Access to healthcare and prevalence of anxiety and depression in persons with epilepsy during the COVID-19 pandemic: A multicountry online survey☆☆☆

Stijn Van Hees a,b, Joseph Nelson Siewe Fodjo a, Veerle Wijtvliet b, Rafael Van den Bergh a, Edlaine Faria de Moura Villela c, Carolina Ferreira da Silva c, Sarah Weckhuysen d,e, Robert Colebunders a,*

a Global Health Institute, University of Antwerp, Belgium
b Laboratory of Experimental Medicine and Pediatrics, University of Antwerp, Belgium
c School of Medicine, Health Sciences Unit, Federal University of Jataí, Brazil
d Department of Neurology, University Hospital Antwerp, Belgium
e VIB-Center for Molecular Neurology, University of Antwerp, Belgium

Abstract

Objective: The objective of this study was to assess access to healthcare and to estimate the prevalence of depression and anxiety among persons with epilepsy (PWE) during the ongoing coronavirus disease 2019 (COVID-19) pandemic.

Methods: We conducted a multicountry online survey among PWE. Persons with epilepsy were invited to participate through various social media channels. The Hospital Anxiety and Depression Scale (HADS) and 9-item Patient Health Questionnaire (PHQ-9) scale were used to score anxiety and depression. Logistic regression modeling was used to investigate factors associated with anxiety and depression.

Results: Three hundred ninety-nine PWE were included (mean age: 38.22 ± 12.09 years), the majority were female (80.2%) and living in high-income countries (83.2%). Two hundred three PWE reported symptoms of a cold since January 2020. Nine (25%) of the 36 PWE tested for COVID were positive. A total of 72 PWE (19.6%) reported problems to obtain antiseizure medication (ASM), which in 25% of cases was directly COVID-related. Of the 399 PWE, 201 (50.4%) screened positive for anxiety according to the HADS; 159 (39.8%) and 187 (46.9%) PWE screened positive for depression based on the HADS and PHQ-9 scale, respectively. Female gender and financial problems were associated with both depression and anxiety. A planned follow-up consultation with the treating physician was associated with a lower risk of depression, whereas difficulties to access ASM treatment increased the odds of depression. In 65/137 (47.4%) PWE with a planned follow-up visit with the treating physician, this consultation was canceled.

Conclusions: Innovative approaches are needed to ensure continuity in access to ASM treatment. Healthcare workers should ensure continued follow-up, either through inperson or telehealth appointments, to timely identify symptoms of anxiety and depression and act accordingly.

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1. Introduction

Epilepsy is a chronic neurological disease, affecting more than 50 million people worldwide [1]. Besides the physical impact of epileptic seizures, epilepsy also has important neurobiological, cognitive, and psychosocial consequences [2]. Psychiatric comorbidities are up to three times more prevalent in persons with epilepsy (PWE) than in the general population. Depression and anxiety are the most frequently encountered, with reported lifetime prevalence rates as high as 30–35% and 22.8%, respectively [3–5]. Psychiatric comorbidities are associated with poor quality of life (QoL) [6], seizure severity [7], more severe adverse effects of antiseizure medication (ASM) [8], therapy resistance [9], and poor outcomes after epilepsy surgery [10]. Despite their tremendous impact, psychiatric comorbidities in PWE are often overlooked. Persons with epilepsy represent a vulnerable population because of increased rates of unemployment, in turn, resulting in financial problems and social stigma [11].

The ongoing coronavirus disease 2019 (COVID-19) pandemic puts the global population under important psychosocial distress. Reports
from China, Turkey, and Italy suggest that during the COVID-19 pandemic, up to 25% and 45% of the general population show symptoms of depression and anxiety, respectively [12–14]. The coronavirus crisis leads to feelings of loneliness, boredom, anger, and anxiety and poses a threat to the public mental health [15]. Previous studies have shown higher incidence rates of mental disorders including depression and anxiety during outbreaks of acute respiratory infections, such as the influenza virus [16,17]. Furthermore, in survivors of severe acute respiratory syndrome coronavirus 1 (SARS-CoV-1), an increased prevalence of neuropsychiatric disorders was noted including posttraumatic stress disorder, depression, and panic disorder [18]. Next to posing a mental burden, COVID-19 may also delay access to and provision of care for non-COVID-related health problems, causing deaths that could have been prevented [19].

