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ORIGINAL RESEARCH

Treatment-Seeking Behavior Towards Epilepsy Among Rural Residents in Ethiopia: A Cross-Sectional Study

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Background: Even though a large number of patients live with epilepsy, few of them receive effective treatment. Most people with epilepsy, particularly those from rural communities, do not seek medical care as they are convinced that solutions lie only with traditional healers (traditional leaders, prophets and community elders). Therefore, studying treatment-seeking behavior regarding this major neurological problem would provide additional knowledge and help to identify a gap which needs to be addressed when tackling related problems.

Participants and Methods: This rural community-based cross-sectional study was conducted in Gedeo zone, Ethiopia, among 755 participants. Data were collected using face-toface interview-based questionnaires. Collected data were entered into EpiData version 3.01 and analyzed using SPSS version 20. Bivariate and multivariate logistic regression analysis was conducted to identify associated factors. The strengths of associations were presented as adjusted odds ratios with 95% confidence intervals.

Results: The prevalence of poor treatment-seeking behavior was 54.6% (95% CI 51.4, 58.2). Poor knowledge (adjusted odds ratio [AOR]=3.21, 95% CI 2.14, 4.81), poor social support (AOR=2.48, 95% CI 1.12, 5.53), unfavorable attitude (AOR=1.84, 95% CI 1.34, 2.54) and having no history of experiencing others' seizures (AOR=2.17, 95% CI 1.47, 3.2) were variables strongly associated with poor help-seeking behavior towards epilepsy.

Conclusion: The study showed that more than half of the participants had poor treatmentseeking behavior towards epilepsy. This indicates the need to implement measures to raise community awareness regarding treatment options for epilepsy.

Keywords: epilepsy, Ethiopia, rural residents, treatment-seeking behavior

Introduction

Epilepsy is a common chronic non-communicable disease of the brain that affects people of all age groups. It carries neurological, cognitive, psychological and social consequences, and globally more than 65 million peoples are affected by epilepsy.^{1–3} Developing countries carry 90% of the financial burden of epilepsy and 85% of the world's people with epilepsy live in developing countries with little or no access to medical treatment.³

Despite the availability of effective and low-cost anti-seizure medicines, more than 80% of people with epilepsy in low-income countries do not have access to treatment.^{2,4} Evidence also shows that up to 70% of people with epilepsy could become seizure free with the appropriate use of anti-seizure medicines.^{4,5}

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© 2020 Molla et al. This work is published and licensed by Dove Medical Press Limited. The full terms of this license are available at https://www.dovepress.com/terms.php and incorporate the Creative Commons Attribution – Non Commercial (unported, v3.0) License (http://treativecommons.org/licenses/by-mc/3.0/). By accessing the work you hereby accept the firms. Non-commercial uses of the work are permitted without any further permission from Dove Medical Press Limited, provided the work is properly attributed. For permission for commercial use of this work, please see paragraphs 4.2 and 5 of our Terms (http://www.dovepress.com/terms.php). In Ethiopia about 81% of people with epilepsy and their relatives suffer from perceived epilepsy stigma, which has a direct influence on treatment options.^{6,7} For centuries, religious and cultural taboos have influenced the type of care and treatment available for people with epilepsy.⁸ In many African communities the myths regarding epilepsy continue to be a cause of wide treatment gaps. Sociocultural attitudes continue to have a negative impact on the management of epilepsy in many African nations.⁹ Wrong attitudes and beliefs about epilepsy create serious negative social and psychological consequences for people with epilepsy, such as fear, humiliation and limitations in social interactions.¹⁰ As a result, people with epilepsy are burdened by social, psychological and economic consequences, which lead to poor quality of life.^{11,12}

Considering this, it is important to know more about treatment perceptions towards epilepsy which can help to improve people's treatment-seeking behavior, to reduce the psychosocial and economic impacts related to disease among community residents.^{10,13} Therefore, this study was conducted to assess treatment-seeking behavior towards epilepsy among rural residents in southern Ethiopia.

