

Original Research Article

Clinician's Perspectives and Prescription Practice of sodium valproate in the Management of migraine, bipolar disorder, and epilepsy in Indian Healthcare settings

1 ABSTRACT

Objective: To gather clinicians' perspectives on the management of neurological disorders such as migraine, bipolar disorder (BPD), and epilepsy, as well as the prescription practice of sodium valproate in Indian settings.

Methodology: This was a cross-sectional study carried out among clinicians specializing in migraine, BPD, and epilepsy management. The selected participants completed a 29-question questionnaire distributed via email or online platforms. The survey explored prescription practices, clinical observations, and preferences regarding sodium valproate and neurological disorder management. Data analysis involved descriptive statistics, with responses presented as frequencies and percentages.

Results: The survey involved 340 participants, and nearly half (45.88%) of them indicated that sodium valproate was their preferred choice for managing migraine, seizures, and BPD. Around 45% of the respondents noted that women are most frequently affected by migraine. Approximately 55% of the clinicians identified anxiety as the most common comorbid condition associated with migraine. About 47% of the participants preferred sodium valproate for migraine prophylaxis, while the majority (76.76%) of the clinicians chose it as the preferred treatment for BPD. Roughly 40% of the respondents reported that a daily dose of 300 mg of sodium valproate was commonly used for migraine. Approximately 54% of participants found that sodium valproate was most often prescribed to individuals in the 18-45 years age group. Additionally, 65% of the participants reported headaches or dizziness as common adverse effects of sodium valproate.

Conclusion: This study revealed that sodium valproate was a widely preferred treatment for migraine, epilepsy, and BPD among Indian clinicians, particularly for women and individuals aged 18-45 years. The findings highlighted the prevalent use of 300 mg daily dose for migraine and the common association of anxiety with migraine.

2
3 *Keywords: BPD, Epilepsy, Migraine, seizures, sodium valproate*

4 5 1. INTRODUCTION

6
7 The burden of migraine, bipolar disorder, and seizures is significant, affecting millions
8 worldwide. These conditions not only impair individuals' quality of life but also contribute to
9 broader societal challenges, including economic costs and healthcare demands [1-3].

10 Migraine accounts for 4.9% of global ill health measured in years lived with disability (YLDs),
11 impacting over 1 billion people, with India reporting the highest prevalence in 2019 at
12 approximately 213,890,208 cases [4,5]. Globally, around 70 million people live with epilepsy,
13 with nearly 12 million in India [6]. The World Mental Health survey estimates a global
14 prevalence of BPD at 0.8%, with the incidence among adolescents and young adults
15 increasing from 79.21 per 100,000 in 1990 to 84.97 per 100,000 in 2019 [7,8]. In India, a
16 population-based study found current and lifetime prevalence rates of BPAD at 0.3% and
17 0.5%, respectively [8].

18 Sodium valproate (valproic acid), which belongs to the class of drugs known as
19 anticonvulsants or antiepileptics, was approved by the Food and Drug Administration (FDA)
20 for treating complex partial seizures, simple and complex absence seizures, and as an
21 adjunctive therapy for multiple seizure types in both adults and pediatric patients. Sodium
22 valproate inhibits voltage-gated sodium channels to reduce neuronal excitability and seizure
23 activity. It also inhibits gamma-aminobutyric acid (GABA) transaminase, thereby boosting
24 GABA levels and enhancing inhibitory effects. Additionally, sodium valproate promotes
25 GABA synthesis by increasing glutamic acid decarboxylase activity and inhibits histone
26 deacetylases (HDACs), influencing gene expression. Furthermore, sodium valproate
27 modulates calcium channels, impacting neuronal signaling and pain pathways [9].

28 Understanding the prescription practice of sodium valproate was crucial for optimizing its
29 use, improving patient outcomes, and enhancing neuronal health management. The present
30 survey aims to collect expert opinions on the clinical application of sodium valproate
31 maintenance therapy for migraine, epilepsy, and BPD within Indian healthcare settings.

