
Postacute Sequelae of SARS-CoV-2 in University Setting

Megan Landry, Sydney Bornstein, Nitasha Nagaraj, Gary A. Sardon Jr., Amanda Castel, Amita Vyas, Karen McDonnell, Mira Agneshwar, Alyson Wilkinson, Lynn Goldman

Postacute sequelae of SARS-CoV-2 infection, commonly known as long COVID, is estimated to affect 10% to 80% of COVID-19 survivors. We examined the prevalence and predictors of long COVID from a sample of 1,338 COVID-19 cases among university members in Washington, DC, USA, during July 2021–March 2022. Cases were followed up after 30 days of the initial positive result with confidential electronic surveys including questions about long COVID. The prevalence of long COVID was 36%. Long COVID was more prevalent among those who had underlying conditions, who were not fully vaccinated, who were female, who were former/current smokers, who experienced acute COVID-19 symptoms, who reported higher symptom counts, who sought medical care, or who received antibody treatment. Understanding long COVID among university members is imperative to support persons who have ongoing symptoms and to strengthen existing services or make referrals to other services, such as mental health, exercise programs, or long-term health studies.

It is estimated that 1 in 3 Americans who have SARS-CoV-2 infection will experience symptoms related to postacute sequelae of SARS-CoV-2 (1), also referred to as long COVID (other terms include long-haul coronavirus disease, post–COVID-19 conditions, or chronic COVID-19) (2). The length of time that a person must experience symptoms to be considered to have long COVID is not universally accepted; definitions range from 28 days to 6 months after acute SARS-CoV-2 infection (3–7). A recent World Health Organization working group used a Delphi process to conclude that “a post-COVID-19 condition occurs in individuals with a history of probable or confirmed SARS-CoV-2 infection, usually 3 months from the onset of COVID-19 with symptoms that last for at least 2 months and cannot be explained by an alternative diagnosis” (8).

Regardless of a universally agreed upon length of time a person must experience symptoms to be characterized as long COVID, this sequela has been suggested to be the “next national health disaster” (9), and because of discrepancies in symptoms and long-term effects on quality of life, there seem to be more questions than answers. Although long COVID manifests differently in each person, nearly 50 signs and symptoms have been linked to the condition (10). The most common signs and symptoms are fatigue, shortness of breath, muscle pain, joint pain, headache, cough, chest pain, altered smell, altered taste, and diarrhea (11). Other reported signs and symptoms include cognitive impairment (known as brain fog), memory loss, palpitations, anxiety, sore throat, sleep disorders, runny nose, sneezing, hoarseness, ear pain, thoughts of self-harm and suicide, seizures, and bladder incontinence (8,11), as well as cardiac effects, such as myocardial inflammation (12).

Although some investigators have reported that long COVID occurs at rates that are independent of symptom severity (11–13), others have found long COVID is more common among patients hospitalized for COVID-19 or those who experienced moderate-to-severe symptoms (6,11,14–20). However, long COVID has been observed in patients who were asymptomatic (2) or only experienced mild symptoms, and it has been reported that symptoms can fluctuate or relapse (7–8,21–23). Furthermore, little is known about long COVID signs and symptoms and predictors on a college campus, where most of the population is young and healthy, but among whom potential complications of long COVID could be detrimental to academic learning and overall quality of life.

Long COVID signs and symptoms might vary by sex, age, and initial illness severity. For example, nervous system symptoms such as headaches and dizziness are more common among women, but men are more likely to have musculoskeletal system symptoms such as pain in the muscles or joints and numbness of the limbs (24). Younger patients have reported

Author affiliation: The George Washington University Milliken Institute School of Public Health, Washington DC, USA

DOI: <https://doi.org/10.3201/eid2903.221522>

more headaches, abdominal symptoms, and anxiety/depression, and older patients were more likely to have breathing difficulties, cognitive symptoms, pain, and fatigue (19).

Aside from the medical illness long COVID poses, persistent signs and symptoms can negatively affect leisure and work, causing further strain on one's quality of life. Persons who have long COVID frequently experience a substantial reduction or impairment in the ability to engage in preillness levels of occupational, educational, social, or personal activities that persist for >6 months (14). They might also experience difficulty sticking to daily routines, dealing with stress, getting household tasks done, and caring for/supporting others (25). Abnormal scores on mental and cognitive health questionnaires have also been observed among patients who have long COVID (7). Our study builds on the existing knowledge base by examining the prevalence and predictors of long COVID among a sample of university members, including students, faculty, and staff, who tested positive for COVID-19 over an 18-month period.

Methods

COVID-19 Case Identification

The George Washington University COVID-19 surveillance and testing program identified 4,800 COVID-19 cases during August 2020–February 2022. COVID-19 positivity at George Washington University was determined on the basis of PCR tests that were performed in the George Washington University Clinical Laboratory Improvement Amendments, or Clinical Laboratory Improvement Amendment-certified, Public Health Laboratory (n = 3,228); other cases were identified through results uploaded to the

person's medical portal from external positive tests, either PCRs from an external Clinical Laboratory Improvement Amendment-certified laboratory or self-administered antigen tests (n = 1,572). Only antigen tests approved for emergency use under the Food and Drug Administration emergency use authorization were accepted (26).

COVID-19 Case Investigation Data Collection

As COVID-19 cases were identified, the George Washington University Campus COVID-19 Support Team (CCST), which is responsible for campus-related COVID-19 case management (27), completed case investigations within 24–48 hours of the person receiving a positive test result. Among the 4,800 positive results during August 2020–February 2022, a total of 133 initial case investigations were incomplete because of loss to follow-up, meaning they could not be reached by telephone or electronic survey; because the case was already cleared by a medical provider (because of not being able to reach the person during their isolation period); or because the person refused to complete the interview. Furthermore, 1,072 persons were missing case investigation data, such as missing data for symptoms or underlying conditions. Those exclusions resulted in 3,595 positive test results (with corresponding completed case investigation data) for which CCST had obtained complete case investigation data (Table 1).

Long COVID Follow-up Data Collection

During July 2021–March 2022, all 4,800 positive COVID-19 test results reported during August 2020–February 2022 were followed up with confidential electronic surveys sent to each patient at least 28 days after their initial positive result that included

Table 1. Initial case investigation and long COVID data collection for postacute sequelae of SARS-CoV-2 in university setting, Washington, DC, USA*

Characteristic	Value
Positive test results reported during 2020 Aug–2022 Feb	4,800
GWU PCR	3,228
External CLIA PCR or self-administered antigen	1,572
Initial case investigation incomplete	133
Initial case investigation data missing	1,072
Total positive test results with case investigation data	3,595
Long COVID surveys sent out, 2021 Jul–2022 Mar	4,800
Repeat infection duplicates	143
Persons surveyed	4,657
Responses received	1,493
Duplicate responses/multiple responses	11
Persons responding	1,482
Response rate, 1,482/4,657	31.8%
Responders with incomplete case investigation	141
Responders with incomplete long COVID questions	3
Total valid responses	1,338

*Values are no. except as indicated. CLIA, Clinical Laboratory Improvement Amendments; GWU, George Washington University.

questions about long COVID. Those data were merged with the COVID-19 case investigation data.

