JAMA Open.

Original Investigation | Infectious Diseases

Perceived Cognitive Deficits in Patients With Symptomatic SARS-CoV-2 and Their Association With Post-COVID-19 Condition

Teresa C. Liu, MD, MPH; Sun M. Yoo, MD, MPH; Myung S. Sim, DrPH; Yash Motwani, BS; Nisha Viswanathan, MD; Neil S. Wenger, MD, MPH

Abstract

IMPORTANCE Neuropsychiatric symptoms are common in acute SARS-CoV-2 infection and in post-COVID-19 condition (PCC; colloquially known as long COVID), but the association between early presenting neuropsychiatric symptoms and PCC is unknown.

OBJECTIVE To describe the characteristics of patients with perceived cognitive deficits within the first 4 weeks of SARS-CoV-2 infection and the association of those deficits with PCC symptoms.

DESIGN, SETTING, AND PARTICIPANTS This prospective cohort study was conducted from April 2020 to February 2021, with follow-up of 60 to 90 days. The cohort consisted of adults enrolled in the University of California, Los Angeles, SARS-CoV-2 Ambulatory Program who had a laboratoryconfirmed symptomatic SARS-CoV-2 infection and were either hospitalized in a University of California, Los Angeles, hospital or one of 20 local health care facilities, or were outpatients referred by a primary care clinician. Data analysis was performed from March 2022 to February 2023.

EXPOSURE Laboratory-confirmed SARS-CoV-2 infection.

MAIN OUTCOMES AND MEASURES Patients responded to surveys that included questions about perceived cognitive deficits modified from the Perceived Deficits Questionnaire, Fifth Edition, (ie, trouble being organized, trouble concentrating, and forgetfulness) and symptoms of PCC at 30, 60, and 90 days after hospital discharge or initial laboratory-confirmed infection of SARS-CoV-2. Perceived cognitive deficits were scored on a scale from 0 to 4. Development of PCC was determined by patient self-report of persistent symptoms 60 or 90 days after initial SARS-CoV-2 infection or hospital discharge.

RESULTS Of 1296 patients enrolled in the program, 766 (59.1%) (mean [SD] age, 60.0 [16.7] years; 399 men [52.1%]; 317 Hispanic/Latinx patients [41.4%]) completed the perceived cognitive deficit items at 30 days after hospital discharge or outpatient diagnosis. Of the 766 patients, 276 (36.1%) perceived a cognitive deficit, with 164 (21.4%) having a mean score of greater than 0 to 1.5 and 112 patients (14.6%) having a mean score greater than 1.5. Prior cognitive difficulties (odds ratio [OR], 1.46; 95% CI, 1.16-1.83) and diagnosis of depressive disorder (OR, 1.51; 95% CI, 1.23-1.86) were associated with report of a perceived cognitive deficit. Patients reporting perceived cognitive deficits in the first 4 weeks of SARS-CoV-2 infection were more likely to report symptoms of PCC than those without perceived cognitive deficits (118 of 276 patients [42.8%] vs 105 of 490 patients [21.4%]; χ_{1}^2 , 38.9; *P* < .001). Adjusting for demographic and clinical factors, perceived cognitive deficit socre of >0 to 1.5: OR, 2.42; 95% CI, 1.62-3.60; patients with cognitive deficit score >1.5: OR, 2.97; 95% CI, 1.86-4.75) compared to patients who reported no perceived cognitive deficits.

Key Points

Question Among hospitalized and ambulatory patients with SARS-CoV-2 infection, how many perceived cognitive deficits early in the course of their infection and were these perceived deficits associated with post-COVID-19 condition (PCC)?

Findings In this cohort study of 766 patients with SARS-CoV-2 infection, 276 patients (36.1%) perceived cognitive deficits within 4 weeks of hospital discharge or outpatient infection, and patients with perceived cognitive deficits at 30 days were twice as likely as patients without perceived deficits to report symptoms of PCC at 60 to 90 days. Severity and persistence of perceived cognitive deficits were associated with symptoms of PCC.

Meaning These findings of perceived cognitive deficits early during the course of SARS-CoV-2 infection suggest that there is an affective component to PCC in some patients.

Supplemental content

Author affiliations and article information are listed at the end of this article.

(continued)

Open Access. This is an open access article distributed under the terms of the CC-BY License.

Abstract (continued)

CONCLUSIONS AND RELEVANCE These findings suggest that patient-reported perceived cognitive deficits in the first 4 weeks of SARS-CoV-2 infection are associated with PCC symptoms and that there may be an affective component to PCC in some patients. The underlying reasons for PCC merit additional exploration.

JAMA Network Open. 2023;6(5):e2311974. doi:10.1001/jamanetworkopen.2023.11974

Introduction

The world continues to grapple with the diverse clinical manifestations of SARS-CoV-2 infection and patients who have persistent symptoms. A key question is why some patients with SARS-CoV-2 have persistent symptoms, which the Centers for Disease Control and Prevention defines as symptoms that extend beyond 4 weeks after the initial infection.¹ Post-COVID-19 condition (PCC; colloquially known as long COVID) is characterized by many symptoms, of which cognitive impairment is a frequent complaint.²⁻⁸ So-called brain fog in particular has been a common and debilitating symptom⁹ affecting all age groups. Other neurocognitive symptoms associated with PCC include memory problems, difficulty concentrating, trouble focusing, and posttraumatic stress disorder.¹⁰⁻¹³ Furthermore, anxiety and depression are commonly reported alongside PCC.¹⁴⁻¹⁶

Most studies evaluating cognitive dysfunction in patients with SARS-CoV-2 have focused on the clinical characterization of cognitive symptoms associated with acute SARS-CoV-2 and PCC.^{4,5,9,17-19} With respect to both the acute and postacute sequelae of SARS-CoV-2, few studies have examined potential factors associated with the development of cognitive dysfunction. Given the potential long-term impact of neurocognitive deficits on quality of life and productivity,⁵ it is important to understand potential factors associated with cognitive dysfunction during the acute phase of SARS-CoV-2. Furthermore, some retrospective studies²⁰⁻²² and a 2022 prospective analysis²³ demonstrate that premorbid distress is associated with PCC symptoms at 4 weeks after initial infection or later, yet the association between early development of cognitive dysfuncts and the development of PCC is not understood. In this cohort study, we evaluated data from a clinical cohort of patients with SARS-CoV-2 who were surveyed concerning perceived cognitive deficits.²⁴ We explored the perceived level of cognitive deficit, factors associated with those deficits in the acute phase of SARS-CoV-2 infection, and the association of those deficits with the development of PCC.

