@gmail.com

DOI: 10.70034/ijmedph.2025.3.472

Source of Support: Nil, Conflict of Interest: None declared

Int J Med Pub Health

2025; 15 (3); 2572-2577

ABSTRACT

Background: Epilepsy is a common pediatric neurological disorder with a global prevalence of 0.5–1%, significantly higher in low- and middle-income countries. Beyond seizures, epileptic children frequently exhibit neurobehavioral challenges and sleep disturbances, often underrecognized in clinical care. This study aimed to compare the neurobehavioral profiles and sleep patterns in epileptic children versus non-epileptic controls to emphasize the need for holistic epilepsy management.

Materials and Methods: A hospital-based case-control study was conducted over one year in a tertiary care center in Navi Mumbai. Fifty children aged 1–12 years were enrolled—25 with epilepsy (Group A) and 25 with non-neurological illnesses (Group B). Behavioral assessment was conducted using the Child Behavior Checklist (CBCL), while sleep disturbances were screened using the BEARS questionnaire. Statistical analysis was performed using t-tests and Chi-square tests with p<0.05 considered significant.

Results: Epileptic children had significantly higher CBCL scores across all domains, including Anxious/Depressed (64.4 vs. 51.16), Withdrawn/Depressed (67.88 vs. 53.52), and Attention Problems (65.08 vs. 52.04) with p<0.001. Sixty percent of epileptic children had clinically significant behavioral issues versus 0% in controls (p<0.001). Sleep disturbances were more prevalent in the epilepsy group (68%) compared to controls (44%), though not statistically significant (p=0.087). However, all epileptic children with behavioral issues had co-existing sleep disturbances (p<0.001).

Conclusion: Children with epilepsy exhibit significantly more behavioral problems and sleep disturbances than non-epileptic peers. These findings highlight the importance of integrating behavioral and sleep assessments into routine epilepsy care for better developmental outcomes.

Keywords: Epilepsy, Child Behavior Checklist, Sleep Disorders, BEARS Questionnaire, Neurobehavioral profile, Neurodevelopment.

INTRODUCTION

Epilepsy is one of the most prevalent chronic neurological disorders in children, affecting approximately 0.5–1% of the global pediatric population, with over 11 million children under the age of 15 currently living with the condition worldwide. The burden is disproportionately higher in low- and middle-income countries, including India, where prevalence rates among children vary

between 2.02 to 22.2 per 1,000 depending on region and healthcare access. [1,2]

Defined by the International League Against Epilepsy (ILAE), epilepsy is characterized by recurrent unprovoked seizures or the diagnosis of an epilepsy syndrome. [3] While seizures are the hallmark of the disease, epilepsy also imposes significant neurodevelopmental, cognitive, and psychosocial burdens, often exacerbated by antiepileptic drugs

¹Associate Professor, Department of Paediatrics, Dr D. Y. Patil Medical College, Nerul Navi Mumbai, India.

²Professor, Department of Community Medicine, Dr D. Y. Patil Medical College, Nerul Navi Mumbai, India.

³Professor and Head of Department, Department of Paediatrics, Dr D. Y. Patil Medical College, Nerul Navi Mumbai, India.

⁴Junior Resident, Department of Paediatrics, Dr D. Y. Patil Medical College, Nerul Navi Mumbai, India.

(AEDs), socioeconomic status, and family dynamics.^[4,5]

Children with epilepsy frequently demonstrate behavioral issues such as aggression, hyperactivity, anxiety, and attention deficits, with studies reporting psychological disorders in up to 58% of such children, particularly when brain abnormalities are present. [6] Cognitive impairments are also common, with reduced memory, language function, and school attendance cited in numerous studies. [7,8] These issues are not isolated to seizure activity but may persist interictally and worsen over time, especially in those with early-onset or refractory epilepsy. [9]