32

33 **2. MATERIALS AND METHODS**

34

35 We carried out a cross-sectional, multiple-response questionnaire-based study involving
36 clinicians with expertise in managing migraine, seizures, and BPD in the major Indian cities
37 from June 2023 to December 2023.

38 **2.1 Questionnaire**

39 The questionnaire booklet named SCORE (Sodium Valproate Efficacy & Tolerability Profile)
40 study was sent to the clinicians who were interested in participating in this study. The
41 SCORE study questionnaire included 29 questions focusing on current prescription
42 practices, clinical observations, and preferences related to sodium valproate, as well as
43 experiences with migraine, epilepsy, and BPD in routine practice.

44

45 **2.2 Participants**

46

47 An invitation was sent to leading clinicians in treating migraine, epilepsy, and BPD in March
48 2023 for participation in this Indian survey. About 340 doctors from major cities of all Indian
49 states representing the geographical distribution shared their willingness to participate and
50 provided necessary data. Participants were asked to complete the questionnaire without
51 discussing it with their peers.

52 **2.3 Statistical Methods**

53

54 Descriptive statistics were employed for data analysis with categorical variables presented
55 as percentages. The frequency of each variable and its corresponding percentage was
56 calculated to illustrate its distribution. Graphs and pie charts were created using Microsoft

57 Excel 2013 (version 16.0.13901.20400) to visually depict the distribution of categorical
58 variables.

59 3. RESULTS

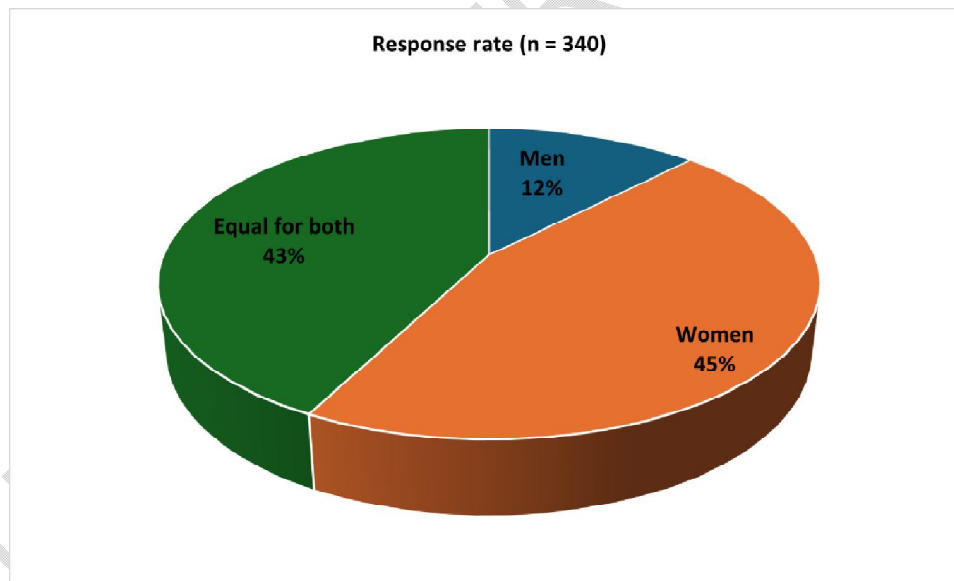
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61 Out of 340 clinicians surveyed, nearly half (45.88%) of the respondents reported that sodium
62 valproate was the preferred choice for migraine prophylaxis, seizures, and BPD (Table 1).
63 Over half (52.65%) of the participants indicated that an average of 6-10 patients present with
64 migraine headaches per day. Approximately 47% of the clinicians noted that fewer than 5
65 patients present with a history of seizures daily, and about 58% of the participants reported
66 that fewer than 5 patients present daily with BPD. Approximately 45% of participants noted
67 that women were most affected by migraine, while 43% of the clinicians observed that both
68 genders equally experience migraine (Fig. 1) and about 55% of the participants identified
69 anxiety as the most common comorbid condition in patients with migraine (Table 2).
70