Methods

This study was approved by the University of California, Los Angeles (UCLA) institutional review board with no requirement of informed consent due to the nature of the study as a retrospective analysis of deidentified clinical data in accordance with 45 CFR § 46. This study also follows the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) reporting guideline.²⁵The UCLA SARS-CoV-2 Ambulatory Program enrolled a longitudinal, prospective cohort of adults with laboratory-confirmed SARS-CoV-2 infection from April 2020 to February 2021.²⁴ Patients were followed up for clinical purposes with standardized questionnaires administered by nurses via telephone at 30, 60, and 90 days after hospital discharge or from the date of a positive SARS-CoV-2 test for nonhospitalized patients. Patients hospitalized for SARS-CoV-2 were discharged from UCLA hospitals or 20 other local health care facilities. Ambulatory patients were referred by a primary care clinician. A multidisciplinary team followed this cohort to address persistent symptoms associated with SARS-CoV-2.

The 30-day, 60-day, and 90-day questionnaires (eTable 1 in Supplement 1) assessed baseline functional activity level and perceived symptoms in the 4 weeks prior to each survey. Each survey asked whether the patient felt that their health was back to normal. Baseline functional activity and

maximal exertion level before COVID-19 were assessed by asking whether the patient could complete vigorous activities (eg, running), moderate activities (eg, moving a table), climb 1 flight of stairs, walk 1 block, lift or carry groceries, and bathe or dress independently.²⁶ Patients were asked about 9 symptom clusters during the previous 4 weeks: fever, chills or night sweats; loss of smell or taste; fatigue; shortness of breath; chest pain; numbness or tingling; nausea, vomiting or diarrhea; muscle aches; and rash. Perceived cognitive deficits were evaluated with 3 questions modified from the Perceived Deficits Questionnaire, Fifth Edition²⁷ that asked patients whether they had trouble getting things organized, whether they had trouble concentrating on activities like watching TV or reading a book, and whether they forgot what they talked about during a phone conversation during the previous 4 weeks. Likert scale response options included never, rarely, sometimes, often, and almost always. The Perceived Deficits Questionnaire, Fifth Edition has been studied in patients with multiple sclerosis, whiplash, and soft-tissue work injuries and has been found to be not associated with objective cognitive impairment, but instead with anxiety, depression, and self-efficacy.^{28,29}

Demographic characteristics (age, sex, race, and ethnicity) were obtained from the electronic health record, as were a history of diabetes, organ transplant, body mass index (BMI; calculated as weight in kilograms divided by height in meters squared), Elixhauser Comorbidity Index,³⁰ and facility of care for the initial SARS-CoV-2 infection (ie, inpatient facility or outpatient facility). Race and ethnicity (African American or Black, Asian, or White races, Hispanic or Latinx ethnicity, other race and/or ethnicity, or unknown) information were analyzed in this study because race and ethnicity have been associated with outcomes of SARS-CoV-2. Using the International Statistical Classification of Diseases and Related Health Problems, Tenth Revision (ICD-10), history of depression (ICD-10 code F32), anxiety (ICD-10 code F41), and cognitive difficulties (ICD-10 codes F1, F2, and F3 for dementia; code R41 for cognitive decline; and code G31 for cognitive impairment) were obtained from encounter-associated codes in the electronic health record. Insurance was collapsed into commercial, Medicare, Medicaid, and none or other. A Social Vulnerability Index (SVI)³¹ was calculated and split into quartiles. Patients were characterized as having PCC if they noted persistent SARS-CoV-2 symptoms among the 9 symptom clusters noted above (none of which were cognitive or affective symptoms) on the 90-day survey or the 60-day survey if the 90-day survey was incomplete.

Statistical Analysis

The 3 perceived cognitive deficit items from the questionnaire were scored from 0 to 4, and a mean of the 3 items was computed (α = .90). For the cohort of patients who completed the perceived cognitive deficits questions at the 30-day survey, we reported demographic and clinical characteristics, baseline functional activity status, history of depression, anxiety, or cognitive difficulties, and perceived cognitive deficits. We compared characteristics of patients reporting any perceived cognitive deficits vs no perceived cognitive deficits on the 30-day survey using *t* tests and χ^2 tests.

To identify factors independently associated with perceived cognitive deficits on the 30-day survey, we trichotomized the perceived cognitive deficit score (O, >O to 1.5, and >1.5 to 4) and performed ordinal logistic regression using this dependent variable. The proportional odds assumption was not violated by the χ^2 score test. Independent variables included age in years (18-39, 40-59, and 60 or above), sex, race or ethnicity, health insurance, baseline functional activity status, clinical characteristics (diabetes, organ transplant, and BMI), SVI (in quartiles), inpatient vs outpatient care facility, and history of depression, anxiety, or cognitive difficulties. Multiple imputation was used for missing BMI (4 values), organ transplant (3 values), and Elixhauser Comorbidity Index score (81 values) after confirming that the data were missing at random. We also performed logistic regression analysis on the complete cases and observed no difference in the estimates (ie, the direction) and statistical significance (eTable 2 in Supplement 1).

We evaluated the association between patient-reported perceived cognitive deficits on the 30-day survey with their reports of PCC on later surveys. We compared characteristics of patients

who reported PCC symptoms and those who did not report PCC symptoms using χ^2 and *t* tests. A multivariable logistic regression model evaluated factors associated with report of PCC on 20 imputed data sets. The final odds ratio (OR) estimates were obtained by pooling the parameter estimates and associated covariance matrices for each imputation set. The prespecified factors included in the model²⁴ included the aforementioned variables plus the trichotomized perceived cognitive deficits score. A 2-sided *P* < .05 was considered statistically significant. Analyses were performed using SAS statistical software version 9.4 (SAS Institute) from March 2022 to February 2023. To evaluate the longitudinal association of perceived cognitive deficits with PCC, we plotted the proportion of patients reporting any level of deficit for each cognitive deficit item, stratified by whether the patient reported PCC symptoms.