For the long COVID follow-up survey data collection, we determined that 143 persons had ≥ 2 COVID-19 diagnoses during August 2020–February 2022; those persons were only included once in the long COVID follow-up data collection, resulting in a total of 4,657 persons who were COVID-19 positive during the study period. The follow-up survey had a response rate of 32% (1,482/4,657). We observed major differences in age, university affiliation, underlying conditions, and vaccination status at the time of test between follow-up survey respondents and nonrespondents (Appendix 1, <https://wwwnc.cdc.gov/EID/article/29/3/22-1522-App1.pdf>). A total of 11 respondents completed the follow-up survey twice but were only counted once for the response rate. Not all responses were usable in the final analysis: 141 did not have a complete initial case investigation, and 3 did not provide responses to the survey questions about long COVID, removing them from the final sample. Thus, the final analytic sample consisted of 1,338 respondents (Table 1).

Instrument and Measures

Survey Instrument

The long COVID survey was designed as a follow-up telephone interview (Appendix 2, <https://wwwnc.cdc.gov/EID/article/29/3/22-1522-App2.pdf>); interviews were administered by CCST during July 2021–March 2022, and all survey responses were stored on REDCap, a secure web application for online surveys and databases (28). Initially, CCST interviewers exclusively administered the follow-up survey by telephone calls. However, after 3 months, a link to an electronic survey was sent to all remaining cases in addition to calling. Three call attempts were made over a period of 5 weeks, prompting case-patients to complete an anonymous survey. The long COVID survey consisted of close-ended questions pertaining to symptoms during the postisolation period and behavior changes from preisolation to postisolation periods (Appendix 1). The survey took ≈ 15 –20 minutes to complete, and at the conclusion, a list of resources to assist with long COVID symptoms was provided.

Measures

We defined long COVID as experiencing ≥ 1 of the following symptoms lasting for >28 days after a respondent's 10-day isolation period ended (2): difficulty driving, difficulty having conversations, difficulty making decisions, difficulty thinking, fatigue, feeling

anxious, feeling depressed or sad, loss of smell, loss of taste, memory loss, muscle pain, muscle weakness, shortness of breath or difficulty breathing, trouble sleeping, worsening of symptoms after physical activity, worsening of symptoms after mental activity, or other symptoms. In addition, respondents were considered to have long COVID if they reported still experiencing COVID-19–related symptoms at the time of the long COVID survey.

Sociodemographic Characteristics

We calculated age from the respondent's date of birth extracted from their health record. Sex and race were self-reported at the time of the case investigation. We determined school affiliation by asking respondents their primary university affiliation at the time of the case investigation.

Symptoms and Underlying Conditions

We measured symptoms at the time of the case investigation by asking if respondents experienced any of the following: chest pain, chills, congestion, cough, diarrhea, fatigue, fever, headache, loss of smell, loss of taste, muscle pain, nausea or vomiting, runny nose, shortness of breath, sore throat, or other symptoms. At the time of the case investigation, respondents self-reported any of the following medical conditions: diabetes, asthma, hypertension, obesity, sickle cell disease, cancer, chronic kidney disease, lung diseases, serious heart conditions, or other conditions. Smoking status was self-reported as current/former smoker or vaper.

Vaccination Status and Severity of COVID-19 Infection

Over the course of the study period, COVID-19 vaccine availability and recommendations shifted dramatically. In December 2020, vaccines were first available but only for select groups such as health-care workers, the elderly, and certain other susceptible populations. During March–April 2021, vaccines were made available to all adults (>16 years of age) across all US states. In June 2021, George Washington University mandated all members of the campus community to be up to date (an up-to-date course of COVID-19 vaccines consisted of either 2 doses of Moderna [<https://www.modernatx.com>] or Pfizer-BioNTech [<https://www.pfizer.com>] vaccines or 1 Johnson & Johnson/Janssen [<https://www.jnj.com>] immunization with the primary series of COVID-19 vaccinations), or to have obtained an exemption. In September 2021, in the United States, COVID-19 booster shots were authorized for administration 6 months after the second dose of Pfizer or Moderna

or 2 months after 1 dose of Johnson & Johnson/Janssen, initially just for persons ≥ 65 years of age, persons living or working in high-risk settings, or persons who had underlying conditions. In November 2021, booster shots were recommended for all adults >18 years of age. In January 2022, George Washington University mandated all members of the campus community to have a booster shot or to have obtained an exemption. Community members uploaded vaccine information including the type of vaccine(s) and dates of vaccinations and boosters as a condition of employment and access to campus. This information was used to determine vaccine status on the date of first positive COVID-19 test. The case investigation interviews also collected data about whether medical care was sought, hospitalizations, and administration of monoclonal antibodies.

Statistical Analysis

We described continuous variables by using medians and interquartile ranges (IQRs) and categorical variables by using frequencies and percentages. We compared characteristics of survey respondents by using χ^2 tests for categorical variables and Wilcoxon rank-sum tests for continuous variables. We used logistic regression to determine unadjusted associations between characteristics of survey respondents and long COVID status. We included characteristics that were found to be significantly associated with long COVID status in bivariate analyses in multivariable logistic regression models. All hypothesis tests were 2-sided, and statistical significance was set at an α of 0.05. We performed analyses by using SAS version 9.4 (SAS Institute, Inc., <https://www.sas.com>).

All university community members provided informed consent to participate in the George Washington University COVID-19 surveillance program. The George Washington University Institutional Review Board concluded that these were non-research-related activities.

Results

Overall, the median age of respondents was 23 (IQR 21–32) years, and the median symptom count was 4 (IQR 1–6) (Table 2). More than half of respondents were female (63.4%) and non-Hispanic White (55.7%). Most (73.4%) respondents were students; 26.6% were faculty/staff. The median days from end of isolation to the follow-up survey was 57 QR (39–158) days.

Most respondents had no underlying conditions (75.2%), never smoked (83.0%), had acute COVID-19 symptoms (79.1%), did not seek medical care at the

time of their first positive COVID-19 result (96.6%), and did not receive monoclonal antibody treatment (94.5%) (Table 2). Approximately 41.5% of respondents had received a booster vaccine, 29.9% were fully vaccinated with an initial vaccine series, and 28.6% were not fully vaccinated at the time of their first positive COVID-19 test result. The most common acute symptom was upper respiratory (e.g., congestion, cough, sore throat, runny nose) (92.0%), followed by headache (51.2%), fatigue (51.1%), and chills/fever (44.9%).