Participants and Methods

This rural community-based cross-sectional study was conducted in Gedeo zone in southern Ethiopia from 1 to 30 March 2019. Gedeo zone is located 365 km from Addis Ababa, the capital city of Ethiopia. All adults aged \geq 18 years and who were available during the data-gathering period were invited as study participants. Individuals who were ill and unable to communicate during the data-collection time were excluded.

Sample size was calculated using a single population proportion formula, considering the following assumptions: proportion of residents having poor practice of seeking modern treatment 66.5%,⁹ margin of error 0.05, 95% confidence interval (CI) and design effect 2. After adding a 10% nonresponse rate, the final calculated sample size was 755. To select samples, a multistage sampling technique was used. First, out of six rural districts in Gedeo zone, three districts were selected at random. In the second step, nine subdistricts (three in each district) were taken, again at random. The list of households in each of the area was obtained from the district offices and health extension workers. The calculated sample size was allocated proportionally in the selected subdistricts based on the number of households belonging to the subdistricts. Finally, participants were interviewed in the household; in the case of more than one participant in the household being eligible, we selected one individual by a lottery method.

Data were collected using a structured and interviewadministered questionnaire. The content of the questionnaire included sociodemographic variables, Stigma Scale of Epilepsy, knowledge- and attitude-related variables, helpseeking behavior questionnaires, Oslo Social Support Scale and Substance-Related Questionnaire. Knowledge and attitude were assessed using the Kilifi Epilepsy Beliefs and Attitude Scale.¹⁴ Epilepsy stigma was assessed using the 24-item Stigma Scale of Epilepsy, which is validated for use in resource-limited countries.¹⁵ Help-seeking behavior was assessed by WHO General Health-Seeking behavior questionnaires and participants who scored greater than or equal to the mean score were considered as having good help-seeking behavior.¹⁶ The Alcohol, Smoking and Substance Involvement Screening Test (WHO ASSIST version 3.0), which is a commonly used and culturally validated tool, was used to assess lifetime and current substance use.¹⁷ Social support was collected by the 3-item Oslo Social Support Scale; this culturally validated tool has a sum score ranging from 3 to 14 in three categories: poor support 3-8, moderate support 9–11 and strong support 12-14.¹⁸

We used nine trained diploma nurses as data collectors and three Bachelor of Science (BSc) psychiatric nurses as supervisors. The questionnaire was translated from the English version to the local language and back to the English language to check its consistency. The data collectors were supervised daily and the completed questionnaires were checked daily by the supervisors and investigators for completeness. Data were entered into the computer using EpiData version 3.1 and exported to SPSS version 20 for analysis. Descriptive data were summarized using tables. Binary logistic regression was used to identify factors associated with the outcome variable. Variables with a P-value of <0.2 in bivariate regression were included in multivariable logistic regression. Finally, a *P*-value ≤ 0.05 in multivariate logistic regression was considered statistically significant and the strength of associations was determined using the adjusted odds ratio (AOR) and 95% CI.

Results

Sociodemographic Distribution of Respondents

A total of 732 participants were included in this study, with a response rate of 97%. The mean±SD age of the

respondents was 33 ± 11.3 years. Among the participants, the majority [316 (43.2%)] were in the age range 18–28 years and 467 (63.8%) were male. Of the total participants, a large proportion [513 (70.1%)] were Protestant in their religion and 632 (86.3%) were from a Gedeo ethnic background. The majority of participants [513 (70.1%)] were married. The educational status of participants showed that 276 (37.7%) had been educated at primary level. Regarding occupation, 232 (31.7%) of participants [718 (98.1%)] could access health centers at a distance of <10 km. The majority of respondents had monthly income of less than \$1.9 [538 (73.5%)] (Table 1).

