Incidence and Severity of Depression Among Recovered African Americans with COVID-19-Associated Respiratory Failure

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Abstract

Background Coronavirus disease (COVID-19) disproportionately affects African Americans, and they tend to experience more severe course and adverse outcomes. Using a simple and validated instrument of depression screening, we evaluated the incidence and severity of major depression among African American patients within 90 days of recovery from severe COVID-19-associated respiratory failure.

Methods African American patients hospitalized and treated with invasive mechanical ventilation for COVID-19-associated respiratory failure in the intensive care unit (ICU) of Grady Memorial Hospital, Atlanta, between April 1, 2020, and June 30, 2020, were screened for depression within 90 days of hospital discharge using the validated patient health questionnaires (PHQ-2) and PHQ-9.

Results A total of 73 patients completed the questionnaire. The median age was 52.5 years [IQR 44–65] and 65% were males. The most common comorbidities were hypertension (66%) and diabetes mellitus (51%). Forty-four percent of the patients had a diagnosis of major depressive disorder (MDD) based on their PHQ-9 questionnaire responses. The incidence of MDD was higher among females (69%, n=18/26) compared to males (29%, n=14/47), in patients > 75 years (66%) and those with multiple comorbidities (45%). Eighteen percent of the patients had moderate depression, while 15% and 22% had moderately severe and severe depression, respectively. Only 26% (n=7/27) of eligible patients were receiving treatment for depression at the time of this survey.

Conclusion The incidence of depression in a cohort of African American patients without prior psychiatric conditions who recovered from severe COVID-19 infection was 44%. More than 70% of these patients were not receiving treatment for depression.

Keywords African Americans • COVID-19 • Depression • ICU • PHQ-9

Introduction

The mental health impact of COVID-19 infection from the novel severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) on its survivors has been reported in the literature [1]. According to surveys conducted during the ongoing COVID-19 pandemic, patients often suffer from anxiety, post-traumatic stress disorder, and trouble sleeping after recovery from COVID-19 infection [2–4].

COVID-19 disproportionately affects African Americans, and they tend to experience more severe course and adverse outcomes [5, 6]. Also, they are more likely to have undiagnosed and untreated major depression which can negatively impact daily functioning and lead to a poor quality of life [7, 8]. Currently, there is paucity of data on the incidence of depression among African Americans who have recovered from COVID-19 infection particularly those who had severe infection requiring intensive care unit (ICU) hospitalization.

Using a simple and validated instrument of depression screening, we evaluated the incidence and severity of major depression among African American patients within 90 days of recovery from severe COVID-19 infection with associated respiratory failure.
Methods

The study was approved by the Morehouse School of Medicine institutional review board, and verbal informed consent was obtained from the study participants. We identified African American patients with confirmed severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), the virus that causes COVID-19 infection, on polymerase chain reaction testing of a nasopharyngeal or tracheal sample. Those who were hospitalized and treated with invasive mechanical ventilation for respiratory failure at the Grady Memorial Hospital ICU, Atlanta, Georgia, between April 1, 2020, and June 30, 2020, were included in the study. Patients who were still hospitalized by June 30, 2020, and those with previous history of psychiatric diagnosis including major depression were excluded. Data on sociodemographics and comorbidities were extracted from the electronic health record system (Epic 2019 version).

The Patient Health Questionnaire (PHQ)-2 and PHQ-9 (supplementary material) are used to reliably assess for depression and its severity in the outpatient and other medical care settings [9]. The PHQ-9 is a nine-item questionnaire, and the PHQ-2 includes the first two items of the PHQ-9. The PHQ-2 evaluates the frequency of depressed mood and anhedonia over the past 2 weeks as a “first step” in the screening process. The PHQ-9 is then administered to patients if the PHQ-2 is positive for depression to determine whether they meet criteria for a depressive disorder [9–11]. The PHQ-2 has a 97% sensitivity and 67% specificity in adults, whereas the PHQ-9 has a 61% sensitivity and 94% specificity in adults [9].

A team of two physicians trained on the depression screening tools administered the PHQ-9 over the phone to the study patients within 30 to 90 days of discharge from the hospital. Verbal informed consent was obtained from the patients prior to administering the questionnaire. Patients who responded “yes” to at least 5 questions in the two right columns (one of which corresponds to question #1 or #2) of the PHQ-9 questionnaire were considered to have an initial diagnosis of major depressive disorder (MDD) [10, 11]. Other depression is diagnosed if there are 2–4 yes answers in the right columns of the PHQ-9 (one of which corresponds to question #1 or #2) [11].

