Post-acute COVID-19 Syndrome Negatively Impacts Physical Function, Cognitive Function, Health-Related Quality of Life, and Participation

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Objective: This report describes persistent symptoms associated with post-acute COVID-19 syndrome (PACS) and the impact of these symptoms on physical function, cognitive function, health-related quality of life, and participation.

Design: This study used a cross-sectional observational study design. Patients attending Mount Sinai’s post-acute COVID-19 syndrome clinic completed surveys containing patient-reported outcomes.

Results: A total of 156 patients completed the survey, at a median (range) time of 351 days (82–457 days) after COVID-19 infection. All patients were prevaccination. The most common persistent symptoms reported were fatigue (n = 128, 82%), brain fog (n = 105, 67%), and headache (n = 94, 60%). The most common triggers of symptom exacerbation were physical exertion (n = 134, 86%), stress (n = 107, 69%), and dehydraton (n = 77, 49%). Increased levels of fatigue (Fatigue Severity Scale) and dyspnea (Medical Research Council) were reported, alongside reductions in levels of regularly completed physical activity. Ninety-eight patients (63%) scored for at least mild cognitive impairment (Neuro-Qol), and the domain of the EuroQol: 5 dimension, 5 level most impacted was Self-care, Anxiety/Depression and Usual Activities.

Conclusions: Persistent symptoms associated with post-acute COVID-19 syndrome seem to impact physical and cognitive function, health-related quality of life, and participation in society. More research is needed to further clarify the relationship between COVID-19 infection and post-acute COVID-19 syndrome symptoms, the underlying mechanisms, and treatment options.

Key Words: Post-acute COVID-19, Fatigue, Cognition, Employment, Quality of life

(Am J Phys Med Rehabil 2022;101:48–52)

After the dramatic influx of patients with persistent, debilitating symptoms after acute SARS-CoV-2 (COVID-19) infection, the National Institutes of Health announced an initiative to fully investigate the post-acute sequelae of COVID-19 (post-acute COVID-19 syndrome [PASC]). Post-acute COVID-19 syndrome can take many forms, from post-intensive care unit syndrome\(^1\) to pulmonary fibrosis secondary to aggressive COVID-19 pneumonia.\(^2\) However, PACS (also known as long COVID) is one of the most troubling manifestations of PASC that has been reported to date. It is characterized by persistent symptoms that are still present at least 4 wks after initial infection and often lasting for several months.\(^3\) Despite the highly debilitating nature of PACS, the long-lasting symptoms often occur in the absence of severe acute infection, medically explainable physical symptoms, or preexisting comorbidities.\(^4\)–\(^6\)

Several studies have documented the most common persistent symptoms after severe COVID-19 infection. These symptoms include fatigue, dyspnea, “brain fog”\(^7\) various cognitive symptoms, pain, anxiety, depression, and gastrointestinal issues.\(^3\)–\(^6\)–\(^9\) In these cohorts, the symptoms arising from COVID-19 increased disability and negatively impacted physical function and quality of life\(^7\) and affected participation in general life activities and the ability to work.\(^9\) There is a critical need to classify the prevalence of specific persistent symptoms.
that follow acute COVID-19 infection and the impact of these symptoms on patient-reported outcomes that are well validated in other conditions. This will facilitate the establishment of diagnostic criteria for PACS and accurate tracking of responses to various prospective therapies.

It has been hypothesized that persistent symptoms after acute COVID-19 infection result from an immune-mediated disruption to the autonomic nervous system.10,11 Similar to other postviral autoimmune conditions (such as Guillain-Barré syndrome), COVID-19 infection seems to act as an immune trigger.12 This immune response, coupled with a lack of access to acute COVID-19 treatments offered only in a hospital setting, may explain why even those with less severe acute infection are still experiencing persistent symptoms.

It is clear that in the wake of the COVID-19 pandemic, a second, longer-term public health emergency has emerged. It is imperative to understand the burden of this novel condition with millions Americans at risk of developing PACS by the end of the pandemic. This study describes the persistent symptoms reported by a cohort of patients with PACS, the majority of whom were infected with COVID-19 in early 2020 and not hospitalized. The impact of these symptoms on physical function, cognitive function, health-related quality of life, and participation is also reported.

### METHODS

This was a retrospective observational study of patients attending Mount Sinai’s PACS clinic. Approval for publication was provided and requirement for patient consent was waived by the Mount Sinai Program for Protection of Human Subjects (IRB 21-01147). This study conforms to all strengthening the reporting of observational studies in epidemiology guidelines and reports the required information accordingly (see Supplementary Checklist, Supplemental Digital Content 1, http://links.lww.com/PHM/B417).

#### Participants

This was a convenience sample exploring symptom characteristics of patients attending the Mount Sinai’s PACS clinic. This is an interdisciplinary clinic consisting of physicians, physical therapists, and dietitians. Patients were either referred by a physician or self-referred. All patients had confirmed (by polymerase chain reaction [PCR] and/or antibody test) or probable (diagnosed by a medical doctor in accordance with World Health Organization recommendations13) previous COVID-19 infection and diagnosis of PACS (defined as experiencing symptoms >12 wks since initial symptom onset). Inclusion criteria for the present study were attending the Mount Sinai’s PACS clinic between March 2020 and March 2021 and completion of the patient-reported outcome survey.14 There were no exclusion criteria.

