



COVID-19 & ADULTS WITH DIFFICULTIES READING, WRITING, OR COMMUNICATING

ONLINE SURVEY REPORT

American Association on Health & Disability

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A. Introduction

The purpose of the COVID-19 & Disability (C-19 & D) Survey was to conduct a rapid, real-time online assessment of the coronavirus pandemic's impact on access to health care and on adults with a range of disabilities, including people with serious difficulties concentrating, remembering, or making decisions due to difficulties reading, writing, or communicating. Online data-collection technologies permit efficient, rapid assessments that, despite limitations, are useful to begin understanding public health issues. People with disabilities experience unique risks during emergencies, disasters, epidemics, and pandemics that can impact health and health care. *Yet, at this time, we do not know the full impact of COVID-19 on people with disabilities.*

Organization of the COVID-19 & Disability Survey

The self-report survey includes demographic questions, including disability identifiers, gender, ethnicity, race, geographic area, and type of health insurance. A Summary Report combining all disability responses was issued by AAHD on May 8, 2020. Disability-specific reports have followed.

To ease respondent burden, we asked respondents for their *primary* disability, knowing that a substantial minority of respondents experience multiple disabilities. The four primary disability questions were drawn from the US Census Bureau and the Centers for Disease Control and Prevention's (CDC) disability definitions of difficulties with Hearing, Vision, Mobility, or Concentrating, Remembering, or Making Decisions (CRD). We added four follow-up questions for persons who identified as having CRD difficulties. This report focuses on persons who identified the underlying condition causing CRD as difficulty reading, writing, or communicating.

The survey includes a set of questions on access to regular (non-COVID-19) health care and services, including access to needed health care treatment, prescriptions, and the use of direct care workers.

The survey also includes questions regarding COVID-19 information and impact, including sources of information about the Coronavirus pandemic, adherence to public health COVID-19 recommendations, testing, and access to emotional supports during the pandemic.

B. Methodology

We reviewed existing COVID-19 surveys and either modified existing items or created new ones consistent with our areas of interest. Print, electronic and social media platforms were used to solicit responses from people with disabilities on AAHD's dissemination networks. AAHD also distributed the survey through existing partnerships and requested that other organizations and individuals distribute the survey. The survey was available online from April 17, 2020, until May 1, 2020.

C. Results

A total of 2,469 adults with disabilities responded to the survey, including 34 respondents who experienced serious difficulties concentrating, remembering, or making decisions due to difficulties reading, writing, or communicating. The survey completion rate varied by item, ranging from a low of 74 to 100%. The overall completion rate was 86% (exclusive of skip patterns). Specific results are described below. All results are rounded.

Demographics

Main Disability

As displayed in Table 1, 34 of respondents experienced serious difficulties concentrating, remembering, or making decisions due to difficulties reading, writing, or communicating or 1% of the total sample.

Table 1. Respondents with Difficulties Reading, Writing, or Communicating (N=34)

Disability	Number	Percentage
Serious Difficulty Reading, Writing, or Communicating	34	1%

Age

As displayed in Figure 1, respondents with difficulties reading, writing, or communicating selected from the following age groups: a) 18 to 44; b) 45-64; or c) 65 and above. The majority of the respondents were adults that were 18-44 (56%); followed by adults 45-64 (38%) and adults 65 and older (6%).

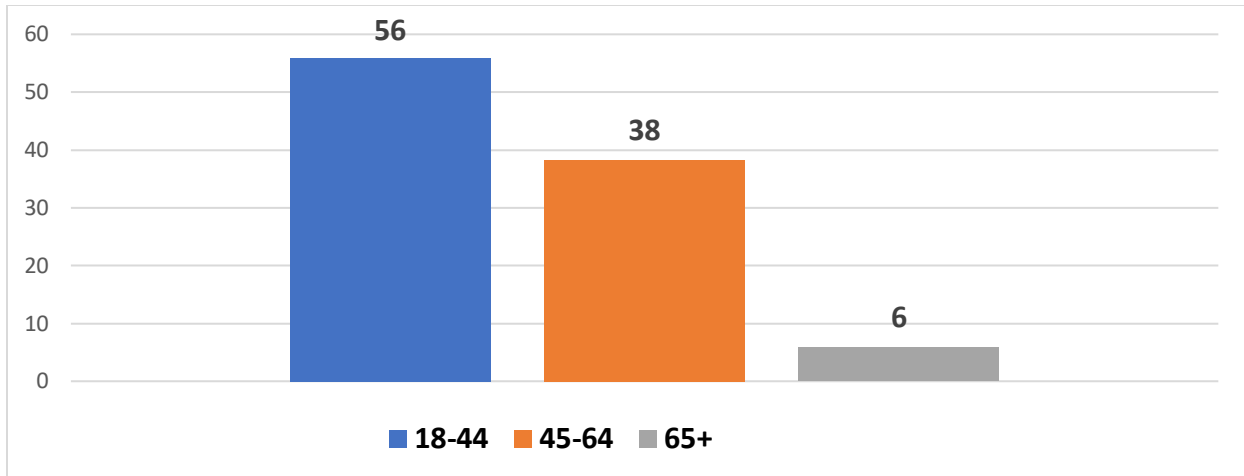


Figure 1. Percentage of Respondents by Age Group (N = 34)

Gender

As displayed in Table 2, the majority of adult respondents reported being either Female (72%) or Male (25%), with Gender Variant/Non-Conforming (0%), Other (0%), and Transgender Male (0%) and Transgender Female (0%). Three percent of the sample preferred not to answer.