Nearly 36% of survey respondents reported experiencing symptoms of long COVID (Table 2). Respondents who had underlying conditions (44.7%; $p = 0.003$), who were not fully vaccinated (47.7%; $p < 0.0001$), who were female (40.9%; $p = 0.002$), who were former/current smokers (45.3%; $p = 0.028$), who experienced acute COVID-19 symptoms (43.3%; $p < 0.0001$), who reported higher symptom counts (mean 5; $p < 0.0001$), who sought medical care (73.9%; $p < 0.0001$), or who received antibody treatment (72.0%; $p = 0.0012$) were significantly more likely to report symptoms of long COVID. All symptom categories were strongly associated with long COVID status, except for upper respiratory and other symptoms (Table 2; Figure).

Unadjusted associations between characteristics of survey respondents and long COVID status showed that sex, race/ethnicity, underlying conditions, smoking status, vaccination status, any symptoms, symptom type, symptom count, seeking out medical care, and receiving antibody treatment were strongly associated with long COVID (Table 3). Multivariable models adjusting for statistically significant characteristics in the bivariate analyses found several significant associations: smoking history (former/current smokers versus never smokers) (model 1: adjusted odds ratio [aOR] 1.59, 95% CI 1.13–2.25); experiencing any symptoms at the time of positive test (model 1: aOR 1.92, 95% CI 1.01–3.62); experiencing fatigue (model 1: aOR 1.80, 95% CI 1.32–2.47); and experiencing chest pain/shortness of breath (model 1: aOR 2.18, 95% CI 1.48–3.22). Immunization status was significantly associated with long COVID; those fully vaccinated had higher odds of long COVID than those who had also received a booster (model 1: aOR 2.10, 95% CI 1.51–2.90), and those who were not fully vaccinated had higher odds than those fully vaccinated and those given a booster (model 1: aOR 2.71, 95% CI 1.94–3.77). We found similar results after using symptom count in lieu of any symptoms (versus no symptoms) in model 2.

Table 2. Characteristics of survey respondents by long COVID status for postacute sequelae of SARS-CoV-2 in university setting, Washington, DC, USA*

Characteristic	Total, n = 1,338	No long COVID, n = 833	Long COVID,† n = 475	p value
Age, y median (IQR)	23 (21–32)	23 (21–33)	23 (21–30)	0.265
Sex, n = 1,327				0.002
F	841 (63.4)	497 (59.1)	344 (40.9)	
M	486 (36.6)	328 (67.5)	158 (31.5)	
Race/ethnicity, n = 1,319				0.089
Non-Hispanic White	734 (55.7)	439 (59.8)	295 (40.2)	
Asian	175 (13.3)	119 (68.0)	56 (32.0)	
Non-Hispanic Black	175 (13.3)	117 (66.9)	58 (33.1)	
Hispanic	117 (8.9)	77 (65.8)	40 (34.2)	
Other	70 (5.3)	37 (52.9)	33 (47.1)	
Multiracial	48 (3.6)	32 (66.7)	16 (33.3)	
Affiliation				0.115
Students	982 (73.4)	599 (61.0)	383 (39.0)	
Faculty/staff	356 (26.6)	234 (65.7)	122 (34.3)	
Underlying conditions‡, n = 1,262				0.003
No	949 (75.2)	613 (64.6)	336 (35.4)	
Yes	313 (24.8)	173 (55.3)	140 (44.7)	
Smoking status, n = 1,259				0.028
Never	1,045 (83.0)	655 (62.7)	390 (37.3)	
Former/current	214 (17.0)	117 (54.7)	97 (45.3)	
Vaccination status at time of positive test result				<0.0001
Fully vaccinated with booster	555 (41.5)	411 (74.1)	144 (25.9)	
Fully vaccinated	400 (29.9)	213 (55.6)	170 (44.4)	
Not fully vaccinated	383 (28.6)	209 (52.3)	191 (47.7)	
Any symptoms at time of positive test result, n = 1,328				<0.0001
No	278 (20.9)	231 (83.1)	47 (16.9)	
Yes	1,050 (79.1)	595 (56.7)	455 (43.3)	
Symptom type at time of positive test result, n = 1,050				
Congestion/cough/sore throat/runny nose	966 (92.0)	549 (56.8)	417 (43.2)	0.713
Headache	538 (51.2)	275 (51.1)	263 (48.9)	0.0002
Fatigue	537 (51.1)	256 (47.7)	281 (52.3)	<0.0001
Chills/measured fever/subjective fever	471 (44.9)	240 (51.0)	231 (49.0)	0.0008
Muscle pain	326 (31.1)	148 (45.4)	178 (54.6)	<0.0001
Chest pain/shortness of breath	189 (18.0)	67 (35.5)	122 (64.5)	<0.0001
Diarrhea/nausea/vomiting	181 (17.2)	78 (43.1)	103 (56.9)	<0.0001
Loss of taste/smell	180 (17.1)	86 (47.8)	94 (52.2)	0.008
Other	59 (5.6)	28 (47.5)	31 (52.5)	0.142
Symptom count, median (IQR)	4 (1–6)	3 (0–5)	5 (3–8)	<0.0001
Sought medical care, n = 1,336				<0.0001
No	1,290 (96.6)	819 (63.5)	471 (36.5)	
Yes	46 (3.4)	12 (26.1)	34 (73.9)	
Received monoclonal antibodies, n = 1,331				0.0012
No	1,258 (94.5)	788 (62.6)	470 (37.4)	
Unknown	48 (3.6)	33 (68.8)	15 (31.2)	
Yes	25 (1.9)	7 (28.0)	18 (72.0)	

*Values are no. (%) except as indicated. IQR, interquartile range.

†Long COVID was defined as experiencing ≥1 of the following symptoms lasting for >28 d after a respondent’s 10-day isolation period ended: difficulty driving, difficulty having conversations, difficulty making decisions, difficulty thinking, fatigue, feeling anxious, feeling depressed or sad, loss of smell, loss of taste, memory loss, muscle pain, muscle weakness, shortness of breath or difficulty breathing, trouble sleeping, worsening of symptoms after physical activity, worsening of symptoms after mental activity, or other symptoms.

‡Includes diabetes, asthma, hypertension, obesity, sickle cell disease, cancer, chronic kidney disease, lung diseases, serious heart conditions, and other conditions.