#### Data Collection and Outcomes

Data were collected using REDCap (Research Electronic Data Capture) electronic data capture tools hosted at Mount Sinai Health System. REDCap is a secure, web-based application designed to support data capture for research studies. Participants were provided with a link to the survey via e-mail on March 14, 2021, as part of their clinical care.

Baseline demographic data included sex, age, body mass index, race, and comorbidities. COVID-19 clinical data included duration of COVID-19 symptoms (at survey completion), PCR (obtained from nasopharyngeal swab), and antibody test completion and results, need for hospitalization at time of COVID-19 infection, and vaccination status.

Patient-reported outcomes included current persistent symptoms and triggers of symptom exacerbation, and screening tools for fatigue (Fatigue Severity Scale, Fatigue Visual Analog Scale), breathlessness (Medical Research Council Breathlessness Scale), completion of regular moderate and vigorous intensity physical activity (author developed), cognitive function (Neuro-Qol), health-related quality of life (EuroQol: 5 dimension, 5 level [EQ-5D-5L]), anxiety (generalized anxiety disorder scale [GAD-7]), depression (patient health questionnaire-2 [PHQ-2]), disability (World Health Organization Disability Assessment Schedule), and pre- and post-COVID-19 employment status (author developed).

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**TABLE 1. Patient (N = 156) baseline demographic and COVID-19-related data**

|                        | All Patients (N = 156) | Confirmed COVID-19 (n = 87) | Presumed COVID-19 (n = 69) |
|------------------------|------------------------|-----------------------------|-----------------------------|
| Female                 | 107 (69)               | 54 (62)                     | 53 (77)                     |
| Age, median (range), yr| 44 (13–79)             | 45 (13–79)                  | 44 (14–79)                  |
| BMI, median (range), kg/m² | 24 (16–52)         | 24 (17–52)                  | 24 (16–42)                  |
| Race                   |                        |                             |                             |
| White                  | 119 (76)               | 65 (75)                     | 54 (78)                     |
| Asian                  | 8 (5)                  | 3 (3)                       | 5 (7)                       |
| Black or African American | 6 (4)                 | 4 (5)                       | 2 (3)                       |
| American Indian or American Native | 2 (1) | 1 (1)                  | 1 (1)                       |
| Native Hawaiian or Pacific Islander | 0 (0) | 0 (0)                  | 0 (0)                       |
| Other                  | 15 (10)                | 8 (9)                       | 7 (10)                      |
| Hispanic or Latinx     | 10 (7)                 | 3 (4)                       | 7 (10)                      |
| Duration of symptoms, median (range), d | 351 (82–457) | 350 (157–424) | 355 (82–457) |
| PCR completed          | 98 (63)                | 57 (66)                     | 41 (59)                     |
| PCR positive           | 34 (22)                | 34 (39)                     | 0 (0)                       |
| Antibody test completed| 149 (96)               | 86 (99)                     | 63 (91)                     |
| Antibody positive      | 80 (51)                | 80 (92)                     | 0 (0)                       |
| PCR and/or antibody positive | 87 (56)          | 87 (100)                    | 0 (0)                       |
| Hospitalized for COVID-19 | 17 (11)            | 16 (18)                     | 1 (1)                       |
| Received COVID-19 vaccination | 87 (56)          | 45 (52)                     | 42 (61)                     |
| Most prevalent comorbidities |           |                             |                             |
| Cancer (any type)      | 30 (20)                | 10 (11)                     | 20 (29)                     |
| Asthma                 | 30 (20)                | 13 (15)                     | 17 (25)                     |
| Anxiety                | 18 (12)                | 12 (14)                     | 6 (9)                       |
| Depression             | 13 (8)                 | 8 (9)                       | 5 (7)                       |
| Hypertension           | 11 (7)                 | 7 (8)                       | 4 (6)                       |

Data are presented as n (%) unless otherwise indicated.

*All COVID-19 vaccination occurred after COVID-19 infection.

BMI, body mass index.

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Statistical Analyses

Statistical analyses were undertaken with Stata (Stata Statistical Software Release: V14; StataCorp). Data were analyzed using descriptive statistics and reported using number and percentage, or median and range.

RESULTS

The survey was sent to 386 patients, with 156 (48%) responding. The median (range) time to follow-up time since the onset of COVID-19 infection was 351 days (82–457 days; Table 1). The most common symptoms reported were fatigue \((n = 128, 82\%)\), brain fog \((n = 105, 67\%)\), headache \((n = 94, 60\%)\), sleep disturbance \((n = 92, 59\%)\), and dizziness \((n = 85, 54\%)\); Fig. 1). The most common triggers of symptom exacerbation reported were physical exertion \((n = 134, 86\%)\), stress \((n = 107, 69\%)\), dehydration \((n = 77, 49\%)\), weather changes \((n = 58, 37\%)\), consuming large meals \((n = 44, 28\%)\), premenstrual period \((n = 34, 22\%)\), and alcohol consumption \((n = 34, 22\%)\).