Results

Participant Characteristics

Of 1296 patients enrolled in the program from April 2020 to February 2021, 1038 patients completed any follow-up survey, and 766 patients (59% of the full cohort) completed the perceived cognitive deficits items on the survey approximately 30 days after hospital discharge or outpatient diagnosis. A total of 740 patients completed the 60-day survey, and 496 patients completed the 90-day survey. Of the 766 patients who completed the 30-day survey (mean [SD] age, 60.0 [16.7] years; 399 men [52.1%]; 317 Hispanic/Latinx patients [41.4%]; mean [SD] BMI, 30.0 [7.4]; median [IQR] SVI, 0.46 [0.20-0.76]), 293 patients (38.3%) had diabetes, and 90 patients (11.7%) had received an organ transplant. A total of 325 patients (42.4%) had commercial insurance. At baseline, 180 patients (23.5%) reported being able to complete vigorous activities and 368 patients (48.0%) reported being able to complete activities. On the basis of encounter data, 109 patients (14.2%) had a history of cognitive difficulties, 153 patients (20.0%) had a diagnosis of depression, and 213 patients (27.8%) had a diagnosis of anxiety (**Table 1**).

Perceived Cognitive Deficits Among Patients With SARS-CoV-2

During the 4 weeks following SARS-CoV-2 diagnosis, largely during the acute phase of illness, 490 of 766 patients (63.9%) reported no perceived cognitive deficits on the 3 cognitive survey items. A total of 231 patients (30.2%) reported trouble getting things organized, 220 patients (28.7%) reported trouble concentrating on activities like watching TV or reading a book, and 198 patients (25.8%) reported having forgotten what they had talked about during a telephone conversation. Of the 276 patients (36.1%) who perceived a cognitive deficit, 63 patients (22.8%) responded yes to only 1 item, 53 patients (19.2%) responded yes to 2 items, and 160 patients (58.0%) responded yes all 3 items. Overall, the mean (SD) perceived cognitive deficit score on the 30-day survey was 0.51 (0.82; median [IQR], 0.00 [0.00-1.00]); 164 patients (21.4%) had a mean score between 0.1 and 1.5, and 112 patients (14.6%) had a mean score above 1.5. During this period, the most common symptoms of acute COVID-19 were fatigue (432 patients [62%]), shortness of breath (475 patients [56%]), and muscle aches (345 patients [45%]) (eTable 2 in Supplement 1).

In bivariable analyses, no demographic or clinical factors other than diagnosis of depressive disorder, anxiety disorder, or cognitive difficulties and 1 aspect of physical function (moderate activities)were associated with patient report of a perceived cognitive deficit (Table 1). The logistic model demonstrates that a history of a cognitive difficulties (adjusted OR [aOR], 1.46; 95% CI, 1.16-1.83) and history of depressive disorder (aOR, 1.51; 95% CI, 1.23-1.86) were associated with a patient reporting cognitive deficits during the first 4 weeks after SARS-CoV-2. Women were more likely than men (aOR, 1.19; 95% CI, 1.01-1.40) and patients aged 40 to 59 years were more likely than younger patients (aOR, 1.35; 95% CI, 1.06-1.72) to report perceived cognitive deficits (**Table 2**).

Association of Perceived Cognitive Deficits With PCC

Patients reporting cognitive deficits in the first 4 weeks after SARS-CoV-2 were more likely to report PCC symptoms at 60 to 90 days than those without perceived cognitive deficits (118 patients [42.8%] vs 105 patients [21.4%]; χ_{1}^2 38.9; *P* < .001). Of the 223 patients with SARS-CoV-2 who

Fable 1. Demographic and Clinical Characteristi	cs of Patients			
	Patients, No.	(%)		
Characteristic	Total (N = 766)	Reporting a perceived cognitive deficit (n = 276) ^a	Reporting no perceived cognitive deficit (n = 490)	OR (95% CI)
Age, mean, (SD), y	60.0 (16.7)	59.6 (17.1)	60.2 (16.5)	NA
Age range, y				
18-39	105 (13.7)	36 (13.0)	69 (14.1)	1 [Reference]
40-59	257 (33.6)	104 (37.7)	153 (31.2)	1.30 (0.81-2.09)
≥60	404 (52.7)	136 (49.3)	268 (54.7)	0.97 (0.62-1.53)
Sex				
Female	367 (47.9)	144 (52.2)	223 (45.5)	1.31 (0.97-1.76)
Male	399 (52.1)	132 (47.8)	267 (54.5)	1 [Reference]
Race and ethnicity				
African American or Black	58 (7.8)	29 (10.5)	29 (5.9)	1.56 (0.88-2.78)
Asian	65 (8.5)	24 (8.7)	41 (8.4)	0.91 (0.52-1.61)
Hispanic or Latinx	317 (41.4)	99 (35.9)	218 (44.7)	0.71 (0.50-1.01)
White	233 (30.4)	91 (33)	142 (29.4)	1 [Reference]
Other race and ethnicity or unknown ^b	93 (12.1)	33 (12)	60 (11.6)	0.86 (0.52-1.42)
Comorbidities ^c				
Diabetes	293 (38.3)	101 (36.6)	192 (39.2)	0.90 (0.66-1.22)
Organ transplant	90 (11.7)	25 (9.1)	65 (13.3)	0.65 (0.40-1.06)
Body mass index, mean (SD) ^d	30.0 (7.4)	30.5 (7.6)	29.7 (7.2)	1.01 (0.99-1.04)
History of neuropsychiatric condition ^c				
Depressive disorder	153 (20.0)	81 (29.3)	72 (14.7)	2.41 (1.68-3.46)
Anxiety disorder	213 (27.8)	95 (34.4)	118 (24.1)	1.66 (1.20-2.29)
Cognitive difficulties (dementia, cognitive decline, or cognitive impairment)	109 (14.2)	57 (20.7)	52 (10.6)	2.19 (1.46-3.30)
Health care facility				
Outpatient	187 (24.4)	63 (22.8)	124 (25.3)	1 [Reference]
Inpatient	579 (75.6)	213 (77.2)	366 (74.7)	1.15 (0.81-1.62)
Social Vulnerability Index, percentile				
0-25	226 (29.5)	86 (31.2)	140 (28.6)	1 [Reference]
25.1-50	163 (21.3)	61 (22.1)	102 (20.8)	0.87 (0.64-1.48)
50.1-75	148 (19.3)	48 (17.4)	100 (20.4)	0.78 (0.51-1.21)
75.1-100	193 (25.2)	66 (23.9)	127 (25.9)	0.85 (0.57-1.26)
Missing	36 (4.7)	15 (5.4)	21 (4.3)	1.16 (0.57-2.38)
Elixhauser Comorbidity Index score, mean (SD) ^d	10.9 (12.5)	11 (13.5)	10.8 (12)	NA
Health insurance,				
Commercial	325 (42.4)	123 (44.6)	202 (41.2)	1 [Reference]
Medicare	290 (37.9)	98 (35.5)	192 (39.2)	0.84 (0.60-1.17)
Medicaid	126 (16.5)	45 (16.3)	81 (16.5)	0.91 (0.60-1.40)
Other or none	25 (3.3)	10 (3.6)	15 (3.1)	1.10 (0.48-2.51)
Baseline functional activity status				
Vigorous	180 (23.5)	77 (27.9)	103 (21.0)	1 [Reference]
Moderate	368 (48.0)	107 (38.8)	261 (53.3)	1.82 (1.26-2.64)
Able to climb 1 flight stairs or walk 1 block	53 (8.2)	22 (8.0)	31 (6.3)	1.05 (0.57-1.96)
Able to carry groceries, bathe, or dress	151 (19.7)	62 (22.5)	89 (18.2)	1.07 (0.69-1.66)