71 **Table 1: Distribution of response to the preferred use of sodium valproate in clinical**
72 **disorders**

Preference	Response rate (n = 340)
Migraine prophylaxis	9.41%
Seizures	19.71%
BPD	15.88%
All of the above	45.88%

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74

75 **Fig. 1: Distribution of response to the predominant gender presenting with migraine**
76 **headaches in clinical settings**

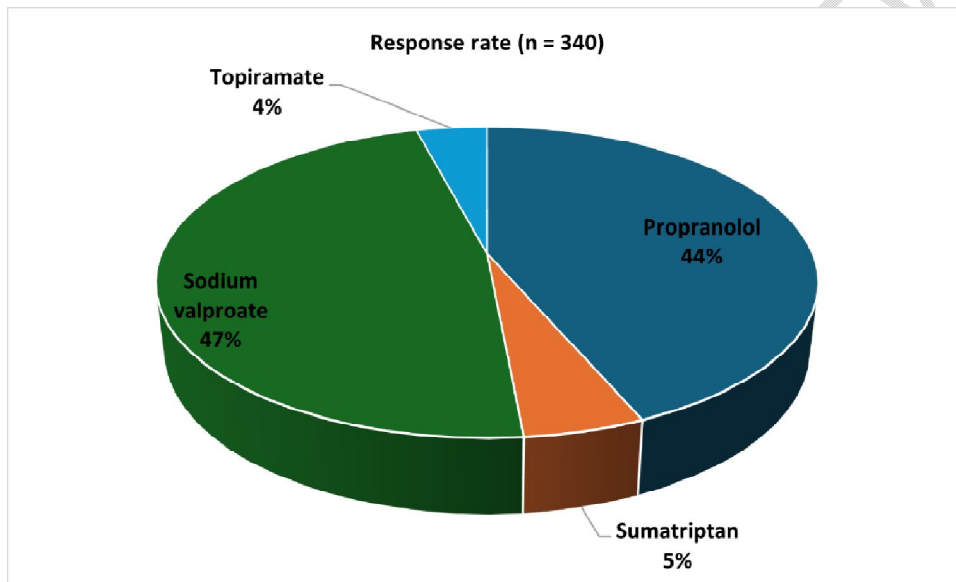
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78 **Table 2: Distribution of response to most common comorbid condition noted in**
79 **patients diagnosed with migraine**

Common comorbid condition	Response rate (n = 340)
Anxiety	55.29%
Panic disorder	14.71%

Hypertension	10%
Diabetes	0.88%
Depression	11.76%
All of the above	7.35%

80 Approximately 27% of clinicians noted that men were most affected by BPD, while about
81 61% of respondents stated that the 18-35 age group was most affected by migraine. About
82 49% of participants indicated that the 36-50 age group was most affected by BPD. About
83 47% of experts preferred sodium valproate for migraine prevention, while 44% of participants
84 preferred propranolol for the same purpose (Fig. 2), while nearly 77% selected it as the
85 preferred treatment for BPD (Table 3).



86

87 **Fig. 2: Distribution of response to preferred drug choice for patients with migraine**
88 **prophylaxis**