Sleep disturbances form a critical underrecognized comorbidity in pediatric epilepsy. Sleep and seizures have a bidirectional relationship disrupted sleep can precipitate seizures, while seizures (especially nocturnal) can impair sleep architecture. Epileptiform discharges such as centrotemporal spikes and generalized spike-wave discharges often occur or intensify during non-REM sleep, affecting memory consolidation and cognitive performance.[10,11] Syndromes such as Electrical Status Epilepticus during Sleep (ESES) exemplify the severity of this interaction. Tools like the BEARS questionnaire help screen for common sleep disorders including insomnia, night awakenings, and sleep apnea, all of which are more prevalent in children with epilepsy than their peers.^[12]

The role of AEDs in modulating both behavior and sleep are significant. While newer AEDs like lamotrigine and levetiracetam have improved side effect profiles, many traditional agents are linked to sedation, cognitive dulling, and mood changes. Polytherapy, in particular, exacerbates behavioral and cognitive disturbances. Studies have shown that children on multiple AEDs are more prone to attention deficits, irritability, and academic difficulties. [14]

Despite these profound implications, routine clinical care often focuses solely on seizure control, overlooking associated behavioral and sleep-related comorbidities. This narrow focus delays early diagnosis and intervention, reducing the overall quality of life and developmental outcomes in these children. Early recognition through standardized tools like the Child Behavior Checklist (CBCL) and BEARS questionnaire can aid in identifying at-risk individuals and tailoring holistic interventions.^[15,16] This study aims to compare the neurobehavioral profiles and prevalence of sleep disorders in epileptic children and their age-matched non-epileptic counterparts. By identifying significant differences, the study emphasizes the need for integrative pediatric epilepsy care that addresses not only seizure control but also the broader spectrum of developmental and behavioral health.

MATERIALS AND METHODS

A case-control study and was conducted over a period of one year, from January 2024 to January 2025, in the Department of Paediatrics at a tertiary care centre at Navi Mumbai. A total of 50 children, aged between 1 to 12 years, were enrolled in the study. Ethical clearance was obtained from the Institutional Ethics Committee prior to the commencement of the study. Informed written consent was collected from parents or legal guardians of all participants. All data collected during the study was handled confidentially, and privacy was ensured at every stage of the research process.

The participants were divided into two groups: Group A (cases) comprised 25 children diagnosed with epilepsy based on the International League Against Epilepsy (ILAE) criteria, while Group B (controls) included 25 children admitted for non-neurological illnesses such as acute gastroenteritis, viral fever, or respiratory tract infections. A simple random sampling technique was employed to select participants who fulfilled the inclusion and exclusion criteria.

Children were included in the study if they were aged 1 to 12 years, of either sex, and if informed consent was obtained from their parents or legal guardians. Children with known neurodevelopmental disorders such as autism spectrum disorder, cerebral palsy, intellectual disability, and those with febrile or metabolic seizures or congenital structural abnormalities were excluded from both groups to eliminate confounding variables.

The sample size was calculated assuming an epilepsy prevalence of 1.6% based on a study by Santhosh et al,^[17] at a confidence interval of 95% and an absolute precision of 5%, the minimum required sample size was 24.18. This was rounded off to 25 participants per group.

Data was collected through structured interviews and clinical evaluations conducted during hospital admission. Each participant underwent a thorough history-taking and physical examination. Sociodemographic details, clinical diagnosis, antiepileptic medication use, EEG and imaging findings were recorded where applicable.

To assess the neurobehavioral profile, the Child Behavior Checklist (CBCL) was administered to parents or guardians. The CBCL is a standardized, validated tool that evaluates emotional and behavioral problems in children and provides both cumulative scores and T-scores for comparison with normative data. The internalizing and externalizing behavioral domains were analyzed.

For sleep disorder assessment, the BEARS questionnaire was used. This is a brief screening tool that evaluates five domains of pediatric sleep: Bedtime problems, Excessive daytime sleepiness, Awakenings during the night, Regularity of sleep, and Snoring. A positive response in any of these

domains was indicative of a potential sleep disturbance.

The collected data was entered into Microsoft Excel and analyzed using IBM SPSS Statistics for Windows, Version 26.0. Quantitative variables were expressed as means and standard deviations, while categorical data was presented in frequencies and percentages. Comparisons between groups were made using the independent t-test for continuous variables and the Chi-square or Fisher's exact test for categorical variables. A p-value of less than 0.05 was considered statistically significant.