Abbreviations: NA, not applicable; OR, odds ratio.

- ^a Perceived cognitive deficits were reported at the 30-day interview conducted approximately 4 weeks after hospital discharge or ambulatory infection.
- ^b Other was defined as patients who chose multiple races or ethnicities or who chose other race or ethnicity.
- ^c Reference for diabetes is no diabetes, for organ transplant is no organ transplant, and for neuropsychiatric conditions is no condition.
- ^d Unit for body mass index and Elixhauser Comorbidity Index is 1 unit.

Table 2. Factors Associated With Perceived Cognitive Deficits at 30-Day Interview After Initial SARS-CoV-2 Infection^a

Factor	Adjusted OR (95% CI)
Age range, y	
18-39	1 [Reference]
40-59	1.35 (1.06-1.72)
≥60	0.88 (0.67-1.16)
Sex	
Female	1.19 (1.01-1.40)
Male	1 [Reference]
Race and ethnicity	
African American or Black	1.55 (1.00-2.40)
Asian	0.97 (0.62-1.51)
Hispanic or Latino	0.78 (0.52-1.18)
White	1 [Reference]
Other race and ethnicity or unknown ^b	0.97 (0.66-1.42)
Comorbidities ^c	
Diabetes	0.88 (0.63-1.24)
Organ transplant	0.74 (0.43-1.28)
Body mass index ^d	1.01 (0.99-1.03)
Health insurance	
Commercial	1 [Reference]
Medicare	0.79 (0.55-1.12)
Medicaid	1.05 (0.72-1.54)
Other or none	1.20 (0.64-2.28)
Social Vulnerability Index, percentile	
0-25	1 [Reference]
25.1-50	0.85 (0.63-1.16)
50.1-75	0.82 (0.59-1.13)
75.1-100	1.06 (0.75-1.48)
Missing	1.21 (0.70-2.11)
Elixhauser Comorbidity Index score ^d	1.00 (0.98-1.01)
Health care facility	
Inpatient	1.40 (0.96-2.05)
Outpatient	1 [Reference]
Baseline functional activity status	
Vigorous	1 [Reference]
Moderate	0.49 (0.21-1.15)
Able to climb 1 flight stairs or walk 1 block	1.22 (0.85-1.75)
Able to carry groceries, bathe, or dress	0.58 (0.39-0.88)
Missing	1.74 (0.74-4.06)
History of neuropsychiatric condition ^c	
Cognitive difficulties (dementia, cognitive decline, or cognitive impairment)	1.46 (1.16-1.83)
Depressive disorder	1.51 (1.23-1.86)
Anxiety disorder	1.08 (0.90-1.31)

Abbreviation: OR, odds ratio.

^a Perceived cognitive deficits were reported at the 30-day interview conducted approximately 4 weeks after hospital discharge or ambulatory infection. Ordinal logistic regression was performed for 766 patients.

^b Other was defined as patients who chose multiple races or ethnicities or who chose other race or ethnicity.

^c Reference for diabetes is no diabetes, for organ transplant is no organ transplant, and for neuropsychiatric conditions is no condition.

 $^{\rm d}$ Unit for body mass index and Elixhauser Comorbidity Index is 1 unit.

reported PCC symptoms at 60 to 90 days, 118 patients (52.9%) reported a perceived cognitive deficit on the 30-day survey, whereas of the 543 patients with SARS-CoV-2 who did not report PCC symptoms at 60 to 90 days, 158 patients (29.1%) reported perceived cognitive deficits on the 30-day survey. Among patients reporting any perceived cognitive deficit, 64 of 164 patients (39.0%) with a mean perceived cognitive deficit score below 1.5 reported PCC symptoms as did 54 of 112 patients (48.2%) with a mean perceived cognitive deficit score above 1.5. The association of patient demographic and clinical characteristics and whether patients reported PCC symptoms is displayed in **Table 3**. History of depressive disorder, cognitive difficulties, and diabetes were associated with reporting of PCC symptoms, and organ transplant was negatively associated with PCC symptoms. Perceived cognitive score was most associated with reporting PCC symptoms at the 60-day to 90-day survey (Table 3).