Table 2. Gender of Respondents (N=32)

Gender	Percentage	Number
Female	72	23
Male	25	8
Transgender Female	0	0
Transgender Male	0	0
Gender Variant/Non-Conforming	0	0
Other	0	0
Prefer Not to Answer	3	1

Race and Ethnicity

We combined an ethnicity identifier (Hispanic, Latino, or of Spanish Origin) with race for the results, as shown in Figure 2. The sample was primarily White (80%), followed by Black or African-American (10%), Hispanic (3%), American Indian or Alaska Native (3%), Asian (0%), and Native Hawaiian or Other Pacific Islander (0%). Seven percent of respondents preferred not to answer.

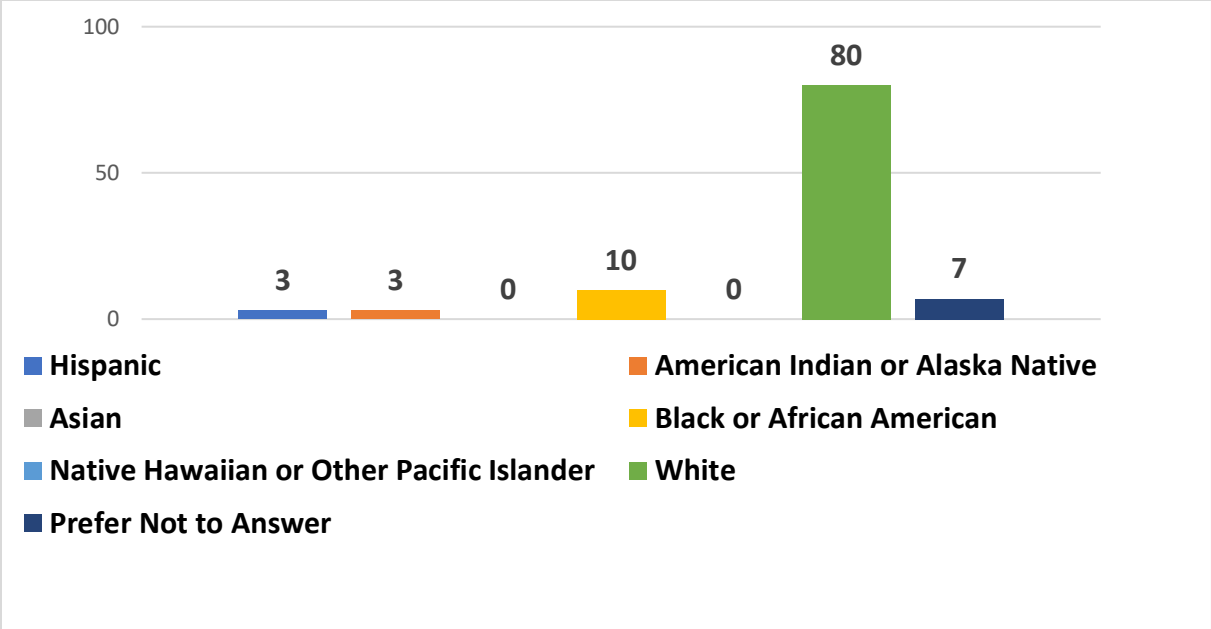


Figure 2. Percentage of Respondents by Ethnicity & Race (N = 30)

Geographic Location

Respondents selected their location as Urban, Suburban, Rural, or Frontier. As shown in Figure 3, the largest percentage of respondents were in Suburban locations (50%), followed by Rural (23%), Urban (23%), and Frontier (3%) locations.