Psychosocial and Substance-Related Factors of Respondents

Regarding the psychosocial characteristics of the respondents, 347 participants (47.4%) reported stigma with respect to epilepsy and 451 (61.6%) reported poor social support. Of the participants, 173 (23.6%) reported current alcohol use, 40 (5.5%) that they chew khat and 39 (5.3%) that they currently smoke cigarettes (within 3 months of the study) (Table 2).

Clinical and Related Factors

Although 54 respondents (7.4%) reported a previous history of epilepsy, only 12 (1.6%) of them had obtained help for their problem. Among participants who received help for their epilepsy, seven (58.33%) obtained help from religious centers and three (25%) from traditional healers. Only two respondents (16.66%) accessed help from health centers. Among the respondents who did not seek help for epilepsy, most [31 (73.8%)] reported that epilepsy has no modern treatment. Furthermore, 27% and 51.6% of participants had poor knowledge and an unfavorable attitude, respectively. About 21% of study participants had a history of witnessing seizures. Regarding family history, 61 respondents (8.3%) and 76 (10.4%) reported a family history of epilepsy and mental illness, respectively. Furthermore, 58 participants (7.9%) reported a history of head injury (Table 3).

Factors Related to Treatment-Seeking Behavior Towards Epilepsy

More than half of the participants [400 (54.6%)] scored below the mean score, which is an indicator of poor help-seeking behavior towards epilepsy according to the WHO general help-seeking questionnaire. Regarding the place in which TableIFrequencyDistributionofParticipantsbySociodemographicFactorsinRuralGedeoZone,Ethiopia,2019(n=732)

Variable	Category	Frequency	Percent (%)
Age	18–28 years	316	43.2
-	29–39 years	231	31.6
	40–50 years	112	15.3
	>50 years	73	10
Sex	Female	265	36.2
	Male	467	63.8
Religion	Protestant	513	70.1
	Christian		
	Orthodox	117	16
	Christian		
	Muslim	42	5.7
	Catholic	40	5.5
	Others ^a	20	2.7
Marital status	Married	513	70.1
	Single	147	20.1
	Widowed	35	4.8
	Divorced	25	3.4
	Separated	12	1.6
Ethnicity	Gedeo	632	86.3
	Oromo	45	6.1
	Amhara	28	3.8
	Gurage	20	2.7
	Others ^b	8	1.1
Educational status	Cannot read and write	196	26.8
	Primary	276	37.7
	Secondary	151	20.6
	Preparatory	51	7
	College and above	58	7 7.9
Occupational status	Government employed	96	13.1
	Farmer	232	31.7
	Business	105	14.3
	Private employed	116	15.8
	Housewife	94	12.8
	Daily laborer	24	3.3
	Others ^c	65	8.9
Distance from health	<10 km	718	98.1
center	≥10 km	14	1.9
Average monthly	<\$1.9	538	73.5
income	≥\$1.9	194	26.5

Notes: ^aJehovah's Witnesses and no religion; ^bTigre, Wolaita and Silte; ^cjobless and students.

they sought treatment, most respondents [416 (56.8%)] preferred religious centers for epilepsy treatment, and 305

Variable	Category	Frequency	Percent (%)
Stigma related to	Yes	347	47.4
epilepsy	No	385	52.6
Social support	Poor	451	61.6
	Moderate	249	34
	Strong	32	4.4
Ever alcohol use	Yes	300	41
	No	432	59
Ever cigarette use	Yes	70	9.6
	No	662	90.4
Ever khat use	Yes	88	12
	No	644	88
Current alcohol use	Yes	173	23.6
	No	599	76.4
Current cigarette use	Yes	39	5.3
	No	693	94.7
Current khat use	Yes	40	5.5
	No	692	94.5

Table 2Distribution of Psychosocial and Substance-RelatedFactors Among Rural Residents in Gedeo Zone, 2019 (n=732)

participants (41.7%) reported that they did not prefer health centers as a treatment option for epilepsy. Among community participants, 147 (20.1%) reported they would not seek any treatment options for epilepsy (Table 4).