To assess depression severity, all patients were categorized into none or minimal, mild, moderate, moderately severe, and severe based on their severity scores on a scale of 0–27 [10, 11]. Using descriptive analysis, we determined the association between the severity of depression and the ICU length of stay (LOS) and discharge disposition. Lastly, we determined the number of patients on treatment for depression as a percentage of the total eligible for treatment based on their PHQ-9 scores. Summary statistics were reported as medians and proportions for various sociodemographic and clinical characteristics of patients. Due to limited sample size and descriptive nature of the study, analysis for statistical significance was performed. Data analysis was performed using R version 3.6.3 (R Foundation).

Results

A total of 91 patients met the inclusive criteria, and 73 completed the questionnaire (response rate of 80%). Table 1 describes the sociodemographic and clinical characteristics of the patients with their PHQ-9 score distribution. The median age of the participants was 52.5 years [IQR 44–65]. There were more male participants than females (65% vs. 35%). Eighty-five percent of the patients had more than 1 comorbidity, and the most common comorbidities were hypertension (66%) and diabetes mellitus (51%). Forty-four percent of the patients had a diagnosis of MDD based on their responses on the PHQ-9 questionnaire. The incidence of MDD was higher among females (69%, n=18/26) compared to males (29%, n=14/47). Sixty percent (n=18/30) of patients aged 55–74 years, and 66% (n=4/6) of patients > 75 years had PHQ-9 scores consistent with MDD. Among patients with multiple comorbidities, 45% (n=28/62) had a diagnosis of MDD compared to 36% (n=4/11) in patients with only one or no comorbidity. Comorbidities with the highest proportion of patients MDD diagnosis were cerebrovascular accident (62.5%), obstructive sleep apnea (60%), solid tumors (57.5%), hypertension (58.3%), and diabetes mellitus (51.4%). Eighteen percent of the patients had moderate depression, while 15% and 22% had moderately severe and severe depression, respectively.

The median LOS of patients in the ICU was 10 days [IQR 4–18.5]. Among patients who spent 10 days or more in the ICU, 58% had a PHQ-9 score reflecting MDD compared to 32.5% in those who spent less than 10 days. Seventy-five percent of the patients were discharged home, out of which 38% had a diagnosis of MDD (Table 2). Though only 8% of the patients were discharged to a long-term acute care facility (LTAC), MDD diagnosis was disproportionately higher (67%, n=4/6) relative to those discharged home (38.2%) or skilled nursing facilities (SNF) (58.3%).

Figure 1 shows that 37% (n=27/73) of patients in the study had a PHQ-9 score of at least 14 and were therefore eligible for depression treatment using antidepressants, psychotherapy, or a combination of both [10]. However, only 26% (n=7/27) were on some form of depression treatment at the time of this survey. Two of the patients were on cognitive behavioral therapy (CBT), while 5 patients were treated with a combination of CBT and antidepressant medications—selective serotonin receptor inhibitors (SSRIs). The remaining 74% said they had not been screened for depression since discharge from the hospital. Though the research team at the time of the survey implored all the study patients to follow up with their primary care
**Table 1** Characteristics of patients and distribution of their PHQ-9 scores