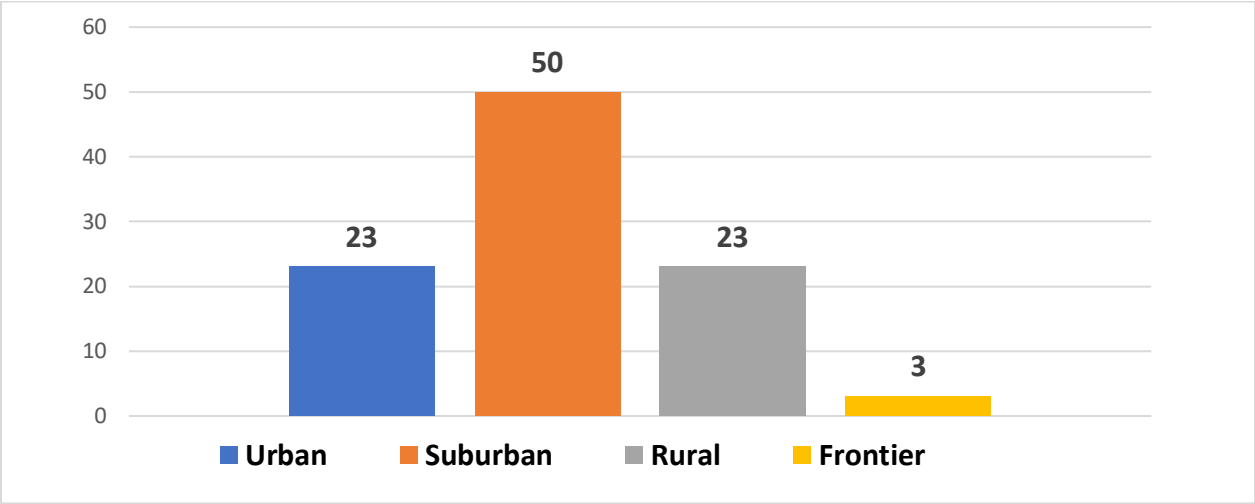


Figure 3. Percentage of Respondents by Geographic Location (N = 30)

Health Insurance

As displayed in Figure 4, the largest percentage of respondents with difficulties reading, writing, or communicating have Private Insurance (57%), followed by Medicare (33%), Medicaid/CHIP (27%), VA/Government insurance (0%) and Other (3%). Respondents were allowed to select “all that apply.” Three percent of respondents had no insurance.

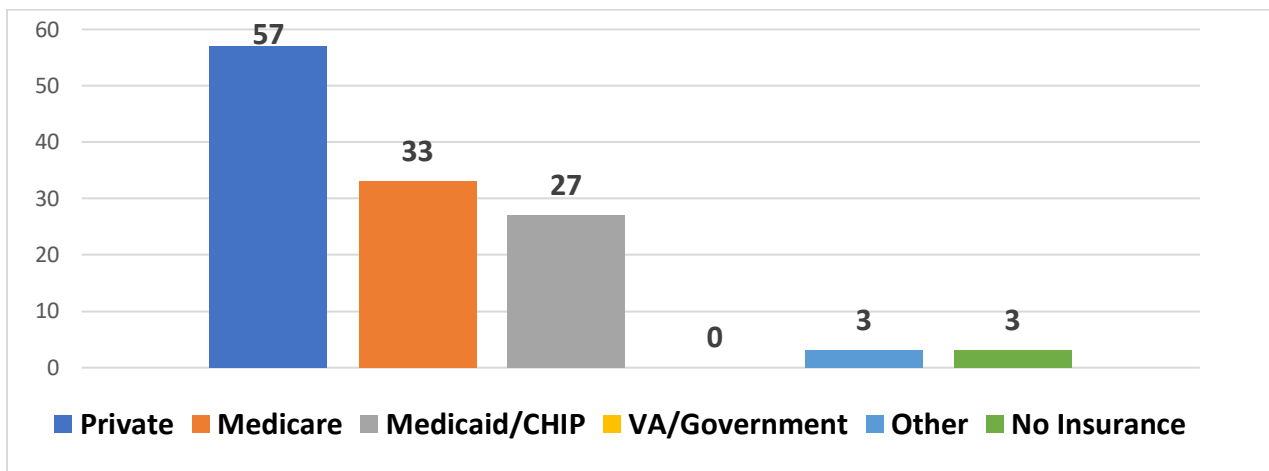


Figure 4. Percentage of Respondents by Type of Health Insurance (N = 30)

Access to Regular Health Care & Services

We developed a set of questions to determine if the coronavirus pandemic was impacting access to non-COVID-19 (regular) health care and services, including access to needed health care treatment, prescriptions, and the use of direct care workers.

Use of Direct Care Workers & Impact of COVID-19

Respondents were asked if they used a Direct Care Worker (e.g., Home Health Aide, Personal Care Aide, or Unpaid Family Caregiver), if their Direct Care Worker was continuing to provide care, and if respondents had been able to maintain a safe distance from their Direct Care Worker (e.g., six feet).

Forty-one percent of respondents reported having a Direct Care Worker in their home, while 59% did not (see Figure 5). As shown in Figure 6, among respondents with a Direct Care Worker, 64% of respondents continued to receive these home services, while 34% of respondents were no longer receiving these services.