The prevalence of depression and anxiety among PWE as well as access to healthcare by PWE during the COVID-19 pandemic has not been well studied so far. A study from southwest China by Hao et al. reported that PWE experience more psychosocial distress compared with healthy subjects during the COVID-19 pandemic [20]. In the latter study, the diagnosis of refractory epilepsy and time spent to follow the evolution of the COVID-19 pandemic were predictive of severe psychological distress [20]. In a bid to expand the knowledge regarding the health and medical follow-up of PWE during the COVID-19 pandemic, we conducted a multicountry online survey to evaluate their access to healthcare and the prevalence of anxiety and depression in this population.

2. Methods

2.1. Study design

An online questionnaire survey among PWE was organized between April 10th and May 18th, 2020. At the beginning of the study period, Europe was still in full lockdown with healthcare services focused on COVID-19 and nonpharmaceutical interventions such as physical distancing highly recommended by local governments. By the first half of May, Europe had started to gradually lift lockdown measures. In Brazil and North America, there were large differences between regions in terms of lockdown and healthcare situations.

Respondents were invited to participate by clicking on an URL link that was shared through various websites or social media channels of associations that support persons with epilepsy and epilepsy research such as the Epilepsy Liga Flanders, the Brazilian Federation of Epilepsy (EPIPBrasil), the International League Against Epilepsy (ILAE), the Brazilian Association of Epilepsy (ABE), and Citizens United for Research in Epilepsy (CURE). The link was accessible to anyone.

The questionnaire consisted of five parts, inquiring on demographic data, epilepsy-related data, COVID-related data, and anxiety- and depression-related data. The survey was available in English, Dutch, French, and Portuguese. Respondents were included if they provided e-consent, reported to be a PWE or a caretaker/parent of a PWE, and were older than 18 years. Caretakers/parents of a PWE were asked to complete the questionnaire from the perspective of the PWE. A duplicate of the survey template can be found in the supplementary data of this paper. Respondents were categorized as living in low- to middle-income countries or high-income countries according to the World Bank Classification [21].

This study was approved by the ethics committee of the Antwerp University Hospital. All participants provided e-consent.

2.2. Assessment of anxiety and depression

The prevalence of anxiety and depression among the participants was estimated using the Hospital Anxiety and Depression Scale (HADS) and the 9-item Patient Health Questionnaire (PHQ-9) [22–29]. Both screening tools have been validated as a self-reporting tool for use in PWE [22–29]. The HADS consists of two main sections of seven questions each, to screen for depression and anxiety. The answer options followed a 0–3 Likert scale format, with 0 corresponding to no sign of anxiety or depression and 3 for the highest level of anxiety or depression for a particular item in the screening tool. The overall scores ranged from 0 to 21 for anxiety and depression screening, respectively. The PHQ-9 screening tool scores each of the 9 items of the Diagnostic and Statistical Manual of Mental Disorders (DSM-IV) criteria for depression between 0 (not at all) and 3 (nearly every day).

Persons with epilepsy with HADS scores of 7 and above in each section of the scale (anxiety and depression) were considered as screened positive for that condition. A cutoff value of 9 was used to define depression according to the PHQ-9 scale, such that any PWE scoring 9 and above was classified as suffering from depression. These cutoffs were shown to result in the best sensitivity and specificity [22–29].

2.3. Statistical analyses

Continuous variables are reported as mean with standard deviation (mean ± SD) or median with interquartile range (IQR), categorical variables as numbers with percentages. Continuous variables were compared between groups using a Student's t- or Mann–Whitney U test as appropriate, categorical variables using a chi-square or Fisher’s exact test. Multiple logistic regression modeling was used to investigate factors associated with anxiety and depression (both based on HADS). Factors with a p < 0.100 in bivariate analysis (uncorrected for multiple testing, with a view on being inclusive) were included in multivariate analyses. All statistical tests were two-sided, and p < 0.05 was considered significant. Statistical analyses were performed in Statistical Product and Service Solutions (SPSS) 26.0.