RESULTS

The present study was conducted among 50 children, 25 children admitted with epilepsy seizures (study group) and 25 children with non-neurological illnesses (control group) admitted to a tertiary care hospital in Nerul, Navi Mumbai. The mean age of the study participants was 7.26 + 3.08 years. The majority of the children in the study group (52%) and control group (52%) were boys while 48% were females.

Among the cases, 60% had generalised epilepsy, 28% had focal epilepsy, 8% had generalised idiopathic epilepsy while 4% had syndromic epilepsy. 60% of the children were on Levetiracetam, 44% were on Valproate and 4% were on Lamotrigine. (Figure 1)

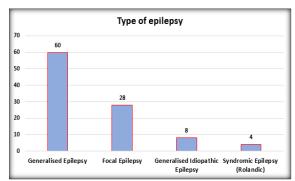


Figure 1: Distribution of cases based on Type of epilepsy

While a subset of children with epilepsy had identifiable abnormalities such as gliosis or cortical dysplasia, the majority showed normal imaging

underlining that the neurobehavioral problems in epilepsy may occur even in the absence of visible anatomical lesions, pointing to functional disruptions as contributing factors. (Figure 2)

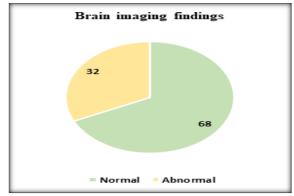


Figure 2: Distribution of cases based on Brain imaging findings

Among the CBCL subscales used to assess the neurobehavioral problems, the maximum clinical significance was observed for Withdrawn/Depressed (56%) followed by Anxious/Depressed (52%) and Thought Problems (52%), Somatic Complaints (44%) and Rule-Breaking Behavior (44%), Social Problems (40%) and Rule-Breaking Behavior (40%). The least clinical significance was observed for Aggressive Behavior (4%).

Table 1 compares the mean CBCL subscale scores between epileptic and non-epileptic children. The study group (epileptic children) had significantly higher scores across all domains including Anxious/Depressed (64.4 \pm 10.6 vs. 51.16 \pm 2.57), Withdrawn/Depressed (67.88 \pm 12.75 vs. 53.52 \pm 5.42), Somatic Complaints (66.04 ± 11.45 vs. 55.92 \pm 6.33), Social Problems (64.76 \pm 10.22 vs. 54.40 \pm 5.78), Thought Problems (65.72 \pm 11.18 vs. 54.92 \pm 5.58), Attention Problems (65.08 \pm 10.53 vs. 52.04 \pm 3.97), Rule-Breaking Behavior (63.72 \pm 10.14 vs. 52.44 ± 4.65), and Aggressive Behavior (58.12 ± 7.59 vs. 52.08 ± 3.13). All differences were statistically significant (p < 0.001), indicating that epileptic children demonstrate significantly greater emotional and behavioral disturbances than their non-epileptic peers. (Table 1)

Table 1: Cl	BCL subscale	score among	cases and controls
-------------	--------------	-------------	--------------------

CBCL subscale score	Group	Mean	Std. Deviation	t-value	p-value
Amvious/Dommossod	Cases	64.400	10.661	(025	<0.001*
Anxious/Depressed	Controls	51.160	2.576	6.035	
Withdrawn/Depressed	Cases	67.880	12.758	5.179	0.001
withdrawn/Depressed	Controls	53.520	5.424	5.1/9	
Somatic Complaints	Cases	66.040	11.454	3.865	0.001
Somatic Complaints	Controls	55.920	6.337		
Social Problems	Cases	64.760	10.223	4.409	0.001
Social Problems	Controls	54.400	5.787		
Thought Duckland	Cases	65.720	11.182	4.321	<0.001*
Thought Problems	Controls	54.920	5.582		
A., .: D. 11	Cases	65.080	10.531	5.700	<0.001÷
Attention Problems	Controls	52.040	3.973	5.792	<0.001*
Rule-Breaking Behavior	Cases	63.720	10.146	5.052	<0.001*