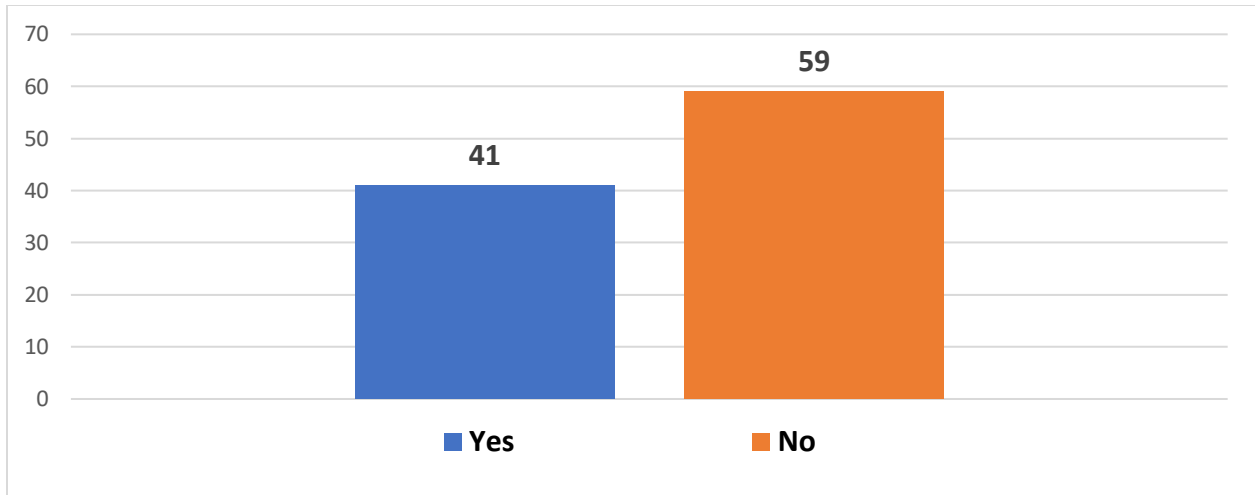


Figure 5. Percentage of Respondents Who Use a Direct Care Worker (N = 27)

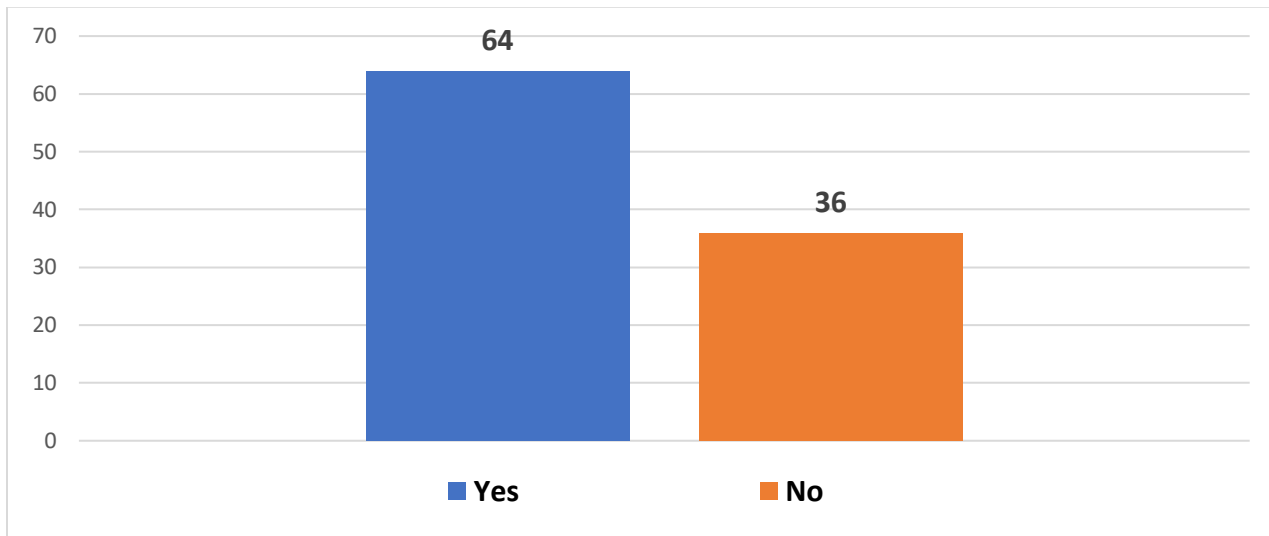


Figure 6. Percentage of Respondents with Direct Care Worker Home Visits During COVID-19 (N = 11)

Safe Distance Practices and Direct Care Workers

Respondents receiving Direct Care Worker services during the COVID-19 pandemic were asked if they were able to maintain a safe distance from their provider. As displayed in Figure 7, 29% of respondents reported that they were unable to maintain a safe distance from their Direct Care Worker, while 71% of respondents were able to maintain a safe distance.

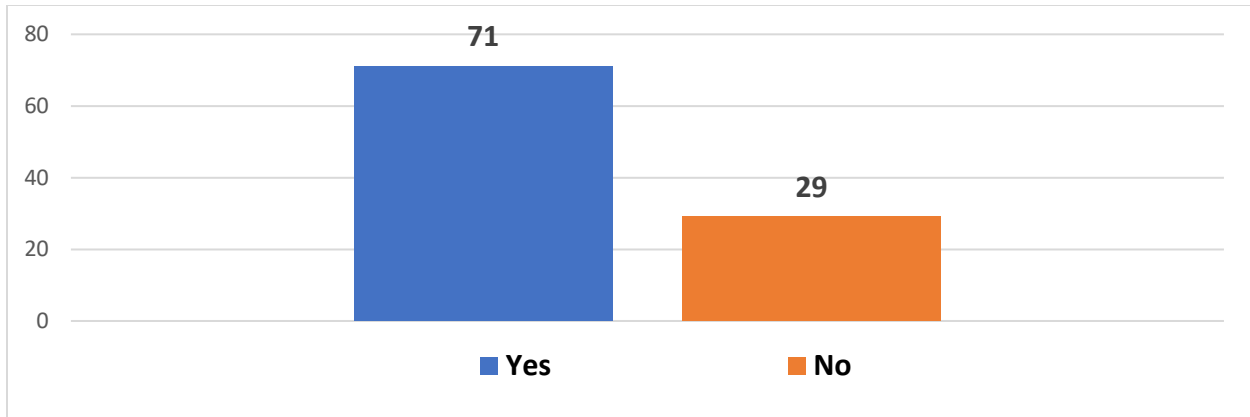


Figure 7. Percentage of Respondents Maintaining A Safe Distance from Direct Care Worker (N = 7)

Access to Regular Health Care Treatment

Respondents were asked if they were able to get regular health care treatment (such as physical therapy, dialysis, bloodwork, etc.) during the COVID-19 outbreak. Among respondents who needed regular health care, 50% reported access had been disrupted and 50% reported that access had not been affected (see Figure 8).