3. Results

3.1. Respondent characteristics

A total of 460 responses were collected. Of these, 37 respondents who reported not to be a PWE nor a parent/caretaker of a PWE and 24 who were younger than 18 years old were excluded from analysis. Three hundred ninety-nine PWE from 18 countries (57.6% from Belgium, 14.5% from Brazil, 9.8% from The Netherlands, 7.9% from the United States, 11.2% from other countries; Supplementary Table 1) were included in the study; the mean age was 38.22 ± 12.09 years, and 80.2% were female (Table 1). A minority of the PWE lived in a low- to middle-income country (67/399; 16.8%). Of the 399 respondents, 35 reported to be a parent or caretaker of a PWE. We compared responses provided by the caretaker or parent with those directly provided by PWE (data not shown) and did not identify any significant difference. Therefore, we pooled the data of both groups for further analysis.

3.2. Epilepsy characteristics

The majority of the PWE were on ASM (368/399; 92.2%). There was no difference between the proportion of PWE who reported being seizure-free during the last three months before the onset of COVID-19 in their respective countries (238/399; 59.7%) and during the last three months preceding completion of the questionnaire (229/399; 57.4%; p = 0.518). Prior to the COVID-19 outbreak and associated restrictions, 137 PWE had a planned follow-up consultation with the treating physician. For 27 (19.7%) of them, this follow-up visit took place as planned; for 65 (47.4%), this visit was canceled because of the COVID-19 restrictions; and for 45 (32.8%), this visit was changed to a telephone or online consult.

3.3. COVID-19 symptoms and testing

Of the total of 399 responding PWE, 203 reported at least one symptom of a cold since January 2020. The most frequently reported symptoms
(Fig. 1) were a stuffy and/or running nose (167/203 PWE; 82.3%), headache (141/203 PWE; 69.5%), and a sore throat (120/203 PWE; 59.1%).

**Table 1**

| Demographic data | Total n = 399 |
|------------------|--------------|
| Age In years (mean ± standard deviation) | 38.22 ± 12.09 |
| Gender Male; n (%) | 79 (19.8%) |
| Country of residence Low- to middle-income country; n (%) | 67 (16.8%) |
| High-income country; n (%) | 332 (83.2%) |
| Marital status Single; n (%) | 133 (33.3%) |
| Highest educational level Primary school; n (%) | 16 (4.0%) |
| Secondary school; n (%) | 158 (39.6%) |
| University undergraduate degree; n (%) | 159 (39.8%) |
| University postgraduate degree; n (%) | 53 (13.3%) |
| None; n (%) | 13 (3.3%) |

**Epilepsy data**

| Seizure-free during the last three months before COVID | Yes; n (%) | 238 (59.6%) |
|--------------------------------------------------------|-----------|------------|
| If not seizure-free, number of seizures over the last three months; median (IQR) | 3 (1–9) |

**On antiseizure medication**

| Yes; n (%) | 368 (92.2%) |
| 2; n (%) | 100 (27.2%) |
| 3; n (%) | 47 (12.8%) |
| 4; n (%) | 18 (4.9%) |
| 5; n (%) | 4 (1.1%) |

**On antidepressants or anxiolytics**

| No; n (%) | 313 (78.4%) |
| Yes, since a long time; n (%) | 77 (19.3%) |
| Yes, since the start of COVID-19; n (%) | 7 (1.8%) |
| No answer reported; n (%) | 2 (0.5%) |
| FU consultation with neurologist planned before onset of corona Yes; n (%) | 137 (34.3%) |

**If yes, what happened to the consultation**

| Canceled; n (%) | 65 (41.7%) |
| Changed to telephone or online consult; n (%) | 45 (23.8%) |
| Took place as planned; n (%) | 27 (19.7%) |