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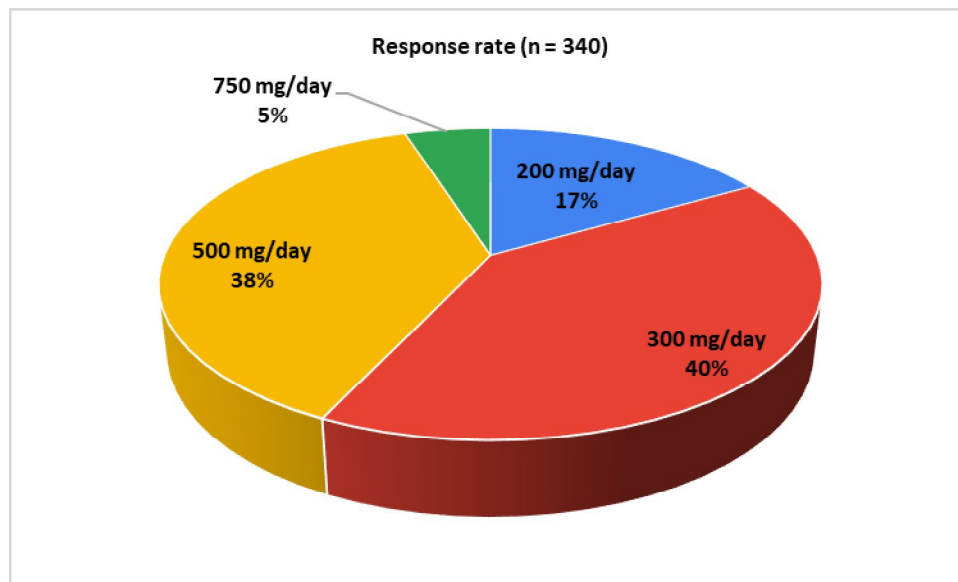
Table 3: Distribution of response to preferred drug choice for patients with BPD

Preference	Response rate (n = 340)
Olanzapine	7.94%
Quetiapine	4.41%
Sodium valproate	76.76%
Lithium	9.12%
No preference	1.76%

91

92 Approximately 54% of participants reported that levetiracetam was the most effective
93 antiepileptic based on its efficacy in seizures. Almost half (48.82%) of the respondents noted
94 that 20-40% of patients were treated with sodium valproate for migraine prophylaxis, and
95 about 43% reported the same proportion of patients receiving the drug as treatment for BPD.
96 Nearly half (50.88%) of the clinicians indicated that 20-40% of patients were treated with
97 sodium valproate for seizures. Approximately 40% of participants indicated that a daily dose

98 of 300 mg of sodium valproate was commonly used for migraine, while 38% reported that
 99 the daily dose was 500 mg (Fig. 3). Approximately 48% of the clinicians stated that a daily
 100 dose of 500 mg was commonly used for BPD, and 56% recommended the same dosage for
 101 seizures. Nearly 54% of the participants found that the 18-45 age group was most
 102 prescribed with sodium valproate (Table 4).



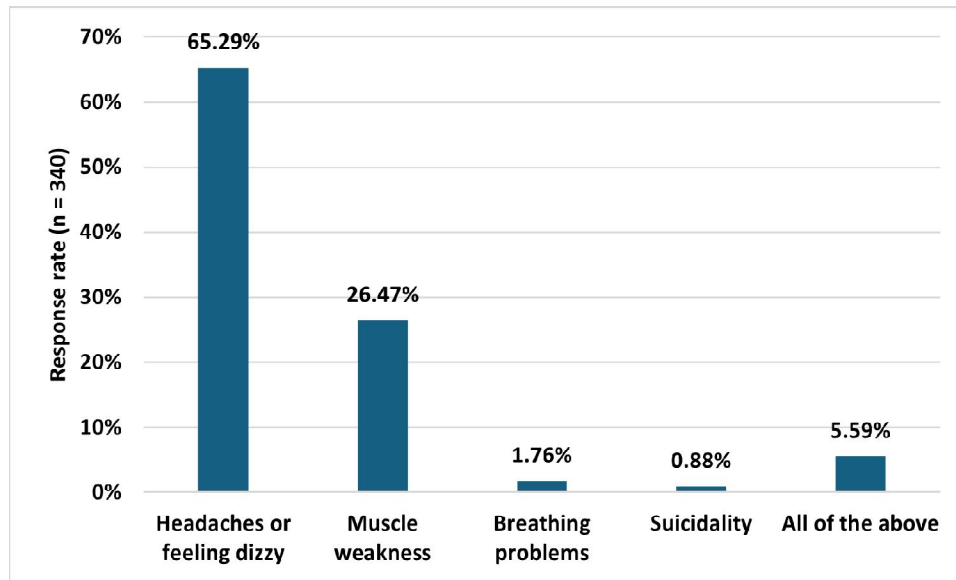
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104 **Fig. 3: Distribution of response to the daily dose of sodium valproate commonly used**
 105 **for migraine**