In the logistic model of PCC, perceived cognitive deficits were associated with reporting of PCC symptoms, and there was a dose-response association between severity of perceived cognitive deficits and likelihood of reporting PCC symptoms. Compared with patients reporting no perceived cognitive deficits, patients reporting a perceived cognitive deficit with a mean score of 1.5 or less were 2.42 times (95% CI, 1.52-3.60) more likely to report PCC symptoms and those with a score above 1.5 were 2.97 times (95% CI, 1.86 – 4.75) more likely to report PCC symptoms. Patients with diabetes (OR, 1.60; 95% CI, 1.11-2.30) were more likely to report PCC symptoms, whereas those with organ transplants (OR, 0.49; 95% CI, 0.26-0.92) or Medicaid insurance (OR, 0.54; 95% CI, 0.31-0.95) were less likely to report PCC symptoms (**Table 4**). We also performed logistic regression analysis on the complete cases and observed no difference in the estimates (ie, the direction) and statistical significance (eTable 3 in Supplement 1).

The proportion of patients reporting cognitive deficits at the question level, (**Figure**) shows that for patients who subsequently did not report PCC symptoms, proportions of patients perceiving persistent cognitive deficits at the 60-day and 90-day surveys decreased. However, among patients reporting PCC symptoms, perception of cognitive deficits remained about the same over the 3-month study period.

Discussion

In this cohort study, more than one-third of patients with SARS-CoV-2 perceived cognitive deficits on the 30-day survey after hospitalization or outpatient infection. Report of perceived cognitive deficit was associated with later reporting of PCC symptoms. To some degree, these findings may help us disentangle the complex construct that is PCC. Prior use of these survey items demonstrates that perceived cognitive deficit is not associated with objective deficient cognition; instead, they are associated with depression, anxiety, and lower perceived functional ability and control.^{28,29} Those findings are consistent with the findings in this cohort study showing that perceived cognitive deficits were associated with a history of anxiety disorder and depressive disorder, although we also found an association with prior cognitive difficulties. These findings suggest a substantial psychological component for long lasting SARS-CoV-2 symptoms for at least some patients.

These findings are also consistent with literature suggesting that PCC is a heterogeneous condition.² Nearly one-half of the patients with PCC reported no perceived cognitive deficits. The temporal trends show that the prevalence of perceived cognitive deficits declines among patients who did not report PCC symptoms while perceived cognitive deficits remained stable among those who reported PCC symptoms at 60 to 90 days. Furthermore, as seen in the model estimating PCC, perceived severity of the reported cognitive deficits is associated with the probability of later reporting PCC symptoms.

Many reports show that cognitive impairment and memory difficulty are common among patients with acute SARS-CoV-2,^{17,32} and among those with PCC.^{2,8,11-14} However, many studies use convenience samples or have no longitudinal data. A 2022 report²³ of a large sample of nurses (albeit with little gender or race heterogeneity) demonstrated that pre-SARS-CoV-2 distress, both general

Table 3. Association of Patient Demographics, Clinical Characteristics, and Perceived Cognitive Deficits With Report of PCC Symptoms at 60-90 Day Survey

	Patients, No. (%) (N	- 766)	_
Characteristic	No PCC symptoms (n = 543)	PCC symptoms (n = 223)	OR (95% CI)
Age range, y	(11 - 545)	(11 - 223)	OK (95% CI)
18-39	77 (14.2)	28 (12.6)	1 [Reference]
40-59	180 (33.1)	77 (34.5)	1.18 (0.71-1.96)
≥60	286 (52.7)	118 (52.9)	1.14 (0.70-1.84)
Sex	200 (32.7)	110 (32.3)	1.11 (0.70 1.01)
Female	255 (47.0)	112 (50.2)	1.14 (0.83-1.56)
Male	288 (53.0)	111 (49.8)	1 [Reference]
Race and ethnicity	200 (33.0)	111 (+9.0)	I [Kelelence]
African American or Black	40 (7.4)	18 (8.1)	1.04 (0.56-1.94)
Asian	46 (8.5)	19 (8.5)	0.95 (0.52-1.74)
Hispanic or Latino	226 (41.6)	92 (41.3)	0.94 (0.65-1.36)
White	164 (30.2)	71 (31.8)	1 [Reference]
Other race and ethnicity or unknown ^a	67 (12.3)	23 (10.3)	0.79 (0.46-1.37)
Comorbidities ^b	/	/	
Diabetes	194 (35.7)	99 (44.4)	1.44 (1.05-1.97)
Organ transplant	73 (13.4)	17 (7.6)	0.53 (0.31-0.92)
Body mass index			
<30	327 (60.2)	122 (54.7)	1 [Reference]
≥30	216 (39.8)	101 (45.3)	1.25 (0.92-1.72)
History of neuropsychiatric condition ^b			
Depressive disorder	94 (17.3)	59 (26.5)	1.72 (1.19-2.49)
Anxiety disorder	141 (26.0)	72 (32.3)	1.36 (0.97-1.91)
Cognitive difficulties (dementia, cognitive decline, or cognitive impairment)	65 (12.0)	44 (19.7)	1.81 (1.19-2.75)
Health care facility,			
Outpatient	140 (25.8)	47 (21.1)	1 [Reference]
Inpatient	403 (74.2)	176 (78.9)	1.30 (0.89-1.89)
Social Vulnerability Index, percentile			
0-25	159 (29.3)	67 (30.0)	1 [Reference]
25.1-50	118 (21.7)	45 (20.2)	0.91 (0.58-1.41)
50.1-75	104 (19.2)	44 (19.7)	1.00 (0.64-1.58)
75.1-100	138 (25.4)	55 (24.7)	0.95 (0.62-1.44)
Missing	24 (4.4)	12 (5.4)	1.19 (0.56-2.51)
Elixhauser Comorbidity Index score,			
<10	262 (48.3)	105 (47.1)	1 [Reference]
≥10	224 (41.3)	94 (42.2)	1.05 (0.75-1.46)
Health insurance			
Commercial	224 (41.3)	101 (45.3)	1 [Reference]
Medicare	204 (37.6)	86 (38.6)	0.94 (0.66-1.32)
Medicaid	98 (18.0)	28 (12.6)	0.63 (0.39-1.03)
Other or none	17 (3.1)	8 (3.6)	1.04 (0.44-2.50)
Baseline functional activity status	17 (011)	0 (010)	1101 (0111 2100)
Vigorous	125 (23.0)	55 (24.7)	1 [Reference]
Moderate	266 (49.0)	102 (45.7)	0.87 (0.59-1.29)
Able to climb 1 flight stairs or walk 1 block	117 (21.5)	50 (22.4)	0.97 (0.61-1.54)
Able to carry groceries, bathe, or dress			1.09 (0.51-2.33)
,,,,	25 (4.6)	12 (5.4)	
Missing	10 (1.8)	4 (1.8)	0.91 (0.27-3.03)
Perceived cognitive deficit score ^c		105 (47.1)	1 [D=(]
0	385 (70.9)	105 (47.1)	1 [Reference]
>0 to 1.5	100 (18.4)	64 (28.7)	2.35 (1.60-3.43)
>1.5 to 4	58 (10.7)	54 (24.2)	3.41 (2.22-5.24)

Abbreviations: OR, odds ratio; PCC, post-COVID-19 condition.