| Variables                      | Total (%) | Major depressive disorder (%) | Other depression (%) | Depression diagnosis | Depression Severity |
|--------------------------------|-----------|-------------------------------|----------------------|----------------------|---------------------|
|                                |           |                               |                      | None–minimal depression PHQ-9 Score 0–4 (%) | Mild depression PHQ-9 Score 5–9 (%) | Moderate depression PHQ-9 Score 10–14 (%) | Moderately Severe depression PHQ-9 Score 15–19 (%) | Severe depression PHQ-9 Score 20–27 (%) |
| Total no. (%)                  | 73 (100)  | 32 (43.8)                     | 41 (56.2)            | 22 (30.1)            | 11 (15.1)           | 13 (17.8)          | 11 (15.1)           | 16 (21.9)            |
| Age (median 52.5y [IQR 44–65]) |           |                               |                      |                      |                     |                     |                     |                      |
| < 55 years                     | 37 (50.7) | 10 (27.1)                     | 27 (73)              | 10 (27)              | 8 (21.6)            | 7 (18.9)           | 5 (13.5)            | 7 (18.9)            |
| 55–74 years                    | 30 (41.1) | 18 (60)                       | 12 (40)              | 10 (33.3)            | 1 (3.3)             | 6 (20)             | 6 (20)              | 7 (23.3)            |
| >75 years                      | 6 (8.2)   | 4 (66)                        | 2 (33.3)             | 2 (33.3)             | 2 (33.3)            | 0 (0)              | 0 (0)               | 2 (33.3)            |
| Gender                         |           |                               |                      |                      |                     |                     |                     |                      |
| Female                         | 26 (35.6) | 18 (69.2)                     | 8 (30.8)             | 9 (34.6)             | 4 (15.4)            | 4 (15.4)           | 3 (11.5)            | 6 (23.1)            |
| Male                           | 47 (64.4) | 14 (29.7)                     | 33 (70.2)            | 13 (27.7)            | 7 (14.9)            | 9 (19.1)           | 8 (17)              | 10 (21.3)           |
| Health insurance               |           |                               |                      |                      |                     |                     |                     |                      |
| Medicaid/Medicare              | 24 (32.9) | 14 (58.3)                     | 10 (41.7)            | 8 (33.3)             | 2 (8.3)             | 3 (12.5)           | 5 (20.8)            | 6 (25)              |
| Private insurance/self-pay     | 21 (28.8) | 8 (38.1)                      | 13 (61.9)            | 7 (33.3)             | 3 (14.3)            | 5 (23.8)           | 3 (14.3)            | 3 (14.3)            |
| Uninsured                      | 28 (38.4) | 10 (35.7)                     | 18 (64.3)            | 7 (25)               | 6 (21.4)            | 5 (17.9)           | 3 (10.7)            | 7 (25)              |
| Comorbid diseases              |           |                               |                      |                      |                     |                     |                     |                      |
| Asthma                         | 9 (12.3)  | 3 (33.3)                      | 6 (66.7)             | 2 (22.2)             | 1 (11.1)            | 3 (33.3)           | 2 (22.2)            | 1 (11.1)            |
| Coronary artery disease (CAD)  | 11 (15.1) | 3 (27.7)                      | 8 (72.7)             | 5 (45.5)             | 1 (9.1)             | 0 (0)              | 1 (9.1)             | 4 (36.4)            |
| Cancer                         | 7 (9.6)   | 4 (57.4)                      | 3 (42.9)             | 1 (14.3)             | 0 (0)               | 0 (0)              | 2 (28.6)            | 4 (57.1)            |
| CHF (congestive heart failure) | 19 (26)   | 8 (42.1)                      | 11 (57.9)            | 5 (26.3)             | 4 (21.1)            | 1 (5.3)            | 6 (31.6)            | 3 (15.8)            |
| Chronic kidney disease (CKD)   | 8 (11)    | 3 (37.5)                      | 5 (62.5)             | 0 (0)                | 1 (12.5)            | 0 (0)              | 2 (25)              | 5 (62.5)            |
| Chronic obstructive pulmonary  | 11 (15.1) | 3 (27.2)                      | 8 (72.7)             | 4 (36.4)             | 1 (9.1)             | 2 (18.2)           | 2 (18.2)            | 2 (18.2)            |
| Cerebrovascular accident       | 8 (11)    | 5 (62.5)                      | 3 (37.5)             | 2 (25)               | 1 (12.5)            | 0 (0)              | 3 (37.5)            | 2 (25)              |
| Diabetes mellitus (DM)         | 37 (50.7) | 19 (51.4)                     | 18 (48.6)            | 10 (27)              | 6 (16.2)            | 6 (16.2)           | 7 (18.9)            | 8 (21.6)            |
| HIV                            | 4 (5.5)   | 1 (25)                        | 3 (75)               | 0 (0)                | 0 (0)               | 2 (50)             | 2 (50)              | 0 (0)               |
| Hypertension                   | 48 (65.8) | 28 (58.3)                     | 20 (41.7)            | 13 (27.1)            | 7 (14.6)            | 7 (14.6)           | 9 (18.8)            | 12 (25)             |
| Obstructive sleep apnea (OSA)  | 10 (13.7) | 6 (60)                        | 4 (40)               | 1 (10)               | 1 (10)              | 5 (50)             | 2 (20)              | 1 (10)              |
| Body mass index (BMI)          |           |                               |                      |                      |                     |                     |                     |                      |
| < 30 kg/m²                      | 45 (61.6) | 19 (42.2)                     | 26 (57.8)            | 8 (17.8)             | 9 (20)              | 8 (17.8)           | 10 (22.2)           | 10 (22.2)           |
| ≥ 30 kg/m²                     | 28 (38.4) | 13 (46.4)                     | 15 (53.6)            | 14 (50)              | 2 (7.1)             | 5 (17.9)           | 1 (3.6)             | 6 (21.4)            |
| Tobacco smoking                |           |                               |                      |                      |                     |                     |                     |                      |
providers to be screened for depression and treatment, they particularly emphasized the need for urgent evaluation in patients with PHQ-9 score of at least 14. No patient expressed homicidal or suicidal ideations (item 9 on the PHQ-9 questionnaire).