106 **Table 4: Distribution of response to common age group prescribed with sodium**
 107 **valproate**

Age (years)	Response rate (n = 340)
<12	2.06%
12-18	11.76%
18- 45	54.12%
45-60	10.59%
>60	0.29%
All age groups	21.18%

108 Around 65% of the participants reported that fewer than 5% of patients experience
 109 osteoporosis (OP) or osteopenia as side effects of sodium valproate, with 31% observing
 110 these effects in patients aged 46-60 years. About 39% of the clinicians noted that OP or
 111 osteopenia typically develops after 12-24 months of treatment. Additionally, 65% of the
 112 participants reported headaches or dizziness as common adverse effects of sodium
 113 valproate (Fig. 4).



114

115 **Fig. 4: Distribution of response to adverse effects reported with the use of sodium**
 116 **valproate**

117

118 About 62% of the clinicians stated that brivaracetam has the advantage of lesser behavioral
 119 disturbances compared to sodium valproate, and more than half (58.24%) of the participants
 120 considered the flavor of sodium valproate syrup important for pediatric patients.
 121 Approximately 53% of the respondents found levetiracetam to be the most effective
 122 antiepileptic syrup based on its efficacy in seizures, and around 68% of the participants
 123 reported that levetiracetam injection was most effective for generalized tonic-clonic seizures.
 124 About 35% of the respondents indicated that 16-40% of individuals show improved
 125 outcomes with sodium valproate therapy for migraine, and 47% of the participants preferred
 126 dose escalation as the strategy for patients failing to respond to the initial sodium valproate
 127 treatment for migraine.

128

129 4. DISCUSSION

130

131 The survey indicated that sodium valproate was a widely preferred treatment for migraine,
 132 seizures, and BPD among clinicians in India, reflecting its established role in neurological
 133 disorder management. Vatzaki et al. highlighted that sodium valproate was widely used for
 134 migraine prophylaxis and was included in primary European guidelines [10]. Similarly,
 135 Silberstein et al. found that the drug was effective for migraine prevention, helping to reduce
 136 the frequency and severity of attacks [11]. In the realm of epilepsy, Marques et al. identified
 137 sodium valproate as the most effective antiepileptic drug for genetic generalized epilepsies
 138 [12]. Furthermore, Balagura et al. highlighted its broad spectrum of activity and diverse
 139 mechanisms of action, which have made it a first-line treatment for most seizure types in
 140 children for the past fifty years [13]. In the context of BPD, Roosen and Sienaert provided
 141 evidence supporting the use of sodium valproate for rapid cycling, while Charles L. Bowden
 142 affirmed its value as an effective treatment for BPD [14].

143 Many survey participants observed that women were more severely affected by migraine.
 144 Nappi et al. reported that migraine was more prevalent in women, with 17% of women
 145 meeting the diagnostic criteria for the condition. They also found that migraine frequency in
 146 women varies with the menstrual cycle and pregnancy, and that combined hormonal

147 contraception (CHC) or hormone replacement therapy (HRT) can either trigger or modify
148 migraine [15]. Similarly, Kalkman et al. noted that migraine prevalence was two to three
149 times higher in women compared to men [16]. Singh et al. described migraine as a common
150 neurological disorder with a higher prevalence in women, marked by painful and debilitating
151 headaches [17]. Kumar and Kadian found that the overall prevalence of migraine was
152 estimated at 16%, with a sex prevalence ratio of 3:1, indicating a significantly higher
153 frequency in women [18].

154 Most survey respondents identified anxiety as the most prevalent comorbid condition in
155 patients with migraine. Cuciureanu et al. emphasized that anxiety was a frequent
156 comorbidity associated with migraine, influencing disease prognosis, treatment, and clinical
157 outcomes [19]. Senaratne et al. observed that symptoms of anxiety disorder were
158 significantly more common in patients with migraine than those without [20]. Additionally,
159 Jeyagurunathan et al. reported that anxiety and mood disorders were approximately two to
160 ten times more prevalent among individuals with chronic migraine compared to the general
161 population [21].