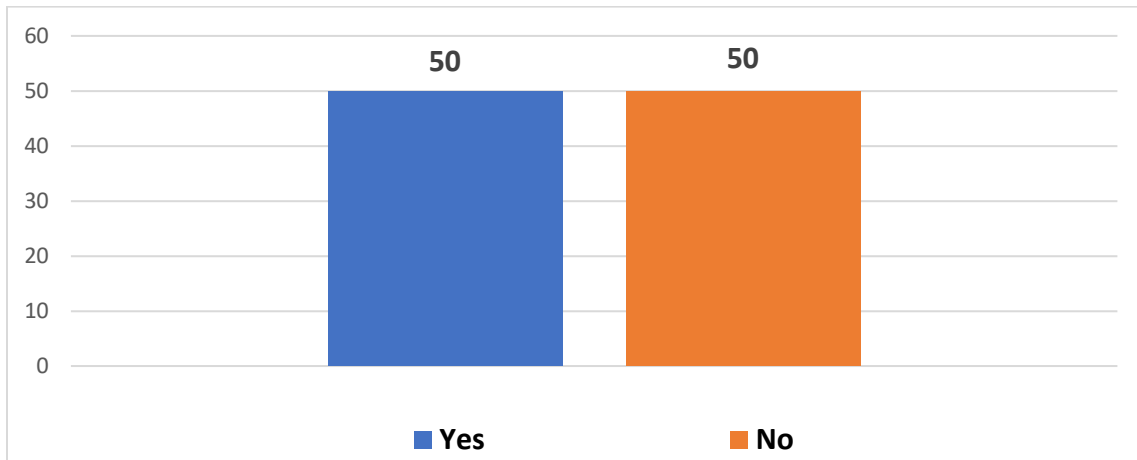


Figure 8. Percentage of Respondents Able to Access Regular Health Care (N = 16)

Access to Prescription Medications

Respondents were asked if they were able to obtain prescription medications during the COVID-19 outbreak. Among respondents taking prescriptions, 95% reported that access had not been affected and 5% reported access had been disrupted (see Figure 9).

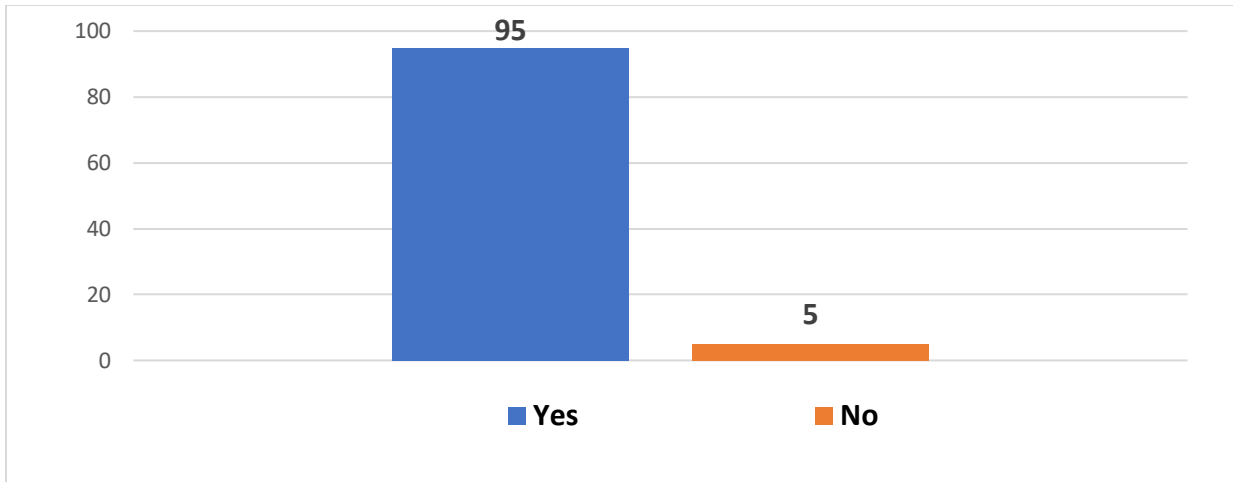


Figure 9. Percentage of Respondents Able to Access Prescriptions (N = 21)

New Challenges to Health

Respondents were asked if they were experiencing new challenges to obtaining health care treatment, health care access, and/or prescriptions during the COVID-19 outbreak. Twenty-three percent of respondents reported experiencing new challenges, while 77% reported that they had not experienced new challenges (See Figure 10).