	Controls	52.440	4.655		
Aggressive Behavior	Cases	58.120	7.590	2 (79	<0.001÷
	Controls	52.080	3.134	3.678	<0.001*

Table 2 provides the overall behavioral burden as assessed by cumulative and T-scores. The mean total cumulative CBCL score was markedly higher in epileptic children (54.04 \pm 37.90) compared to controls (11.96 \pm 11.96), with a significant mean difference of 42.08 (p < 0.001). Similarly, the mean

total CBCL T-score was significantly higher in the case group (58.88 ± 19.41) than in controls (39.92 ± 12.61), with a mean difference of 18.96 (p = 0.026). These results reflect a higher degree of clinical emotional-behavioral difficulties in children with epilepsy. (Table 2)

Table 2: Total CBCL Cumulative Score among cased and controls

CBCL	Score	Mean	SD	Mean difference	t-value	p-value
Total CBCL	Cases	54.040	37.905		5.293	<0.001*
Cumulative Score	Controls	11.960	11.966	42.080		
Total CBCL T	Cases	58.880	19.412	18.960	4.095	0.026*
Score	Controls	39.920	12.612			0.026"
	Independent t-test					

Table 3 shows that among epileptic children, 60% (15 out of 25) had clinically significant behavioral problems, 4% (1 child) were borderline, and 36% (9 children) were normal. In contrast, 96% (24 out of 25) of non-epileptic children were normal, with only 4% (1 child) being borderline and none showing

clinically significant issues. This difference was statistically significant (Chi-square = 24.73, p < 0.001), indicating a higher prevalence of behavioral disturbances in epileptic children compared to controls. (Table 3)

Table 3: Overall Interpretation based on the CBCL score

	Over	Overall CBCL Interpretation			
Group	Normal	Borderline	Clinically Significant	Chi-square value	p-value
Case	9 (36)	1 (4)	15 (60)	24.730	<0.001*
Control	24 (96)	1 (4)	0 (0)	24./30	~0.001 [™]

68% of the cases and 44% of the controls had sleep disorders in our study as per the the BEARS questionnaire. Sleep disorder among both groups were not found to have a statistical significance with a chi-square value of 2.922 and a p-value of 0.087. Table 4 shows a significant association between neurobehavioral problems and sleep disorders in epileptic children. All children (100%) with clinically significant CBCL scores had sleep disorders, while

only 22.2% of those with normal behavior had sleep issues. This association was statistically significant (Chi-square = 18.309, p < 0.001). In contrast, among non-epileptic children, 41.7% of those with normal behavior had sleep disorders, but this was not statistically significant (Chi-square = 1.326, p = 0.440). Thus, sleep disturbances were closely linked to behavioral problems in epileptic children, but not in controls. (Table 4)

Table 4: Association between Neurobehavioral profile and sleep disorder

C	Overall CBCL	Sleep Disorder		Chi-square	
Group	Interpretation	Yes	No	value	p-value
	Normal	2 (22.2)	7 (77.8)		<0.001*
Case	Borderline	0 (0)	1 (100)	18.309	
	Clinically Significant	15 (100)	0 (0)		
	Normal	10 (41.7)	14 (58.3)		
Control	Borderline	1 (100)	0 (0)	1.326	0.440
	Clinically Significant	0 (0)	0 (0)		

DISCUSSION

This study aimed to assess and compare the neurobehavioral profiles and sleep disturbances in epileptic children and non-epileptic controls. The results revealed a significantly higher prevalence of behavioral issues and sleep disturbances among epileptic children, aligning with existing literature on the multidimensional impact of epilepsy on child development.