162 Many of the participants in the survey preferred sodium valproate for migraine prophylaxis,
163 Kumar and Kadian noted that sodium valproate was used for migraine prophylaxis and was
164 considered one of the first-line agents for migraine prevention [18]. Similarly, Vatzaki et al.
165 reported that sodium valproate was commonly used to treat pre-chronic and chronic
166 migraine [10].

167 The majority of the survey respondents identified sodium valproate as the preferred
168 treatment for BPD. Chen et al. confirmed that sodium valproate was a commonly preferred
169 medication for maintaining BPD [22]. Smith et al. concluded that sodium valproate effectively
170 reduces depressive symptoms in acute bipolar depression and is well-tolerated [23].
171 Additionally, Zheng et al. highlighted that sodium valproate was widely used as an
172 anticonvulsant for the maintenance treatment of BPD [24].

173 Many of the survey participants reported that a 300 mg daily dose of sodium valproate was
174 commonly used for migraine. In line with this finding, Rahman et al. noted that the initial
175 dosage of sodium valproate for migraine prophylaxis typically ranges from 250 to 500 mg,
176 administered twice daily for one week [25]. Diener et al. reported that intravenous
177 administration of sodium valproate, at doses of either 300 mg or 800 mg, was effective in
178 treating acute migraine attacks [26]. Pascual et al. reported that the maintenance dose of
179 sodium valproate typically ranges from 300 to 1000 mg daily [27].

180 Many of the participants stated that the 18-45 age group was most prescribed sodium
181 valproate. In a study by Evans et al. conducted in England and Wales, it was reported that
182 87.7 out of every 1,000 individuals prescribed valproate were women or girls aged 14 to 45
183 years [28].

184 Most of the participants identified headaches and dizziness as common adverse effects of
185 sodium valproate. Rahman et al. confirmed that dizziness and headaches were frequently
186 associated with the drug [25]. Montalbano et al. reported additional common side effects
187 including nausea, vomiting, constipation, increased appetite, weight gain, somnolence, and
188 tremor [29]. Philip B. Bradley also noted that sodium valproate can cause dizziness, mild
189 hypotension, and mild thrombocytopenia in some patients [30].

190 The major strengths of the survey include its large sample size and input from neurological
191 disorder specialists regarding the effectiveness of sodium valproate. However, the results
192 may be subject to bias due to reliance on expert opinion, and varying perspectives among

193 clinicians could affect the findings. Additionally, the survey might not fully incorporate
194 emerging evidence or evolving trends in the management of neurological disorders such as
195 migraine, seizures, and BPD. It is crucial to acknowledge these limitations when interpreting
196 the results and to recognize the need for further research to confirm the findings of the
197 survey. Future studies could investigate the long-term effects of sodium valproate and
198 explore strategies for improving patient education.

199 **5. CONCLUSION**

200

201 This study's findings highlight sodium valproate as the preferred treatment for managing
202 neurological disorders such as migraine, BPD, and epilepsy, aligning with earlier studies that
203 recognize its effectiveness and its status as a first-line treatment. The data indicate its
204 prevalent use across various age groups, particularly in the 18-45 age range, with dosage
205 preferences ranging from 300 mg to 500 mg daily.

206 **Ethical Approval:**

207 The study was conducted after getting approval from Bangalore Ethics, an Independent
208 Ethics Committee which was recognized by the Indian Regulatory Authority, Drug Controller
209 General of India.
210

211

212 **Consent:**

213 A written informed consent was obtained from each physician before initiation of the study.
214

215

215 **Disclaimer (Artificial intelligence)**

216 Author(s) hereby declare that NO generative AI technologies such as Large Language
217 Models (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the
218 writing or editing of this manuscript.

219 **COMPETING INTERESTS**

220 Authors have declared that they have no known competing financial interests OR non-
221 financial interests OR personal relationships that could have appeared to influence the work
222 reported in this paper.

223

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