The median (range) Fatigue Severity Scale average score was 5.6 (1–7) of 7, with 122 patients (78%) reporting an Fatigue Severity Scale average score of 4 or greater, indicating problematic fatigue. Sixty-three patients (40%) reported a score of 3 or more (of 5) on the Medical Research Council Breathlessness Scale, suggesting moderate to severe disability due to dyspnea. When compared with pre–COVID-19 infection levels, patients were completing 150 mins/wk of physical activity less frequently after COVID-19 infection, when asked separately about moderate and vigorous intensities (Fig. 2).

Ninety-eight patients (63%) scored for at least mild cognitive impairment on the Neuro-Qol (Fig. 3). The domains of the EQ-5D-5L impacted the most (reported as slight problems or greater) were self-care, anxiety/depression, and usual activities (Table 2). The median (range) EQ-5D-5L Visual Analog Scale score was 64 (6–99) of 100, with a higher score indicating greater health-related quality of life. Twenty-nine patients

![Figure 1](link)

**FIGURE 1.** Most commonly reported persistent symptoms by all patients \((N = 156)\).

![Figure 2](link)

**FIGURE 2.** Levels of moderate (A) and vigorous (B) intensity physical activity regularly completed (150 mins/wk) before and after COVID-19 infection in all patients \((N = 156)\).
(19%) scored 10 or greater on the GAD-7, indicating possible anxiety disorder. Forty-three patients (28%) scored 3 or greater on the PHQ-2, indicating possible major depressive disorder. The median (range) World Health Organization Disability Assessment Schedule total score was 14 (0–44) of 100. A total of 134 patients (86%) answered pre– and post–COVID-19 employment questions; the number of patients in full-time work reduced from 102 (76%) pre–COVID-19 to 55 (41%) at the time of follow-up (Fig. 4).

DISCUSSION

This observational study of a cohort of patients with PACS reported that COVID-19–related symptoms are persistent for at least 2 mos, and often longer than 12 mos, with fatigue, brain fog, sleep disturbance, dizziness, dyspnea, memory loss, and palpitations being identified as the most common. The most common triggers of symptom exacerbation in this cohort were physical exertion, stress, and dehydration. The negative impact of PACS on a variety of patient-reported outcomes has been demonstrated. Just less than 50% of the patients included in this study had a negative PCR test or tested seronegative for antibodies. Issues relating to false-negative rates are well documented with PCR testing. Similarly, there is literature to support the idea that antibody levels in patients who experienced less severe acute infection tend to fade rapidly. In acknowledgement of the potential for health disparities to rapidly arise from an overreliance on COVID-19 tests, the Centers for Disease Control and Prevention has recently recommended against using seropositive status as the sole diagnostic criteria for any postacute sequelae of COVID-19. To fully understand all of the possible presentations of PACS, studies must incorporate and report data from both seronegative and seropositive patient populations.

The most common symptoms observed in this cohort are consistent with those previously reported. The pattern of PACS symptoms resembles other postviral syndromes, including dysautonomia, and postural orthostatic tachycardia syndrome and myalgic encephalomyelitis. It is unsurprising that physical exertion was the most common cause of symptom exacerbation, as this is a feature shared by some of these conditions. The potential for the worsening of symptoms after physical exertion is the most important consideration when prescribing rehabilitation therapies for people with PACS.

The presence of cognitive dysfunction in more than half (63%) of patients, in combination with reduced usual activities and self-care scores on the EQ-5D-5L, highlights that patients with PACS may have a reduced ability to participate in society. Employment was also impacted in most patients; however, it is difficult to determine whether this was specifically due to disability related to COVID-19 infection or possibly to the broader implications of the pandemic on the ability for workplaces to operate as usual. Levels of self-reported physical activity were greatly reduced, likely coinciding with the potential for symptom exacerbation; this raises concerns given the known longer-term health risks of physical inactivity.

The proportion of participants reporting anxiety (19%) and depression (28%) on the PHQ-2 and GAD-7 was slightly higher than reported as part of their medical history. With a lack of pre–COVID-19 PHQ-2 and GAD-7 data available, it is difficult to make conclusions about the impact of PACS on anxiety and depression.

Some limitations of the present study include the use of clinical survey data answered in retrospect, a lack of comparison group, and/or pre–COVID-19 measures. In addition, the
FIGURE 4. Employment status before and after COVID-19 infection in participants who answered employment questions (n = 134).

use of noncondition-specific patient-reported outcomes can increase the risk of inaccuracy and recall bias.

CONCLUSIONS

The presence of persistent symptoms associated with PACS seems to impact physical and cognitive function, health-related quality of life, and participation in society. The data reported contribute to the recognition and research of long COVID as recommended by the World Health Organization [32] and will help inform future rehabilitation strategies.

ACKNOWLEDGMENTS

The authors thank the patients in the study, the frontline healthcare workers at Mount Sinai Health System, and the wider research team at the Abilities Research Center and the Center for Post-COVID Care at Mount Sinai. The authors also thank the RTW Charitable Foundation for supporting multiple PACS initiatives in the Abilities Research Center.

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