Our findings showed that epileptic children had significantly elevated scores in all CBCL subscales, including\ Anxious/Depressed, Withdrawn/Depressed, Somatic Complaints, Social Problems, Thought Problems, Attention Problems, Rule-Breaking, and Aggressive Behavior domains. These results are consistent with the population-based study by Reilly et al., which found that children with epilepsy are at a significantly higher risk of developing emotional and behavioral comorbidities

compared to their peers, with internalizing problems being most common (e.g., anxiety, depression).^[5] Similarly, Rodenburg et al. conducted a meta-analysis confirming that children with epilepsy experience a higher rate of psychopathology than the general pediatric population.^[13]

The current study also demonstrated that 60% of epileptic children had clinically significant behavioral problems, in contrast to none in the control group. This supports findings from the study by Srinivas and Shah, who reported high psychiatric comorbidity rates in pediatric epilepsy, emphasizing the need for regular behavioral screening in this group.[14] Karrasch et al., in a longitudinal cohort study, observed that behavioral and cognitive issues may persist or even worsen over time, particularly in children with early-onset or poorly controlled epilepsy.^[6] This highlights that behavioral changes in epilepsy are not only seizure-related but can be influenced by underlying neurological dysfunction and long-term AED use.

A major contributor to behavioral changes in children with epilepsy is the use of antiepileptic drugs (AEDs). In our study, most children were on Levetiracetam and Valproate. Although newer AEDs like Levetiracetam are considered to have better neurobehavioral profiles, several studies have reported irritability, aggression, and mood changes associated with its use. [13] Polytherapy, in particular, increases the risk of behavioral side effects, as supported by Rodenburg et al. and Owens et al., who found that children on multiple AEDs tend to exhibit greater attention deficits, emotional lability, and academic underperformance. [12,13]

Sleep disturbances were also more prevalent among epileptic children (68%) compared to controls (44%). Though the association between epilepsy and sleep disorder was not statistically significant overall, a strong correlation was seen between clinically significant behavioral scores and the presence of sleep disorders in the epilepsy group (p < 0.001). This suggests a strong interdependence between behavioral and sleep issues in children with epilepsy. Nobili et al. emphasized the bidirectional relationship between epilepsy and sleep disturbances, where nocturnal seizures and interictal epileptiform discharges during non-REM sleep disrupt sleep architecture, contributing to cognitive and behavioral deficits. [10]

The role of sleep disturbances in aggravating behavioral problems has also been well established. Owens highlighted that children with epilepsy often present with insomnia, night awakenings, and daytime sleepiness, which further impair mood and attention regulation. [12] The BEARS questionnaire, used in our study for sleep screening, effectively identified these common disturbances, underscoring its utility in clinical settings.

Despite the strong evidence for neurobehavioral and sleep comorbidities in pediatric epilepsy, these issues are often underrecognized and undertreated. Achenbach and Rescorla have advocated for the routine use of standardized behavioral tools such as the CBCL in pediatric populations to aid in early detection and intervention. [16] Our findings support this recommendation and emphasize the need for integrative care approaches that encompass seizure management along with behavioral and psychosocial support.

This study had several limitations. The sample size was relatively small, which may limit the generalizability of the findings to the broader pediatric population. As a hospital-based study, selection bias may have influenced the results, as more severe cases tend to be admitted. The cross-sectional design prevented the assessment of causal relationships or longitudinal changes in behavior and sleep. Additionally, reliance on parent-reported tools like the CBCL and BEARS may introduce reporting bias. EEG and imaging findings were not correlated in detail with behavioral outcomes, which could have provided further insights into neuroanatomical or functional contributions.

CONCLUSION

This study highlights the significant neurobehavioral and sleep disturbances experienced by children with epilepsy compared to their non-epileptic peers. Epileptic children demonstrated markedly higher scores in all domains of the Child Behavior Checklist (CBCL), with a majority exhibiting clinically significant behavioral issues. Sleep disorders were also more prevalent in this group, and a strong association was observed between behavioral abnormalities and sleep disturbances. These findings emphasize that epilepsy in children is not merely a seizure disorder but a complex condition affecting multiple aspects of development and daily functioning. Despite seizure control being the primary treatment focus, it is crucial to incorporate routine behavioral and sleep assessments into clinical management. Early identification and intervention can substantially improve quality of life, academic performance, and long-term developmental outcomes. A multidisciplinary approach involving pediatricians, neurologists, psychologists, and caregivers is essential to address the broader spectrum of challenges faced by children with epilepsy.