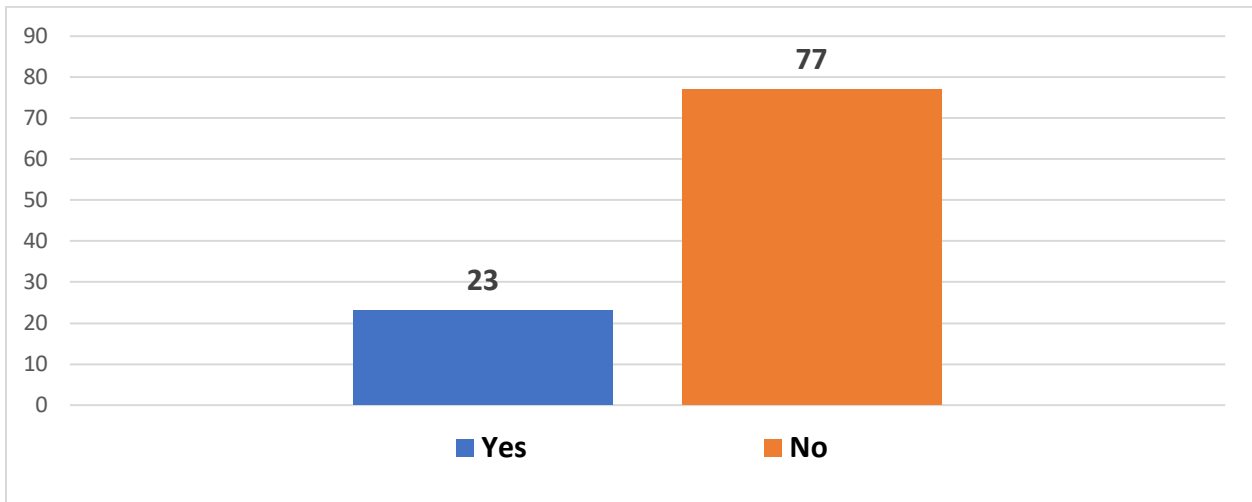


Figure 10. Percentage of Respondents Experiencing New Challenges to Accessing Health Care (N = 26)

COVID-19 Information & Impact

The survey includes a number of questions about the impact of COVID-19, including where respondents obtained information about the Coronavirus pandemic, adherence to COVID-19 public health recommendations, testing, and access to emotional supports during the pandemic.

COVID-19 Information Sources

Respondents were asked to rank their three most important sources of information about COVID-19 out of nine options. As displayed in Table 3, among the nine information sources, 33% of respondents identified the most important source as Television, followed by the Internet (30%), and HCP/HS (15%). Relatives and Other were at 7%, followed by Friends and Social Media (4%), and Radio and Print Media (0%).

Most respondents identified the Television (37%) and Internet (26%), followed by Health Care Providers, Friends, Other, and Print Media at (7%). Other second-most important sources of information were Relatives and Social Media at (4%), and Radio (0%).

Most respondents identified Social Media, Internet, Television and HCP at 15% and Friends and Other (11%) as the third-most important source of information, followed by Radio and Relatives at 7%. Other sources endorsed as the third-most important source of information, included Print Media (4%).

Table 3. Most Important Sources of COVID-19 Information (N = 27)

	HCP/HS	Internet	TV	Radio	Social Media	Print Media	Friends	Relatives	Other
Most Important	15%	30%	33%	0%	4%	0%	4%	7%	7%
Second Most Important	7%	26%	37%	0%	4%	7%	7%	4%	7%
Third Most Important	15%	15%	15%	7%	15%	4%	11%	7%	11%

Following COVID-19 Recommendations

Respondents reported high levels of compliance with COVID-19 public health recommendations. As shown in Figure 12, 96% of respondents reported following recommendations and 4% reported they were not following recommendations.

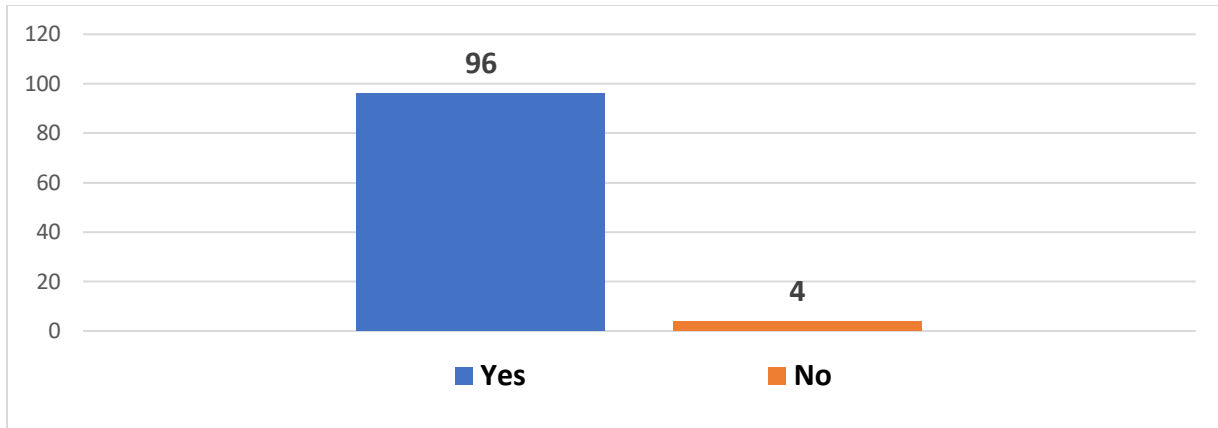


Figure 11. Percentage of Respondents Following COVID-19 Recommendations (N= 27)