^a Other was defined as patients who chose multiple races or ethnicities or who chose other race or ethnicity.

^b Reference for diabetes is no diabetes, for organ transplant is no organ transplant, and for neuropsychiatric conditions is no condition.

^c Perceived cognitive deficits reported at the 30-day survey conducted approximately 4 weeks after hospital discharge or ambulatory infection.

Table 4. Factors Associated With Reporting of PCC Symptoms	
After Ordinal Logistic Regression	

Factor	OR (95% CI)
Age range, y	
18-39	1 [Reference]
40-59	0.98 (0.56-1.72)
≥60	1.01 (0.55-1.85)
Sex	
Female	1.05 (0.73-1.50)
Male	1 [Reference]
Race and ethnicity	
African American or Black	0.89 (0.45-1.78)
Asian	0.96 (0.50-1.86)
Hispanic or Latinx	1.08 (0.68-1.74)
White	1 [Reference]
Other race and ethnicity or unknown ^a	0.81 (0.45-1.47)
Comorbidities ^b	
Diabetes	1.60 (1.11-2.30)
Organ transplant	0.49 (0.26-0.92)
Body mass index ^c	1.01 (0.98-1.03)
Health insurance	
Commercial	1 [Reference]
Medicare	0.92 (0.05-1.43)
Medicaid	0.54 (0.31-0.95)
Other or none	0.99 (0.39-2.51)
Social Vulnerability Index, percentile	
0-25	1 [Reference]
25.1-50	0.90 (0.55-1.47)
50.1-75	1.04 (0.62-1.75)
75.1-100	1.13 (0.65-1.98)
Missing	1.17 (0.52-2.65)
Elixhauser Comorbidity Index score ^c	0.99 (0.98-1.01)
Health care facility	
Inpatient	1.45 (0.95-2.22)
Outpatient	1 [Reference]
Baseline functional activity status	
Vigorous	1 [Reference]
Moderate	0.98 (0.63-1.52)
Able to climb 1 flight stairs or walk 1 block	0.81 (0.47-1.40)
Able to carry groceries, bathe. or dress	1.00 (0.41-2.43)
Missing	0.52 (0.13-2.04)
History of neuropsychiatric	
condition ^b	
	1.48 (0.88-2.46)
condition ^b Cognitive difficulty (dementia, cognitive decline,	1.48 (0.88-2.46) 1.12 (0.70-1.79)

(continued)

Table 4. Factors Associated With Reporting of PCC Symptoms After Ordinal Logistic Regression (continued)

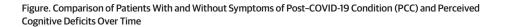
Factor	OR (95% CI)
Perceived Cognitive Deficits score	
0	1 [Reference]
>0 to 1.5	2.42 (1.62-3.60)
>1.5 to 4	2.97 (1.86-4.75)

Abbreviations: OR, odds ratio; PCC, post-COVID-19 condition.

^a Other was defined as patients who chose multiple races or ethnicities or who chose other race or ethnicity.

^b Reference for diabetes is no diabetes, for organ transplant is no organ transplant and for neuropsychiatric conditions is no condition.

^c Unit for body mass index and Elixhauser Comorbidity Index is 1 unit.



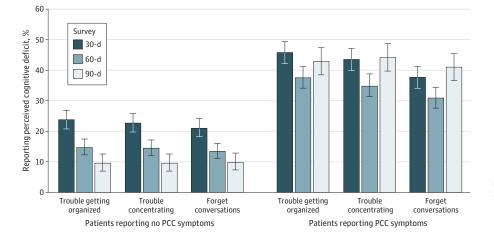


Figure shows patient responses to perceived cognitive deficit items from the 30-day, 60-day, and 90-day surveys, comparing patients who did or did not report PCC symptoms at the 60-day or 90-day survey. Error bars denote 95% Cls.

distress and COVID-19-related distress, was associated with greater likelihood of COVID-19 symptoms persisting for 4 or more weeks. Our findings come from a single health system continuity sample, and we are able to adjust the diverse cohort for demographic and clinical characteristics.²⁴

The finding that more than one-half of patients with PCC perceived cognitive deficits early during the condition is provocative, yet these data likely create more questions than they answer. Do the reported cognitive deficits influence the content or quality of responses to later surveys? Are the early reported cognitive deficits related to the SARS-CoV-2 infection consistent with imaging changes that have been found ?^{33,34} If so, why are these symptoms related to a history of depressive disorder, anxiety disorder, and cognitive difficulties? These data suggest that the constructs of affect and control play a substantial role in the development of PCC for at least some patients. From a clinical perspective, these data might suggest that early evaluation of perceived cognitive deficits might help in identification of patients with acute COVID-19 who should receive more intensive monitoring for persistence of symptoms and perhaps for a focus on intervention.

Limitations

This study has several limitations, including a lack of objective measures of cognition because these clinical surveys aimed to identify patients at risk of clinical deterioration. The principal survey items that elicit subjective responses about perceived cognitive deficits have not been shown to correlate with objective cognitive deficits. Measures of pre-SARS-CoV-2 cognition, depression, and anxiety were obtained from clinical encounter data, which is known to miss these diagnoses.^{35,36} In addition,

the definition of PCC may be biased because it is a subjective rating of a limited number of symptoms. Furthermore, referral bias may exist among the outpatient cohort, because physicians may have referred patients deemed clinically high risk to the program, which may affect generalizability of the outpatient cohort. The patient cohort studied was derived from an academic medical center, indicating that it may not be generalizable to other groups of patients with COVID-19.