Factors Associated with Treatment-Seeking Behavior Towards Epilepsy

In multivariable binary logistic regression, the variables unfavorable attitude, poor knowledge, poor social support and having no history of experiencing others' seizures were statistically significantly correlated with poor helpseeking behavior towards epilepsy (p<0.05).

Participants with poor knowledge were 3.21 times more likely to have poor help-seeking behavior than those with higher knowledge levels (AOR=3.21, 95% CI 2.14, 4.81).

The factor of having no history of experiencing others' seizures was strongly associated with lack of help-seeking behavior in this study. The odds of having poor help-seeking behavior among participants who had no history of experiencing others' seizures were 2.17 times higher than in those with such a history (AOR=2.17, 95% CI 1.47, 3.2).

The odds of having poor help-seeking behavior among respondents with a history of an unfavorable attitude were

Table 3 Descriptions of Clinical-Related Factors Among Rura	l
Residents in Gedeo Zone, 2019 (n=732)	

Variable	Category	Frequency	Percent (%)
Knowledge about	Good	534	73
epilepsy	Poor	198	27
Attitude towards	Favorable	354	48.4
epilepsy	Unfavorable	378	51.6
Pervious history of epilepsy	Yes	54	7.4
	No	678	92.6
Get help for illness	Yes	12	1.6
	No	720	98.4
Place of treatment	Health institutions Religious centers Traditional healers	2 7 3	16.7 58.3 25
Reason not to get help	No treatment No money Does not need treatment	31 1 10	73.8 2.4 23.8
Witnessing seizure	Yes	154	21
	No	578	79
Family history of epilepsy	Yes	61	8.3
	No	671	91.7
Family history of mental illness	Yes	76	10.4
	No	656	89.6
History of head	Yes	58	7.9
injury	No	674	92.1

1.84 times higher than in those without this attitude (AOR=1.84, 95% CI 1.34, 2.54).

The other factor that was strongly associated with poor help-seeking behavior was poor social support. The odds of having poor help-seeking behavior among participants with poor social support were 2.48 times higher than among participants with good social support (AOR=2.48, 95% CI 1.12, 5.53) (Table 5).

Discussion

Epilepsy is a major health problem in Ethiopia.¹⁹ This study aimed to assess treatment-seeking behavior towards epilepsy and associated factors among residents of rural communities in southern Ethiopia.

The study showed that 54.6% of respondents had poor treatment-seeking behavior. This proportion is similar to

Variable	Category	Frequency	Percent (%)
Overall treatment-seeking behavior	Good	332	45.4
	Poor	400	54.6
Seek help from close relatives (family)	Yes	464	63.4
	No	268	36.6
Seek help from friends	Yes	294	40.2
	No	438	59.8
Seek help from religious centers	Yes	416	56.8
	No	316	43.2
Seek help from modern health institutions	Yes	427	58.3
	No	305	41.7
Never seek help from any of the above	Yes	147	20.1
	No	585	79.9

Table 4 Treatment Seeking-Related Factors	s Towards Epilepsy Among Rural Re	sidents in Gedeo Zone, 2019 (n=732)
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Table 5Bivariate and Multivariate Logistic Regression Analysis of Factors Associated with Poor Help-SeekingBehavior Among Rural Residents in Gedeo Zone, Ethiopia, 2019 (n=732)