Discussion

This study aimed to examine the prevalence and predictors of long COVID in a university community. This sample was unique in that it consisted of primarily young adults who had few underlying health conditions and otherwise were considered healthy. Regardless of initial symptoms, nearly 36% of COVID-19 survivors in this study reported experiencing symptoms consistent with long COVID. That

result is within ranges found in other studies reporting a prevalence of long COVID of anywhere from 10% to 80% among COVID-19 survivors (3–5,7,21,29–31). Our study also found an increased odds of reporting symptoms consistent with long COVID for each additional symptom reported during the initial infection. This finding is consistent with recent studies conducted with a high proportion of young adults that also found a higher number of acute symptoms

during a COVID-19 infection predicted ≥ 1 long COVID symptom (32). Monitoring symptoms of initial cases could help identify persons at risk for long COVID.

Our study also found that persons who had the fewest previous COVID-19 vaccines and boosters were at higher risk for development of symptoms consistent with long COVID, supporting other investigations suggesting that vaccination is associated with reduced risk for long COVID (33–36). Many colleges and universities required the COVID-19 vaccine before the fall 2021 semester but offered reasonable medical/religious exemptions. Our results further highlight the need for routine short- and long-term follow-up for persons who test positive for COVID-19 while continuing to advocate and monitor for vaccine and booster adherence to published recommendations.

Although prevention efforts are needed for long COVID, the findings from this study support the need to ameliorate consequences of long COVID. Based on symptomatology, recovery strategies for long COVID include physical rehabilitation, management of preexisting conditions, mental health support, social services support, and exercise programs scaled to the ability of the patient (11,37). Because long COVID can greatly

interfere with the ability to learn or work, classroom or job accommodations, such as modifying academic and workplace policies, flexible scheduling, changing workplace environment, enabling remote or alternative learning, and modifying job responsibilities, are recommended for those having long COVID.

Limitations in conducting this study included the possibility of recall bias, loss to follow-up, and digital literacy challenges, as well as acknowledgment that the results are only for persons who tested positive for SARS-CoV-2. Persons were asked to recall information about their illness after ≥ 28 days had passed. Considering brain fog is a symptom consistent with long COVID and the length of time between isolation and follow-up, some persons who had long COVID might have forgotten details of their health status during a tumultuous time in their life. Although inevitable, this situation was mitigated by providing the person with dates of their illness when asking them to think back to that time.

Loss to follow-up was also a limitation; some persons never completed a case investigation, which made it more likely for them to forgo a follow-up months later. CCST made >3 attempts at different time points throughout the day to reach as many persons as possible. Those strategies, and our achieved response rate, are consistent with other COVID-19 studies conducted during the pandemic (38). Nonetheless, we acknowledge that results could be inflated because persons experiencing symptoms consistent with long COVID might be more likely to respond. Thus, results should be interpreted with caution.

In addition, surveys were conducted by electronic survey and telephone. Although there were no major differences in demographics between telephone and electronic survey completion, some of our participants did not have smartphones, only had landlines, or could not be reached by email, which contributed to loss to follow-up.

Finally, our sample was only of persons who had COVID-19 within our campus community and not of the entire campus population. Thus, it is not possible to know whether symptoms reported in our survey were also increased in the campus population as a whole during this time. Many of the symptoms in our survey are common and might or might not be directly related to SARS-CoV-2 infection or long COVID.

Public health experts and healthcare providers have been gathering data about COVID-19 while simultaneously trying to understand the long-term consequences of SARS-CoV-2 infection. Although preliminary findings of long COVID were anecdotal, researchers continue to gain a clearer picture

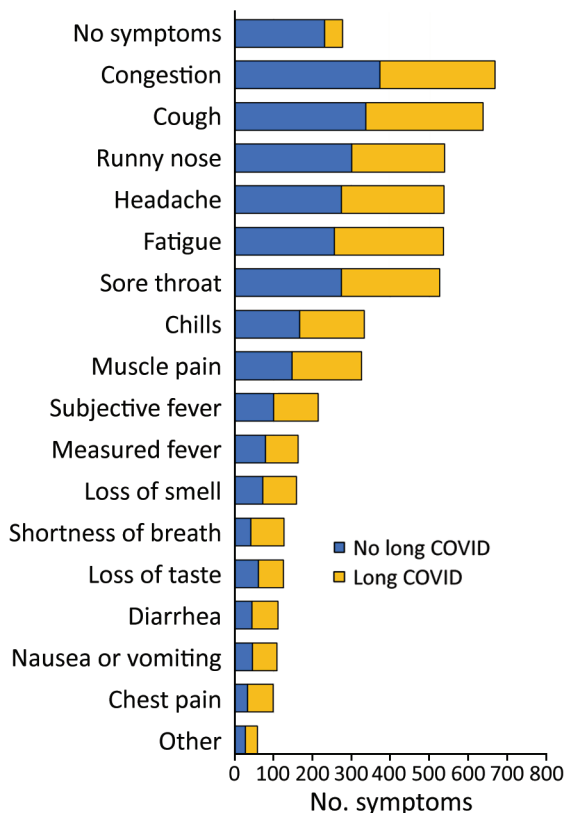


Figure. Frequency of reported acute symptoms among survey respondents for postacute sequelae of SARS-CoV-2 in university setting, by long COVID status, Washington, DC, USA (n = 1,338).

Table 3. Association between characteristics of survey respondents and long COVID for postacute sequelae of SARS-CoV-2 in university setting, Washington, DC, USA*