**Who do you live with?**

| With spouse/partner; n (%) | 221 (55.4%) |
| With siblings or other family relatives; n (%) | 30 (7.5%) |
| With friends; n (%) | 12 (3.0%) |
| Alone; n (%) | 58 (14.5%) |
| With child(ren); n (%) | 160 (40.1%) |
| With friend(s); n (%) | 12 (3.0%) |
| None; n (%) | 13 (3.3%) |

**Country of residence**

| Low- to middle-income country; n (%) | 238 (59.6%) |
| High-income country; n (%) | 131 (33.1%) |

**Marital status**

| Single; n (%) | 133 (33.3%) |
| Married; n (%) | 238 (59.6%) |
| Divorced; n (%) | 18 (4.6%) |
| Widowed; n (%) | 9 (2.3%) |

**Epilepsy data**

| Yes; n (%) | 238 (59.6%) |
| No; n (%) | 313 (78.4%) |

**Access to antiseizure medication and financial difficulties**

Of the 368 PWE who reported the use of ASM, 72 (19.6%) had difficulties to obtain ASM during the COVID-19 pandemic. The reasons for this included: ASM were not available (n = 50/72; 69.4%), COVID-19-related mobility restrictions made a visit to the hospital/pharmacy impossible (n = 9/72; 12.5%), financial problems as a result of COVID-19-related unemployment (n = 9/72; 12.5%), or other reasons, such as difficulties to obtain new prescriptions (n = 3/72; 4.2%).

Seventy-eight (22.8%) of 342 PWE who answered the question reported financial problems. Of these, 61 (78.2%) reported difficulties to pay for housing/bills, 27 (34.6%) difficulties to eat properly, and 30 (38.5%) difficulties to pay for ASM. Compared with PWE without financial problems, PWE with financial problems were more often living in a low- to middle-income country (35.5% vs 14.8%; p < 0.001).

**3.5. Anxiety and depression among PWE**

Of the 399 PWE, 201 (50.4%) screened positive for anxiety according to the HADS, and 159 (39.8%) and 187 (46.9%) PWE screened positive for depression based on the HADS and PHQ-9 scale, respectively.

Multiple logistic regression analysis suggested that female gender and financial problems significantly increased the odds of screening positive for both anxiety and depression. Indeed, of the 66 female PWE, 51 (77.5%) screened positive for anxiety and 44 (66.7%) for depression. In addition, PWE residing in high-income countries had significantly lower odds (0.360, 95% confidence interval [CI]: 0.191–0.682) for anxiety. Importantly, PWE who reported difficulties to access ASM had a higher odds (1.885, 95% CI: 1.083–3.282) for depression.

**4. Discussion**

In this multicountry study, we assessed the access to healthcare and self-reported psychiatric symptoms among PWE during the ongoing COVID-19 pandemic.
COVID-19 pandemic. We detected important problems to access ASM and high prevalence of anxiety and depression. During confinement, follow-up consultations with the treating physician were canceled in almost half of the cases.

Of the 36 PWE (10% of the study population) who were tested for COVID-19, nine reported positive test results, accounting for 2.5% of the total population. However, when we applied the WHO clinical diagnostic criteria on our population, eight times more PWE were classified as cases with suspected COVID-19, suggesting that our participants were being undertested. The COVID-19 infections were asymptomatic in approximately half of the infected PWE, which is in line with previous results [31].

We found that the ongoing COVID-19 pandemic had an impact on the access to ASM. Nearly one-fifth of the PWE on ASM reported problems to obtain ASM; the most frequent constraint to ASM, accounting for approximately 70% of cases, was a shortage of ASM, which is an important problem that is not necessarily related to COVID-19. Our survey does not allow to determine whether COVID-19-related restrictions led to increased problems of ASM availability. Our results, however, do suggest an additional impact of COVID-19 on access to ASM. One-fourth of the PWE with difficulties to obtain ASM indicated COVID-19-related causes such as financial challenges to buy ASM as a consequence of COVID-19-related unemployment (12.5%) and COVID-19-related mobility restrictions making it difficult to go to the hospital/pharmacy for ASM refill (12.5%). Special attention should be paid by governments and healthcare workers to the impact of COVID-19 restrictions on access to ASM. In a longitudinal study conducted in the United States in 2016, ASM shortage caused psychological problems in the PWE related to the fear and anxiety of not being able to get their ASM and the possible consequences thereof [32]. In our study, we observed an increased risk of depression in PWE who reported problems to access ASM. We did not detect an increase in seizure frequency since the start of the COVID-19 pandemic. However, we conducted the survey early during confinement, and a potential impact in case of continuing access problems cannot be excluded.