Explanatory Variable	Help-Seeking Behavior		Crude Odds Ratio	Adjusted Odds Ratio	
	Poor	Good	(95% CI)	(95% CI)	
Educational status					
Cannot read and write	121	75	1.85 (1.03, 3.35)	1.18 (0.6, 2.32)	
Primary	145	131	1.27 (0.72, 2.24)	0.98 (0.52, 1.86)	
Secondary	77	74	1.2 (0.65, 2.2)	1.07 (0.56, 2.07)	
Preparatory	30	21	1.64 (0.77, 3.51)	1.79 (0 0.80, 4.11)	
College and above	27	31	1	I	
Knowledge about epilepsy					
Good	246	288	1	1	
Poor	154	44	4.1 (2.81, 5.97)	3.21 (2.14, 4.81)**	
Having no experience of others' seizures					
Yes	56	98	1	1	
No	344	234	2.57 (1.78, 3.72)	2.17 (1.47, 3.2)**	
Attitude					
Unfavorable	242	136	2.21 (1.64, 2.97)	1.84 (1.34, 2.54)**	
Favorable	158	196	1	I	
Social support					
Poor	255	196	2.86 (1.33, 6.18)	2.48 (1.12, 5.53)*	
Moderate	135	114	2.61 (1.19, 5.73)	2.16 (0.95, 4.90)	
Strong	10	22	1	1	
Average monthly Income					
<\$1.9	307	231	1.44 (1.04, 2.01)	0.99 (0.68, 1.46)	
≥\$1.9	93	101	1	1	

Notes: 1.0=reference, **P<0.01, *P>0.01, Hosmer Lemeshow test P=0.66.

that found in a study conducted in Jimma University, Ethiopia (53.3%),²⁰ but lower than in another study conducted in Ethiopia in which 66.5% had poor practice of

seeking treatment for epilepsy.⁹ It is also lower than in a study conducted in low-income countries in which about 80% of respondents did not seek any treatment.²¹ In the

current study, 41.7% of participants did not seek help from modern health institutions; this result is supported by a study conducted from Nigeria in which 55.8% of participants did not take epileptic patients to a hospital during a seizure.²² This may be a result of cultural influences regarding the causes of and treatment options for seizure, as people in many developing countries believe that epilepsy cannot be cured by modern treatment, but rather by traditional treatments. Another possible reason may be due to stigma, as many people consider that having an epileptic patient is stigmatizing to the community, so they are afraid to take the patient to a public place such as a hospital.²³ Another possible explanation is the difference in the tools used; the study conducted in Nigeria used an author-developed tool, whereas in this study we tried to use tools that have been approved by the WHO to assess help-seeking behavior.

Regarding factors, poor knowledge, having no history of experiencing epileptic seizures, unfavorable attitude and poor social support were strongly associated with poor help-seeking behavior towards epilepsy. This finding is supported by studies conducted in Sudan,²⁴ Ethiopia⁹ and Asia.²⁵ These factors may be due to social isolation and poor social adaptation resulting from perceived stigma, which can make it difficult for people to seek treatment for their illness; however, good social support is vital for good health in reducing the risk of having epilepsy.²⁶ Participants with poor social support may not receive information regarding the treatment options for epilepsy. Another possible reason is that participants with a history of having experienced others' seizures may have had to see different treatment options owing to the frequent impact of seizures on the physical and emotional health of those patients. Most studies also showed that attitude had a direct relationship with treatment-seeking behavior towards epilepsy, particularly in rural communities, and sociocultural attitudes continue to have a negative impact on the management of epilepsy in Ethiopia.⁹ Limitations of this study are that some ideal factors that might have been addressed by a qualitative study were not included, and the nature of the study design could not establish a temporal relationship between the outcome and the independent variables.

Conclusion

The current study showed that participants had poor treatment-seeking behavior. Factors such as poor social support, having no history of experiencing others' seizures, unfavorable attitude and poor knowledge were associated with poor treatment-seeking behavior. Community awareness regarding treatments for epilepsy needs be raised.

Compliance with Ethical Standards

All procedures undertaken during data collection were in accordance with the ethical review board of Dilla University and the Helsinki Declaration. The study was approved by the ethical review board of Dilla University. The participants were informed about the aim of the study and any identification or names were not recorded to maintain confidentiality. The right was also given to the study participants to refuse or discontinue participation at any time during the interview. Finally, data were collected after obtaining written informed consent from the participants.

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Author Contributions

All authors contributed towards data analysis, drafting and revising the manuscript, gave final approval of the paper to be published, and agreed to be accountable for all aspects of the work.

Disclosure

The authors declare that they have no conflicts of interest in this work.

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