Characteristic	Unadjusted OR (95% CI), n = 1,338	Model 1, aOR† (95% CI), n = 1,172	Model 2, aOR† (95% CI), n = 1,175
Age, y	0.99 (0.98–1.00)	–	–
Sex, n = 1,327			
F	1.44 (1.14–1.82)	1.22 (0.92–1.62)	1.16 (0.88–1.54)
M	Referent	Referent	Referent
Race/ethnicity, n = 1,319			
Non-Hispanic White	Referent	Referent	Referent
Asian	0.70 (0.49–0.99)	0.97 (0.64–1.46)	1.00 (0.66–1.50)
Non-Hispanic Black	0.74 (0.52–1.04)	0.79 (0.53–1.19)	0.81 (0.54–1.21)
Hispanic	0.77 (0.51–1.16)	0.82 (0.52–1.19)	0.85 (0.53–1.34)
Other	1.33 (0.81–2.17)	1.54 (0.87–2.74)	1.48 (0.84–2.63)
Multiracial	0.74 (0.40–1.38)	0.78 (0.39–1.59)	0.81 (0.40–1.64)
Affiliation			
Students	1.23 (0.95–1.58)	–	–
Faculty/staff	Referent	–	–
Underlying conditions,‡ n = 1,262			
No	Referent	Referent	Referent
Yes	1.48 (1.14–1.91)	1.23 (0.90–1.66)	1.27 (0.94–1.73)
Smoking status, n = 1,259			
Never	Referent	Referent	Referent
Former/current	1.39 (1.04–1.87)	1.59 (1.13–2.25)	1.60 (1.13–2.25)
Vaccination status at time of positive test result			
Fully vaccinated with booster	Referent	Referent	Referent
Fully vaccinated	2.28 (1.73–3.01)	2.10 (1.51–2.90)	2.19 (1.58–3.03)
Not fully vaccinated	2.61 (1.99–3.43)	2.71 (1.94–3.77)	3.01 (2.16–4.21)
Any symptoms at time of positive test result, n = 1,328			
No	Referent	Referent	–
Yes	3.76 (2.68–5.26)	1.92 (1.04–3.62)	–
Symptom type at time of positive test result			
Fatigue	2.83 (2.25–3.56)	1.80 (1.32–2.47)	1.53 (1.05–2.22)
Chest pain/shortness of breath	3.64 (2.64–5.03)	2.18 (1.48–3.22)	1.70 (1.08–2.67)
Congestion/cough/sore throat/runny nose	2.45 (1.87–3.21)	0.96 (0.58–1.60)	1.07 (0.69–1.65)
Chills/measured fever/subjective fever	2.08 (1.65–2.62)	0.93 (0.69–1.26)	0.73 (0.49–1.08)
Headache	2.21 (1.76–2.77)	1.11 (0.82–1.50)	0.97 (0.69–1.37)
Loss of taste/smell	1.99 (1.45–2.73)	1.04 (0.72–1.49)	0.83 (0.53–1.29)
Muscle pain	2.52 (1.95–3.25)	1.25 (0.91–1.73)	1.10 (0.78–1.57)
Diarrhea/nausea/vomiting	2.48 (1.80–3.41)	1.24 (0.85–1.81)	1.02 (0.66–1.56)
Other	1.88 (1.11–3.17)	0.89 (0.48–1.64)	0.83 (0.44–1.55)
Symptom count	1.22 (1.18–1.27)	–	1.16 (1.00–1.33)
Sought out medical care, n = 1,336			
No	Referent	Referent	Referent
Yes	4.93 (2.53–9.61)	2.17 (0.98–4.77)	2.07 (0.94–4.55)
Received monoclonal antibodies, n = 1,331			
No	Referent	Referent	Referent
Unknown	0.76 (0.41–1.42)	0.74 (0.34–1.64)	0.72 (0.33–1.59)
Yes	4.31 (1.79–10.40)	1.93 (0.63–5.96)	2.06 (0.68–6.23)

*Long COVID was defined as experiencing ≥1 of the following symptoms lasting for >28 d after a respondent's 10-day isolation period ended: difficulty driving, difficulty having conversations, difficulty making decisions, difficulty thinking, fatigue, feeling anxious, feeling depressed or sad, loss of smell, loss of taste, memory loss, muscle pain, muscle weakness, shortness of breath or difficulty breathing, trouble sleeping, worsening of symptoms after physical activity, worsening of symptoms after mental activity, or other symptoms. aOR, adjusted odds ratio; IQR, interquartile range; OR, odds ratio; –, variable omitted for that model.

†Adjusted for all other variables in the column.

‡Includes diabetes, asthma, hypertension, obesity, sickle cell disease, cancer, chronic kidney disease, lung diseases, serious heart conditions, and other conditions.

on who it affects and how it affects certain populations. From a university standpoint, this analysis is key to understanding how administration can fill the needs of the campus population that has long-term complications caused by COVID-19. Paired with the recommendations presented in this article, universities can strengthen existing services or make referrals to prevention and rehabilitation services (i.e., mental health, exercise programs, long-term health studies)

for those who have long COVID that affects their ability to engage in university activities such as classes and work. In addition, universities might benefit from adopting preventive resources for their populations, as well as extended pandemic leave, given the considerable long-lasting effects of long COVID.

Future research avenues should consider following up with long COVID survivors/patients to assess long-term or long-lasting symptoms. Such

analysis could explore the consequences of long COVID for 5–10 years after the initial infection, especially to gain a better understanding of its effect on young, healthy populations. Follow-up could also occur with older populations to assess whether symptoms progress into retirement age and to determine the cost of long-term care resulting from long COVID. Furthermore, research should continue to examine the effect vaccine booster doses have on long COVID symptoms. Such research is vital to clarifying long-term effects of long COVID and how universities can support those dealing with long COVID to promote health and wellness across campus communities.

Acknowledgments

We appreciate our campus community and strive to continue serving our community members. We thank the entire Campus COVID-19 Support Team and their tireless efforts in keeping our community safe. We could not have had a successful return to campus and ongoing surveillance without our team. We also thank the staff of our partners in the Public Health Laboratory, the Student Health Center, and the employee Occupational Health department for their partnership in the successful implementation of the campus surveillance program and their dedication to striving to keep our campus safe and healthy.

About the Author

Dr. Landry is the project director for the Campus COVID-19 Support Team at the George Washington University, Washington, DC. Her primary research interests are public health surveillance and maternal and child health.