In our survey, 201/399 (50.4%) and 159/399 (39.8%) PWE scored positive for anxiety and depression, respectively. The prevalence of depression and anxiety in PWE is higher than in the general population [4,20]. In a meta-analysis of 51 cross-sectional studies among PWE performed between 1999 and 2018, the prevalence of depression varied between 5% and 85%, largely depending on the diagnostic criteria that were used [33]. In 8 studies that used the HADS—Depression score, the point prevalence varied between 5% and 43%. Given the differences in study design, it is difficult to know whether COVID-19 increases the rates of depression among PWE [33]. The same is true for anxiety, for which a wide variation of screening tools and ranges of prevalence (9%-45%) have been reported in the literature [34-37]. It is, however, worth noting that the prevalence of depression reported in our study seems to be higher than what has been reported by other authors when investigating the mental health condition of the general population during the COVID-19 pandemic [12-14]. Telehealth appointments may help to timely identify anxiety and depression. Several recent studies have highlighted the feasibility of telehealth appointments in epilepsy care and suggest an added value of interprofessional collaboration [38-41].

Our study has limitations. First, the design was cross-sectional, which largely prohibited the investigation of causal relationships. Secondly, our survey was web-based, which causes an important sampling bias as people who are more active on social media were more prone to respond and low- to middle-income countries may not have equal access to the internet. Also, people with anxiety and depression may be more prone to complete online questionnaires, which may have resulted in an overestimation of the prevalence of depression and anxiety. Studies with a different design are needed to confirm our results. The link to complete the survey was freely accessible to anyone, making it difficult to estimate the reached sample [42]. Nonetheless, our sample size was small, and only 16.8% of PWE were from low-income countries, prohibiting an in-depth study of relationships with demographic features. In addition, the diagnosis of epilepsy was self-reported. As such, some respondents may not be true PWE according to ILAE diagnostic criteria. As we did not collect information on region of origin, we cannot relate individual responses directly to the actual lockdown and healthcare situation at the time of response. An additional limitation was related to the cancelation of follow-up visits: we do not know whether these were canceled by PWE themselves or by the healthcare providers. Four of the 9 (44.4%) PWE with a positive COVID-19 test reported no symptoms. However, we did not ask why the COVID-19 test was done. Those without symptoms may have been tested because of a contact with a person with a confirmed COVID-19 infection and may have developed symptoms since they completed the survey. Lastly, as more than 40% of the responders reported a seizure during the last 3 months, one would suspect that the COVID-19 epidemic increased the frequency of the seizures. Indeed, an increase of frequency of seizures has been observed with anxiety, depression, and healthcare barriers [43]. A prospective study certainly is needed to investigate the effect of the COVID-19 epidemic and lockdown measures on the frequency of seizures of PWE. Notwithstanding, our study provides important information on the impact of COVID-19 on PWE, which should be taken into account to improve their quality of life.

In conclusion, during the COVID-19 pandemic, PWE are confronted with challenges to access ASM. Half of the PWE screened positive for
depression and slightly less screened positive for anxiety. Almost half of the respondents had their follow-up consultation with the treating physician canceled. While more research is needed, our results call for innovative approaches to ensure continuity in access to ASM. Physicians should ensure continued follow-up, either through in-person or telehealth appointments, to timely identify symptoms of anxiety and depression and act accordingly.

Supplementary data to this article can be found online at https://doi.org/10.1016/j.yebeh.2020.107350.

Declaration of Competing Interest

All authors disclose no conflict of interest related to this study.

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