Conclusions

In a longitudinal cohort study of patients with SARS-CoV-2 in 1 health care system, we found an association between perceived cognitive deficits early in the disease and PCC, which suggests direction for exploration of the underpinnings of PCC.

ARTICLE INFORMATION

Accepted for Publication: March 23, 2023.

Published: May 5, 2023. doi:10.1001/jamanetworkopen.2023.11974

Open Access: This is an open access article distributed under the terms of the CC-BY License. © 2023 Liu TC et al. *JAMA Network Open.*

Corresponding Author: Neil Wenger, MD, MPH, Division of General Internal Medicine & Health Services Research, UCLA Department of Medicine, 1100 Glendon Ave, Ste 850, Los Angeles, CA 90024 (nwenger@mednet.ucla.edu).

Author Affiliations: Division of General Internal Medicine & Health Services Research, Department of Medicine, University of California, Los Angeles (Liu, Yoo, Viswanathan, Wenger); Department of Medicine Statistics Core, University of California, Los Angeles. (Sim, Motwani).

Author Contributions: Dr Sim and Dr Wenger had full access to all of the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

Concept and design: Liu, Yoo, Wenger.

Acquisition, analysis, or interpretation of data: All authors.

Drafting of the manuscript: Liu, Wenger.

Critical revision of the manuscript for important intellectual content: All authors.

Statistical analysis: Liu, Sim, Motwani, Viswanathan.

Administrative, technical, or material support: Liu.

Supervision: Wenger.

Conflict of Interest Disclosures: None reported.

Funding/Support: Research reported in this publication was supported by the National Center for Advancing Translational Science of the National Institutes of Health under the UCLA Clinical and Translational Science Institute grant number UL1TRO01881.

Role of the Funder/Sponsor: The funder had no role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; preparation, review, or approval of the manuscript; and decision to submit the manuscript for publication.

Data Sharing Statement: See Supplement 2.

Additional Contributions: Victor Gonzalez, BA (University of California, Los Angeles), provided administrative support; he was not compensated for this work.

REFERENCES

1. Centers for Disease Control and Prevention. Long-COVID or post-COVID conditions. Accessed October 14, 2022. https://www.cdc.gov/coronavirus/2019-ncov/long-term-effects/index.html

2. Global Burden of Disease Long COVID Collaborators. Estimated global proportions of individuals with persistent fatigue, cognitive, and respiratory symptom clusters following symptomatic COVID-19 in 2020 and 2021. *JAMA*. 2022;328(16):1604-1615. doi:10.1001/jama.2022.18931

3. van Kessel SAM, Olde Hartman TC, Lucassen PLBJ, van Jaarsveld CHM. Post-acute and long-COVID-19 symptoms in patients with mild diseases: a systematic review. *Fam Pract*. 2022;39(1):159-167. doi:10.1093/fampra/cmab076

4. Miskowiak KW, Johnsen S, Sattler SM, et al. Cognitive impairments four months after COVID-19 hospital discharge: pattern, severity and association with illness variables. *Eur Neuropsychopharmacol*. 2021;46:39-48. doi:10.1016/j.euroneuro.2021.03.019

5. Graham EL, Clark JR, Orban ZS, et al. Persistent neurologic symptoms and cognitive dysfunction in non-hospitalized Covid-19 "long haulers". Ann Clin Transl Neurol. 2021;8(5):1073-1085. doi:10.1002/acn3.51350

6. Hellmuth J, Barnett TA, Asken BM, et al. Persistent COVID-19-associated neurocognitive symptoms in non-hospitalized patients. *J Neurovirol*. 2021;27(1):191-195. doi:10.1007/s13365-021-00954-4

7. Jaywant A, Vanderlind WM, Alexopoulos GS, Fridman CB, Perlis RH, Gunning FM. Frequency and profile of objective cognitive deficits in hospitalized patients recovering from COVID-19. *Neuropsychopharmacology*. 2021; 46(13):2235-2240. doi:10.1038/s41386-021-00978-8

8. Liu YH, Chen Y, Wang QH, et al. One-year trajectory of cognitive changes in older survivors of COVID-19 in Wuhan, China: a longitudinal cohort study. *JAMA Neurol*. 2022;79(5):509-517. doi:10.1001/jamaneurol.2022.0461

9. Crook H, Raza S, Nowell J, Young M, Edison P. Long covid-mechanisms, risk factors, and management. *BMJ*. 2021;374(1648):n1648. doi:10.1136/bmj.n1648

10. Davis HE, Assaf GS, McCorkell L, et al. Characterizing long COVID in an international cohort: 7 months of symptoms and their impact. *EclinicalMedicine*. 2021;38:101019. doi:10.1016/j.eclinm.2021.101019

11. Lambert NJ. Survivor Corps. COVID-19 "long hauler" symptoms survey report. Indiana University School of Medicine; 2020. Accessed October 14, 2022. https://dig.abclocal.go.com/wls/documents/2020/072720-wls-covid-symptom-study-doc.pdf

12. Garrigues E, Janvier P, Kherabi Y, et al. Post-discharge persistent symptoms and health-related quality of life after hospitalization for COVID-19. *J Infect*. 2020;81(6):e4-e6. doi:10.1016/j.jinf.2020.08.029

13. Bellan M, Soddu D, Balbo PE, et al. Respiratory and psychophysical sequelae among patients with COVID-19 four months after hospital discharge. *JAMA Netw Open*. 2021;4(1):e2036142. doi:10.1001/jamanetworkopen. 2020.36142

14. Huang C, Huang L, Wang Y, et al. 6-Month consequences of COVID-19 in patients discharged from hospital: a cohort study. *Lancet.* 2021;397(10270):220-232. doi:10.1016/S0140-6736(20)32656-8

15. Xiong Q, Xu M, Li J, et al. Clinical sequelae of COVID-19 survivors in Wuhan, China: a single-centre longitudinal study. *Clin Microbiol Infect*. 2021;27(1):89-95. doi:10.1016/j.cmi.2020.09.023