References

- Calabrese L. Long COVID: what do we know and what should we do? Presented at: Basic and Clinical Immunology for the Busy Clinician; February 26, 2022 (virtual meeting) [cited 2022 Dec 15]. <https://www.clevelandclinicmeded.com/live/courses/Clinical22/agenda.asp>
- Centers for Disease Control and Prevention. National Center for Immunization and Respiratory Diseases (NCIRD), Division of Viral Diseases. Post-COVID conditions, September 16, 2021 [cited 2022 Dec 15]. <https://www.cdc.gov/coronavirus/2019-ncov/long-term-effects/index.html>
- Berger Z, Altiery DE, Jesus V, Assoumou SA, Greenhalgh T. Long COVID and health inequities: the role of primary care. *Milbank Q*. 2021;99:519–41. <https://doi.org/10.1111/1468-0009.12505>
- Jason LA, Islam M, Conroy K, Cotler J, Torres C, Johnson M, et al. COVID-19 symptoms over time: comparing long-haulers to ME/CFS. *Fatigue*. 2021;9:59–68. <https://doi.org/10.1080/21641846.2021.1922140>
- Leviner S. Recognizing the clinical sequelae of COVID-19 in adults: COVID-19 long-haulers. *J Nurse Pract*. 2021;17:946–9. <https://doi.org/10.1016/j.nurpra.2021.05.003>
- Parums DV. Editorial: long COVID, or post-COVID syndrome, and the global impact on health care. *Med Sci Monit*. 2021;27:e933446. <https://doi.org/10.12659/MSM.933446>
- van Kessel SA, Olde Hartman TC, Lucassen PL, van Jaarsveld CH. Post-acute and long-COVID-19 symptoms in patients with mild diseases: a systematic review. *Fam Pract*. 2022;39:159–67. <https://doi.org/10.1093/fampra/cmab076>
- Soriano JB, Murthy S, Marshall JC, Relan P, Diaz JV; WHO Clinical Case Definition Working Group on Post-COVID-19 Condition. A clinical case definition of post-COVID-19 condition by a Delphi consensus. *Lancet Infect Dis*. 2022;22:e102–7. [https://doi.org/10.1016/S1473-3099\(21\)00703-9](https://doi.org/10.1016/S1473-3099(21)00703-9)
- Phillips S, Williams MA. Confronting our next national health disaster: long-haul COVID. *N Engl J Med*. 2021;385:577–9. <https://doi.org/10.1056/NEJMp2109285>
- Lambert, NJ; Survivor Corps. COVID-19 “long hauler” symptoms survey report, 2020. Indiana University School of Medicine [cited 2022 Dec 15]. <https://dig.abclocal.go.com/wls/documents/2020/072720-wls-covid-symptom-study-doc.pdf>
- Aiyegbusi OL, Hughes SE, Turner G, Rivera SC, McMullan C, Chandan JS, et al.; TLC Study Group. Symptoms, complications and management of long COVID: a review. *J R Soc Med*. 2021;114:428–42. <https://doi.org/10.1177/01410768211032850>
- Iqbal FM, Lam K, Sounderajah V, Clarke JM, Ashrafian H, Darzi A. Characteristics and predictors of acute and chronic post-COVID syndrome: a systematic review and meta-analysis. *EClinicalMedicine*. 2021;36:100899. <https://doi.org/10.1016/j.eclinm.2021.100899>
- Anaya JM, Rojas M, Salinas ML, Rodríguez Y, Roa G, Lozano M, et al.; Post-COVID study group. Post-COVID syndrome. A case series and comprehensive review. *Autoimmun Rev*. 2021;20:102947. <https://doi.org/10.1016/j.autrev.2021.102947>
- Mahmud R, Rahman MM, Rassel MA, Monayem FB, Sayeed SKJB, Islam MS, et al. Post-COVID-19 syndrome among symptomatic COVID-19 patients: a prospective cohort study in a tertiary care center of Bangladesh. *PLoS One*. 2021;16:e0249644. <https://doi.org/10.1371/journal.pone.0249644>
- Michelen M, Manoharan L, Elkheir N, Cheng V, Dagens A, Hastie C, et al. Characterising long COVID: a living systematic review. *BMJ Glob Health*. 2021;6:e005427. <https://doi.org/10.1136/bmjgh-2021-005427>
- Nalbandian A, Sehgal K, Gupta A, Madhavan MV, McGroder C, Stevens JS, et al. Post-acute COVID-19 syndrome. *Nat Med*. 2021;27:601–15. <https://doi.org/10.1038/s41591-021-01283-z>
- Sathyamurthy P, Madhavan S, Pandurangan V. Prevalence, pattern and functional outcome of post COVID-19 syndrome in older adults. *Cureus*. 2021;13:e17189.
- Sudre CH, Murray B, Varsavsky T, Graham MS, Penfold RS, Bowyer RC, et al. Attributes and predictors of long COVID. *Nat Med*. 2021;27:626–31. <https://doi.org/10.1038/s41591-021-01292-y>
- Taquet M, Dercon Q, Luciano S, Geddes JR, Husain M, Harrison PJ. Incidence, co-occurrence, and evolution of long-COVID features: a 6-month retrospective cohort study of 273,618 survivors of COVID-19. *PLoS Med*. 2021;18:e1003773. <https://doi.org/10.1371/journal.pmed.1003773>

20. Tirelli U, Taibi R, Chirumbolo S. Post COVID syndrome: a new challenge for medicine. *Eur Rev Med Pharmacol Sci*. 2021;25:4422-5.
21. Berg S. What doctors wish patients knew about long COVID, October 22, 2021. American Medical Association [cited 2022 Dec 15]. <https://www.ama-assn.org/delivering-care/public-health/what-doctors-wish-patients-knew-about-long-covid>
22. Crispo A, Bimonte S, Porciello G, Forte CA, Cuomo G, Montagnese C, et al. Strategies to evaluate outcomes in long-COVID-19 and post-COVID survivors. *Infect Agent Cancer*. 2021;16:62. <https://doi.org/10.1186/s13027-021-00401-3>
23. Singh I, Joseph P, Heerdt PM, Cullinan M, Lutchmansingh DD, Gulati M, et al. Persistent exertional intolerance after COVID-19. *Chest*. 2022;161:54-63. <https://doi.org/10.1016/j.chest.2021.08.010>
24. Sojka A, Machniak M, Andrzejewski W, Kosendiak A, Chwałczyńska A. Changes in physical activity and the occurrence of specific symptoms of “long-COVID syndrome” in men aged 18–25. *Int J Environ Res Public Health*. 2022;19:1199. <https://doi.org/10.3390/ijerph19031199>
25. Lemhöfer C, Sturm C, Loudovici-Krug D, Best N, Gutenbrunner C. The impact of post-COVID-syndrome on functioning: results from a community survey in patients after mild and moderate SARS-CoV-2-infections in Germany. *J Occup Med Toxicol*. 2021;16:45. <https://doi.org/10.1186/s12995-021-00337-9>
26. US Food and Drug Administration. In vitro diagnostic. EUAs: antigen diagnostic tests for SARS-CoV-2, 2021 [cited 2022 Dec 15]. <https://www.fda.gov/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/in-vitro-diagnostics-euas-antigen-diagnostic-tests-sars-cov-2>
27. Towers SA, Gemechu NB, Nagaraj NC, Landry MM, Beane P, Sardon GA Jr, et al.; George Washington University Campus COVID-19 Support Team. SARS-CoV-2 surveillance and outbreak response on an urban American college campus. *J Am Coll Health*. 2022;Mar 8:1-9. <https://doi.org/10.1080/07448481.2022.2034834>
28. Harris PA, Taylor R, Thielke R, Payne J, Gonzalez N, Conde JG. Research electronic data capture (REDCap): a metadata-driven methodology and workflow process for providing translational research informatics support. *J Biomed Inform*. 2009;42:377-81. <https://doi.org/10.1016/j.jbi.2008.08.010>
29. Lewis D. Long COVID and kids: scientists race to find answers. *Nature*. 2021;595:482-3. <https://doi.org/10.1038/d41586-021-01935-7>
30. Lopez-Leon S, Wegman-Ostrosky T, Perelman C, Sepulveda R, Rebolledo PA, Cuapio A, et al. More than 50 long-term effects of COVID-19: a systematic review and meta-analysis. *Sci Rep*. 2021;11:16144. <https://doi.org/10.1038/s41598-021-95565-8>
31. Silverberg JI, Zyskind I, Naiditch H, Zimmerman J, Glatt AE, Pinter A, et al. Predictors of chronic COVID-19 symptoms in a community-based cohort of adults. *PLoS One*. 2022;17:e0271310. <https://doi.org/10.1371/journal.pone.0271310>
32. Ebell MH, Forgacs D, Shen Y, Ross TM, Hulme C, Bentivegna M, et al. High prevalence of both previous infection with SARS-CoV-2 and persistent symptoms. *J Am Board Fam Med*. 2022;35:570-8. <https://doi.org/10.3122/jabfm.2022.03.210348>
33. Antonelli M, Penfold RS, Merino J, Sudre CH, Molteni E, Berry S, et al. Risk factors and disease profile of post-vaccination SARS-CoV-2 infection in UK users of the COVID Symptom Study app: a prospective, community-based, nested, case-control study. *Lancet Infect Dis*. 2022; 22:43-55. [https://doi.org/10.1016/S1473-3099\(21\)00460-6](https://doi.org/10.1016/S1473-3099(21)00460-6)
34. Kuodi P, Gorelik Y, Zayyad H, Wertheim O, Wiegler KB, Jabal KA, et al. Association between BNT162b2 vaccination and reported incidence of post-COVID-19 symptoms: cross-sectional study 2020-21, Israel. *NPJ Vaccines*. 2022;7:101. <https://doi.org/10.1038/s41541-022-00526-5>
35. Hastie CE, Lowe DJ, McAuley A, Winter AJ, Mills NL, Black C, et al. Outcomes among confirmed cases and a matched comparison group in the Long-COVID in Scotland study. *Nat Commun*. 2022;13:5663. <https://doi.org/10.1038/s41467-022-33415-5>
36. Ayoubkhani D, Bermingham C, Pouwels KB, Glickman M, Nafilyan V, Zaccardi F, et al. Trajectory of long covid symptoms after covid-19 vaccination: community based cohort study. *BMJ*. 2022;377:e069676. <https://doi.org/10.1136/bmj-2021-069676>
37. Morley JE. COVID-19: the long road to recovery. *J Nutr Health Aging*. 2020;24:917-9. <https://doi.org/10.1007/s12603-020-1497-y>
38. Amaral S, Dinarte L, Dominguez P, Romero S, Perez-Vincent SM. Talk or text? Evaluating response rates by remote survey method during COVID-19. CESifo working paper no. 9517. 2022 [cited 2022 Dec15]. <https://doi.org/10.2139/ssrn.4009417>