COVID-19 Testing

We were interested in whether people with disabilities were being tested for COVID-19, if not, why not, and if insurance was covering testing. As shown in Figure 12, 7% of respondents had received a test for COVID-19 and 93% had not received testing. Among respondents who had been tested, 50% reported the test had been covered by insurance, 0% reported it had not, and 50% were not sure if the test was covered (see Figure 13).

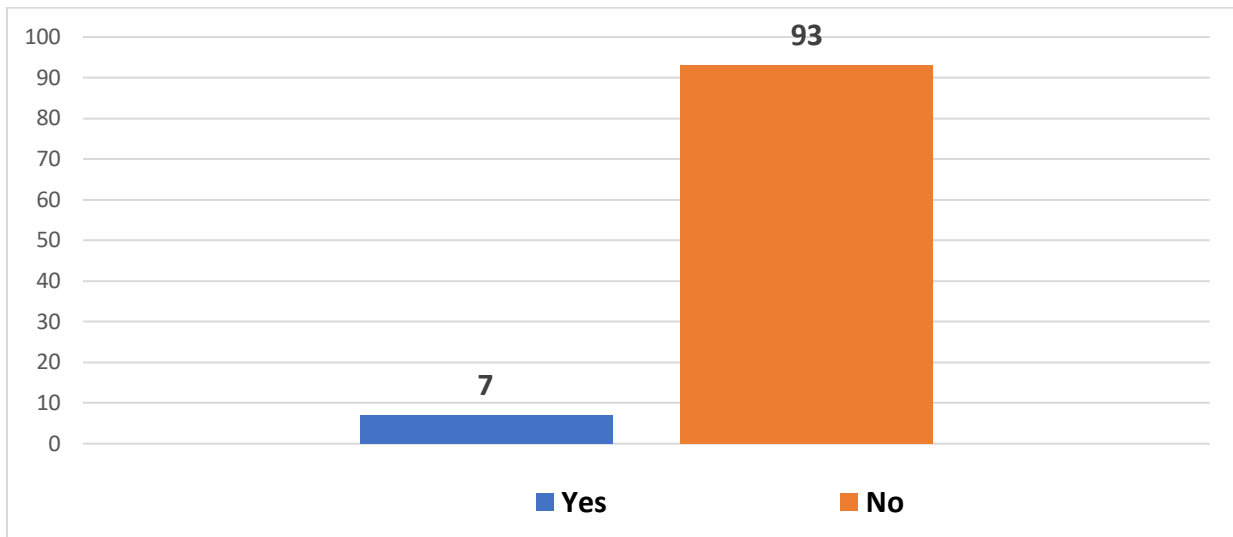


Figure 12. Percentage of Respondents Receiving COVID-19 Testing (N = 27)

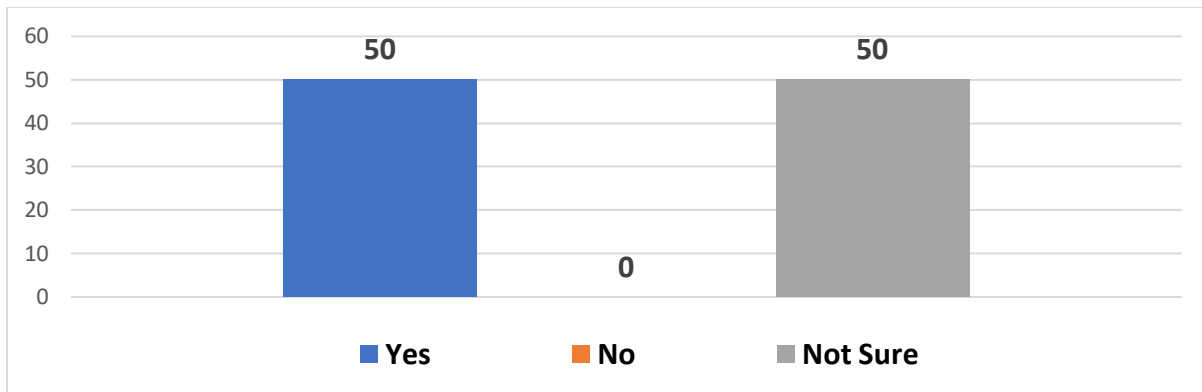


Figure 13. Percentage of COVID-19 Test Takers Covered by Insurance (N=2)

Why No COVID-19 Testing?