REFERENCES

- Singh G, Singh MB, Ding D, Maulik P, Sander JW. Implementing WHO's Intersectoral Global Action Plan for epilepsy and other neurological disorders in Southeast Asia: a proposal. The Lancet Regional Health-Southeast Asia. 2023 Mar 1;10.
- Amudhan S, Gururaj G, Satishchandra P. Epilepsy in India I: Epidemiology and public health. Annals of Indian Academy of Neurology. 2015 Jul 1;18(3):263-77.
- Fisher RS. ChAPTER 22 SEIZURES AND EPILEPSY 301. e1. Neurology Secrets: First South Asia Edition-E-book. 2016 Aug 30:300.
- Reilly C, Atkinson P, Das KB, Chin RF, Aylett SE, Burch V, Gillberg C, Scott RC, Neville BG. Neurobehavioral

- comorbidities in children with active epilepsy: a population-based study. Pediatrics. 2014 Jun 1;133(6):e1586-93.
- Karrasch M, Tiitta P, Hermann B, Joutsa J, Shinnar S, Rinne J, Anttinen A, Sillanpää M. Cognitive outcome in childhoodonset epilepsy: a five-decade prospective cohort study. Journal of the International Neuropsychological Society. 2017 Apr;23(4):332-40.
- Rutter M. Isle of Wight revisited: twenty-five years of child psychiatric epidemiology. Annual Progress in Child Psychiatry & Child Development. 1990:131-79.
- Rejnö-Habte Selassie G. Speech and language dysfunction in childhood epilepsy and epileptiform EEG activity. Institute of Neuroscience and Physiology. Department of Clinical Neuroscience and Rehabilitation; 2010 Mar 5.
- 8. Chatterjee S, Bhattacharjee R, Sinha A, Maiti A, Mukherjee A, Ghosh R, Dubey S. Prevalence and pattern of cognitive dysfunction in young adults and middle-aged patients with type-2 diabetes. Journal of Family Medicine and Primary Care. 2025 Jun 1;14(6):2527-37.
- Wirrell EC. Classification of Seizures and the Epilepsies. Epilepsy. 2021 Jun 14:11-22.
- Nobili L, Proserpio P, Combi R, Provini F, Plazzi G, Bisulli F, Tassi L, Tinuper P. Nocturnal frontal lobe epilepsy. Current neurology and neuroscience reports. 2014 Feb;14(2):424.
- Sinha K, Karimi-Abdolrezaee S, Velumian AA, Fehlings MG. Functional changes in genetically dysmyelinated spinal cord axons of shiverer mice: role of juxtaparanodal Kv1 family K+

- channels. Journal of neurophysiology. 2006 Mar;95(3):1683-95.
- 12. Owens JA. Sleep and sleep disorders in children. In: APA handbook of pediatric psychology, developmental-behavioral pediatrics, and developmental science: Pediatric psychology and developmental-behavioral pediatrics: Clinical applications of developmental science, Vol 2. Washington, DC, US: American Psychological Association; 2025. p. 445–75. (APA Handbooks in Psychology® series).
- Rodenburg R, Stams GJ, Meijer AM, Aldenkamp AP, Deković M. Psychopathology in children with epilepsy: a meta-analysis. Journal of pediatric psychology. 2005 Sep 1;30(6):453-68.
- Srinivas HV, Shah U. Comorbidities of epilepsy. Neurology India. 2017 Mar 1;65(Suppl 1):S18-24.
- 15. Smith MT, McCrae CS, Cheung J, Martin JL, Harrod CG, Heald JL, Carden KA. Use of actigraphy for the evaluation of sleep disorders and circadian rhythm sleep-wake disorders: an American Academy of Sleep Medicine systematic review, meta-analysis, and GRADE assessment. Journal of clinical sleep medicine. 2018 Jul 15;14(7):1209-30.
- Achenbach TM, Rescorla LA. Child behavior checklist for ages 6-18. Burlington, VT: University of Vermont; 2001.
- Santhosh NS, Sinha S, Satishchandra P. Epilepsy: indian perspective. Annals of Indian Academy of Neurology. 2014 Mar 1;17(Suppl 1): S3-11.