Address for correspondence: Megan Landry, The George Washington University Milken Institute School of Public Health, 950 New Hampshire Ave NW, 7th Fl, Washington, DC 20052, USA; email: mmlandry@gwu.edu

Postacute Sequelae of SARS-CoV-2 in University Setting

Appendix 1.

Appendix 1 Table. Characteristics of follow-up survey respondents versus nonrespondents with a completed case interview*

Characteristic	Total, no. (%), (n = 3,595)	Nonrespondents, no. (%), (n = 2,254)	Respondents, no. (%), (n = 1,341)	p value
Age, y, median (IQR)	22 (20–27)	22 (20–26)	23 (21–32)	<0.0001
Sex (n = 3,546)				0.0464
Female	2,173 (61.3)	1,330 (60.0)	834 (63.4)	
Male	1,373 (38.7)	886 (39.9)	487 (36.6)	
Race/Ethnicity (n = 3,538)				0.9185
Non-Hispanic White	1,986 (56.1)	1,251 (56.5)	735 (55.6)	
Asian	464 (13.1)	289 (13.9)	175 (13.2)	
Non-Hispanic Black	443 (12.5)	268 (12.1)	175 (13.2)	
Hispanic	314 (8.9)	195 (8.8)	119 (9.0)	
Other	196 (5.4)	126 (5.7)	70 (5.3)	
Multiracial	135 (3.8)	87 (3.9)	48 (3.6)	
Affiliation				<0.0001
Student	2,930 (81.5)	1,946 (86.3)	984 (73.4)	
Faculty/staff	665 (18.5)	308 (13.7)	357 (26.6)	
Any underlying condition† (n = 3,383)				<0.0001
No	2,668 (78.9)	1,717 (81.1)	951 (75.2)	
Yes	715 (21.1)	401 (18.9)	314 (24.8)	
Smoking status				0.7135
Never	2,876 (83.1)	1,794 (82.9)	1,082 (83.4)	
Current/former	584 (16.9)	369 (17.1)	215 (16.6)	
Vaccination status at time of test (n = 3,592)				<0.0001
Fully vaccinated with booster	1,404 (39.1)	849 (37.7)	555 (41.4)	
Fully vaccinated	1,319 (36.7)	935 (41.5)	384 (28.6)	
Not fully vaccinated	869 (24.2)	467 (20.8)	402 (29.9)	
Any symptoms at time of test‡ (n = 3,574)				0.8788
No	754 (21.1)	475 (21.2)	279 (20.9)	
Yes	2,820 (78.9)	1,768 (78.8)	1,052 (79.0)	
Symptom count, mean (IQR)	4 (1–6)	4 (1–6)	4 (1–6)	0.9871

*IQR, interquartile range.

†Includes diabetes, asthma, hypertension, obesity, sickle cell disease, cancer, chronic kidney disease, lung diseases, serious heart conditions, and other conditions.

‡Includes chest pain, chills, congestion, cough, diarrhea, fatigue, fever, headache, loss of smell, loss of taste, muscle pain, nausea or vomiting, runny nose, shortness of breath, sore throat, and other symptoms.

Article DOI: <https://doi.org/10.3201/eid2903.221522>

EID cannot ensure accessibility for supplementary materials supplied by authors. Readers who have difficulty accessing supplementary content should contact the authors for assistance.

Postacute Sequelae of SARS-CoV-2 in University Setting

Appendix 2

The following pages show the script for the long COVID survey, a follow-up telephone interview given to 4,800 persons with COVID-19 cases identified by The George Washington University COVID-19 surveillance and testing program during August 2020–February 2022. The surveys were administered by The George Washington University Campus COVID-19 Support Team during July 2021–March 2022.

COVID-19 Follow-up Survey

Dear [first_name],

Thank you for participating in this confidential survey. We are aiming to better understand the potential long-term effects and health behaviors surrounding COVID-19 in the GWU community. All of your responses will be grouped together for any reporting purposes and your identity will remain confidential. This survey should take 5-10 minutes to complete.

Thank you for your time!

This set of questions asks about your experience during the two weeks after you tested positive for COVID-19.

SCRIPT:

This first set of questions asks about symptoms you may have experienced DURING the two weeks after you tested positive.