### Discussion

Our study showed that at least 4 out of 10 African Americans without prior psychiatric illness, who were hospitalized in the ICU and received invasive mechanical ventilation for severe COVID-19 infection, will develop MDD within 90 days of discharge from the hospital. Notably, patients older than 55 years, females, patients with multiple pre-existing comorbidities, and those who spent 10 days or more in the ICU had higher rates of MDD diagnosis. Depression is a component of the post-intensive care syndrome (PICS) and is common in ICU survivors. Studies have reported depression rates as high as 45% in patients with respiratory failure which is similar to what we found [12]. However, the rate of depression reported in patients who recovered from COVID-19 infection was lower [1, 3].

Psychiatric complications of SARS-CoV-2 infection have been attributed to the immune response to the virus itself, lingering clinical symptoms, and/or psychosocial stressors from social isolation, concerns about infecting others, and social stigma [3]. It is important to point out that African Americans and social stigmas from social isolation, concerns about infecting others, and social stigma [3]. It is important to point out that African Americans have higher rates of MDD diagnosis, and those who spent 10 days or more in the ICU had also contributed to the higher rate of depression in this study [13].

Expectedly, patients discharged to LTAC facilities and SNF had disproportionately higher incidence of MDD compared to those discharged home. Patients discharged to extended care facilities (SNF and LTACs) are traditionally older, sicker, and more likely to have multiple comorbidities which are risk factors for MDD [14, 15]. Studies show that up to 35% of residents in long-term care facilities may experience major depression [15]. However, depressive symptoms in patients in these facilities are often not recognized by healthcare providers [16]. Given that a significant proportion of African Americans have a significant proportion of African Americans have expressed homicidal or suicidal ideations (item 9 on the PHQ-9 questionnaire)
undiagnosed depression and a relatively smaller proportion of those diagnosed with MDD go on to receive treatment compared to other ethnicities in the USA [8, 17]. Additionally, African Americans and other racial minorities have less access to mental health services compared to whites and are more likely to receive poor quality care when treated [18]. This disparity in mental healthcare services is a deficit of the healthcare system that needs to be addressed.

Depression underdiagnosis and undertreatment among African Americans is rooted in societal stigma associated with mental illnesses, inadequate psychosocial support system, lack of information, and mistrust of the healthcare system [8]. Our study further underscores the need to prioritize the mental health of COVID-19 survivors within the African American community and ensure timely and appropriated linkage to treatment services and interventions. The PHQ-9 questionnaire is sensitive in diagnosing MDD and easy to administer [10, 11, 19–22]. We recommend that medical providers utilize the PHQ-9 to screen COVID-19 survivors for depression during their post-ICU follow-up visits and manage appropriately.

Our study has some limitations. It is cross-sectional in nature, and causality cannot be implied. Also, the findings may not be generalizable to other ethnic groups. The sample size is relatively small, and this is a single center study; therefore, larger and multicenter prospective studies may be needed to further evaluate contextual factors that likely predispose COVID-19 patients to new psychiatric conditions.

**Conclusion**

The incidence of depression in a cohort of African American patients without prior psychiatric conditions who recovered from severe COVID-19 infection was 44%. More than 70% of these patients were not receiving treatment for depression. Given the significantly high burden of COVID-19 infection in the African American community and the debilitating effect of depression, it is important for clinicians to assess for depression during post-ICU discharge follow-up for severe COVID-19 infection–associated respiratory failure. Additionally, public health efforts should be intensified to help African Americans.
Americans recognize the features of depression after COVID-19 infection and seek medical evaluation early.

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**Author Contribution** All authors contributed to the writing of the manuscript.

**Declarations**

**Ethics Approval** This study was approved by the Morehouse School of Medicine Institutional Review Board (IRB).

**Conflict of interest** The authors declare no competing interests.

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