16. Poyraz BÇ, Poyraz CA, Olgun Y, et al. Psychiatric morbidity and protracted symptoms after COVID-19. *Psychiatry Res.* 2021;295:113604. doi:10.1016/j.psychres.2020.113604

17. Daroische R, Hemminghyth MS, Eilertsen TH, Breitve MH, Chwiszczuk LJ. Cognitive impairment after COVID-19: a review on objective test data. *Front Neurol*. 2021;12:699582. doi:10.3389/fneur.2021.699582

18. Delgado-Alonso C, Valles-Salgado M, Delgado-Álvarez A, et al. Cognitive dysfunction associated with COVID-19: a comprehensive neuropsychological study. *J Psychiatr Res*. 2022;150:40-46. doi:10.1016/j.jpsychires. 2022.03.033

19. Hampshire A, Trender W, Chamberlain SR, et al. Cognitive deficits in people who have recovered from COVID-19. *EClinicalMedicine*. 2021;39:101044. doi:10.1016/j.eclinm.2021.101044

20. Tenforde MW, Kim SS, Lindsell CJ, et al; IVY Network Investigators; CDC COVID-19 Response Team; Symptom duration and risk factors for delayed return to usual health among outpatients with COVID-19 in a multistate health care systems network—United States, March-June 2020. *MMWR Morb Mortal Wkly Rep.* 2020;69(30):993-998. doi:10.15585/mmwr.mm6930e1

21. Blackett JW, Wainberg M, Elkind MSV, Freedberg DE. Potential long coronavirus disease 2019 gastrointestinal symptoms 6 months after coronavirus infection are associated with mental health symptoms. *Gastroenterology*. 2022;162(2):648-650.e2. doi:10.1053/j.gastro.2021.10.040

22. Garjani A, Middleton RM, Nicholas R, Evangelou N. Recovery from COVID-19 in multiple sclerosis: a prospective and longitudinal cohort study of the United Kingdom Multiple Sclerosis Register. *Neurol Neuroimmunol Neuroinflamm*. 2021;9(1):e1118. doi:10.1212/NXI.00000000000001118

23. Wang S, Quan L, Chavarro JE, et al. Associations of depression, anxiety, worry, perceived stress, and loneliness prior to infection with risk of post-COVID-19 conditions. *JAMA Psychiatry*. 2022;79(11):1081-1091. doi:10.1001/jamapsychiatry.2022.2640

24. Yoo SM, Liu TC, Motwani Y, et al. Factors associated with post-acute sequelae of SARS-CoV-2 (PASC) after diagnosis of symptomatic COVID-19 in the inpatient and outpatient setting in a diverse cohort. *J Gen Intern Med.* 2022;37(8):1988-1995. doi:10.1007/s11606-022-07523-3

25. von Elm E, Altman DG, Egger M, Pocock SJ, Gøtzsche PC, Vandenbroucke JP; STROBE Initiative. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. *Ann Intern Med*. 2007;147(8):573-577. doi:10.7326/0003-4819-147-8-200710160-00010

26. Ware JE Jr, Sherbourne CD. The MOS 36-item short-form health survey (SF-36). I. Conceptual framework and item selection. *Med Care*. 1992;30(6):473-483. doi:10.1097/00005650-199206000-00002

27. Cha D. Perceived deficits questionnaire—depression, 5-item (PDQ-D-5). In: McIntyre R, ed. *Cognitive Impairment in Major Depressive Disorder: Clinical Relevance, Biological Substrates, and Treatment Opportunities.* Cambridge University Press; 2016:253-256.

28. Strober LB, Binder A, Nikelshpur OM, Chiaravalloti N, DeLuca J. The Perceived Deficits Questionnaire: perception, deficit, or distress? *Int J MS Care*. 2016;18(4):183-190. doi:10.7224/1537-2073.2015-028

29. Sullivan MJ, Hall E, Bartolacci R, Sullivan ME, Adams H. Perceived cognitive deficits, emotional distress and disability following whiplash injury. *Pain Res Manag*. 2002;7(3):120-126. doi:10.1155/2002/502984

30. Thompson NR, Fan Y, Dalton JE, et al. A new Elixhauser-based comorbidity summary measure to predict in-hospital mortality. *Med Care*. 2015;53(4):374-379. doi:10.1097/MLR.00000000000326

31. Centers for Disease Control and Prevention; Agency for Toxic Substances and Disease Registry. CDC/ATSDR Social Vulnerability Index. 2021. Accessed October 14, 2022. https://www.atsdr.cdc.gov/placeandhealth/svi/data_documentation_download.html

32. McIoughlin BC, Miles A, Webb TE, et al. Functional and cognitive outcomes after COVID-19 delirium. *Eur Geriatr Med*. 2020;11(5):857-862. doi:10.1007/s41999-020-00353-8

33. Manca R, De Marco M, Ince PG, Venneri A. heterogeneity in regional damage detected by neuroimaging and neuropathological studies in older adults with COVID-19: a cognitive-neuroscience systematic review to inform the long-term impact of the virus on neurocognitive trajectories. *Front Aging Neurosci.* 2021;13:646908. doi:10.3389/fnagi.2021.646908

34. Douaud G, Lee S, Alfaro-Almagro F, et al. SARS-CoV-2 is associated with changes in brain structure in UK Biobank. *Nature*. 2022;604(7907):697-707. doi:10.1038/s41586-022-04569-5

35. Mackin RS, Areán PA. Incidence and documentation of cognitive impairment among older adults with severe mental illness in a community mental health setting. *Am J Geriatr Psychiatry*. 2009;17(1):75-82. doi:10.1097/JGP. 0b013e31818cd3e5

36. Cully JA, Jimenez DE, Ledoux TA, Deswal A. Recognition and treatment of depression and anxiety symptoms in heart failure. *Prim Care Companion J Clin Psychiatry*. 2009;11(3):103-109. doi:10.4088/PCC.08m00700

SUPPLEMENT 1.

eTable 1. Clinical and Functional Survey eTable 2. Symptoms During Acute COVID-19 and at 30-Day Interview eTable 3. Complete Case Analysis: Factors Associated With Development of Long COVID

SUPPLEMENT 2. Data Sharing Statement