I see that you tested positive for COVID-19 on [lab_pos_date]. I am going to run through a list of symptoms quickly. Please say "yes" for any you experienced during the two weeks after you tested positive.

You tested positive for COVID-19 on [lab_pos_date] (m/d/y). Thinking back to that time, did you experience any of the following symptoms during the two weeks after you tested positive? (Select all that apply)

- Difficulty driving
- Difficulty having conversations
- Difficulty following instructions
- Difficulty making decisions
- Difficulty thinking (known as "brain fog")
- Fatigue
- Feeling anxious
- Feeling depressed or sad
- Loss of smell
- Loss of taste
- Memory loss
- Muscle pain
- Muscle weakness
- Shortness of breath or difficulty breathing
- Trouble sleeping
- Worsening of symptoms after physical activity (e.g., walking, swimming, running, etc.)
- Worsening of symptoms after mental activity (e.g., work, attending school)
- Other, please specify
- No symptoms

Other symptoms:

Of the symptoms selected, which symptom was the most bothersome to you ?
(Choose only one symptom)

- Difficulty driving
- Difficulty having conversations
- Difficulty following instructions
- Difficulty making decisions
- Difficulty thinking (known as "brain fog")
- Fatigue
- Feeling anxious
- Feeling depressed or sad
- Loss of smell
- Loss of taste
- Memory loss
- Muscle pain
- Muscle weakness
- Shortness of breath or difficulty breathing
- Trouble sleeping
- Worsening of symptoms after physical activity (e.g., walking, swimming, running, etc.)
- Worsening of symptoms after mental activity (e.g., work, attending school)
- Other, please specify

Other symptoms:

After you tested positive for COVID-19, did you ever have to seek medical care in the emergency room or urgent care?

- Yes
- No

After you tested positive for COVID-19, were you ever hospitalized (i.e., had to stay overnight in the hospital)?

- Yes
- No

Did you receive any monoclonal antibodies to treat your COVID-19? (Antibodies assist the immune system to respond more effectively to the virus.)

- Yes
- No
- Unknown

Do you have any of the following conditions?
(Select all that apply)

- Alcohol or substance use disorder
- Asthma
- Cancer
- Cardiovascular diseases (heart attack/failure, stroke, etc.)
- Chronic kidney diseases
- Chronic lung diseases (COPD)
- Depression
- Diabetes
- Hepatitis B virus (HBV)
- Hepatitis C virus (HCV)
- HIV
- Hypertension (high blood pressure)
- Tuberculosis (TB)
- Other mental health condition
- Other chronic condition (specify)
- No underlying conditions

Other conditions:

Are you currently pregnant?

- Yes
- No

SCRIPT:

Thank you for that information.

The next set of questions are about behaviors before you tested positive and during your 10-day isolation period. The answers to these questions are on a scale from...

(READ SLOWLY)

- Rarely or none of the time
- Some or little of the time
- Occasionally or a moderate amount of time
- Most or all of the time

The next set of questions asks about behaviors before you tested positive and during your 10-day isolation period.

	Rarely or none of the time	Some or little of the time	Occasionally or a moderate amount of time	Most or all of the time
BEFORE becoming infected, how often did you practice social distancing? (social distancing = staying 3-6 feet away from other people when out in public or in social settings such as work or school)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
BEFORE becoming infected, how often did you consistently wear any PPE (masks, N95, face shields, etc.) when around others outside of your immediate household members?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
BEFORE testing positive, how often did you stay at home as much as possible?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
AFTER you tested positive, how often did you self-isolate or quarantine for the recommended/required number of days? (i.e., you separated yourself from other people, even those in your own household, to prevent others from getting sick)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

The next set of questions asks about symptoms you may have experienced after your 10-day isolation period ended.

SCRIPT:

The next set of questions are about symptoms AFTER your 10-day isolation period ENDED.

Please think back to the time AFTER your isolation ENDED. Have you had any symptoms that lasted more than 28 days? I am going to run through a list of symptoms quickly again. Please say "yes" for any you experienced during the two weeks after you tested positive.

AFTER your 10-day isolation period ENDED, have you had any of the following symptoms [lasting for more than 28 days after your 10-day isolation period ended]?

(Select all that apply)

- Difficulty driving
- Difficulty having conversations
- Difficulty following instructions
- Difficulty making decisions
- Difficulty thinking (known as "brain fog")
- Fatigue
- Feeling anxious
- Feeling depressed or sad
- Loss of smell
- Loss of taste
- Memory loss
- Muscle pain
- Muscle weakness
- Shortness of breath or difficulty breathing
- Trouble sleeping
- Worsening of symptoms after physical activity (e.g., walking, swimming, running, etc.)
- Worsening of symptoms after mental activity (e.g., work, attending school)
- Other, please specify
- No symptoms

Other symptoms: _____

Have you had to reduce your hours at work or at school due to persistent COVID-19 symptoms AFTER your 10-day isolation period?

- Yes
- No

Are you still having any COVID-19 related symptoms?

- I have recovered and am symptom free
- I am feeling better but not completely recovered (i.e., have COVID-19 related symptoms that remain)
- I am not feeling better

SCRIPT:

The answers to the next question are on a scale from...

- Not at all impacted
- A little impacted
- Moderately impacted
- Extremely impacted

Overall, considering all the possible ways your life may have been affected because of persistent COVID-19 symptoms, how much has this impacted your day-to-day life?

- Not at all impacted
- A little impacted
- Moderately impacted
- Extremely impacted

Have you received the COVID-19 vaccine? Yes
 No

Are you fully vaccinated? (e.g., 1 dose of J&J or 2 doses of AstraZeneca/Moderna/Pfizer/etc.) Yes
 No

Did your symptoms improve at all after receiving the vaccine? Yes
 No

The final set of questions asks about the impact COVID-19 has had on you, family, and friends.

SCRIPT:

The answers to the next question are on a scale from...

- Not at all concerned
- A little concerned
- Moderately concerned
- Extremely concerned

How concerned are you about becoming re-infected with COVID-19? Not at all concerned
 A little concerned
 Moderately concerned
 Extremely concerned

How many people do you know who have been diagnosed with COVID-19 ? _____ people _____

We are interested in learning more about the lasting effects of COVID-19 among the GWU population. May we contact you for additional information about your experience? Yes
 No

SCRIPT:

Thank you for participating in this confidential survey. As a reminder...

- Everything that we talk about will be kept confidential.
- Your responses will be grouped with others to hide your identity.
- This information will not impact your GW status.
- The information you've shared will be very helpful in understanding the long term effects of COVID-19.
- You will receive an email with a list of resources if you or someone else needs them.

Do you have any final questions?

Thank you for your time!

*****IMPORTANT*****

Select: SAVE & MARK SURVEY AS COMPLETE (Do not press SAVE & EXIT FORM)