Respondents who had not received a COVID-19 test identified a range of reasons why they were not tested. As shown in Table 4, the largest percentage was among respondents who said they didn't need a test (88%). Among respondents who did need a test, reasons for not receiving the test included the test was unavailable (4%), followed by Other (4%), Health Care Provider would not approve the test (4%), information about testing was not accessible or available in alternate formats (4%), and lacking transportation to the test site, cost or concern that insurance wouldn't provide coverage, the testing site was physically inaccessible, the site lacked accessible parking, and the site did not have sign language interpreters, all at 0%. Respondents were able to identify multiple reasons.

Table 4. Reasons for Not Obtaining COVID-19 Test (N = 25)

Reason	Percentage
Didn't Need Test	88%
Test Was Unavailable	4%
Other	4%
Health Care Provider Would Not Approve Testing	4%
Information About Testing Not in Accessible Formats	4%
Lacked Transportation to Site	0%
Cost/Insurance Wouldn't Cover	0%
Site Was Physically Inaccessible	0%
Site Had No Accessible Parking	0%
Site Had No Sign Language Interpreters	0%

Anxiety & Depression

We asked respondents that were feeling anxiety or depression as a result of COVID-19 if they were able to access emotional support or services. As displayed in Figure 14, among respondents feeling anxiety or depression, 79% reported having access to emotional support or services and 21% reported not having access.

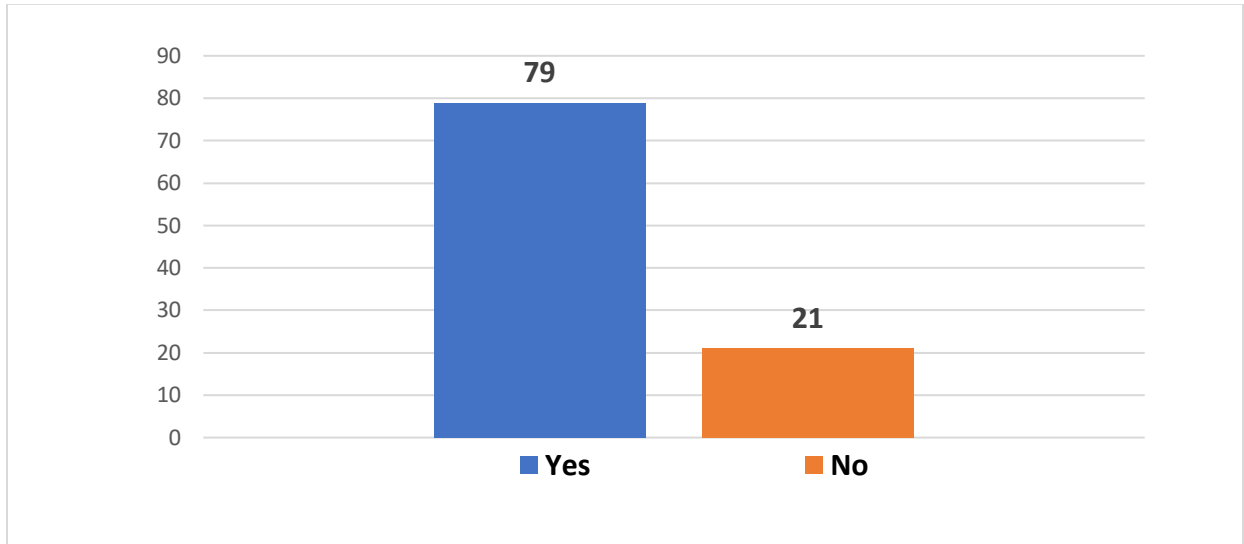


Figure 14. Percentage of Respondents with Access to Emotional Support/Services (N = 19)

D. Discussion

One of the limits of using a convenience sample is that results cannot be generalized to a larger population, although they can be informative (Bethell, et al, 2004). Below, we discuss observations about the survey results in relation to the sample's demographics and responses to the COVID-19 Information and Impact and Access to Health Care and Prescriptions sections.

Demographics

The overall COVID-19 and Disability Survey sample is atypical from population-based disability samples on a number of dimensions. For example, the highest percentage of primary disability in the total sample are persons reporting serious difficulty concentrating, remembering, or making decisions (64%), but only 34 respondents (1%) reported difficulty reading, writing, or communicating. Overall, the sample differs from many population-based results in age (56% of the sample are persons 18-44),

predominately female (72%), White (80%), and Suburban (50%), and have some type of insurance (only 3% of respondents reported not having insurance).

Access to Regular Health Care & Services

A high proportion of the sample used Direct Care Worker services (41%), an unexpected finding considering that 56% of the adult sample were 18-44 and 38% of the adult sample were 45-64. While a majority of respondents (64%) continued to receive these services during the pandemic, over one-third of the sample (36%) had lost critical care services. For the respondents maintaining their Direct Care Worker services, maintaining safe distances was impossible for nearly one-third of the sample (29%).

Of concern is that 50% of respondents who were scheduled to receive regular health care treatment services had those services disrupted during the coronavirus pandemic. In contrast, 95% of respondents taking medications reported that they could access their prescriptions. Yet, for the 5% of the sample who had prescription medication access disrupted, this represents a potentially life-threatening situation.

New Challenges to Health

In many ways, the novel coronavirus pandemic is presenting unfamiliar and unanticipated challenges for persons with and without disabilities. A key difference is that persons with disabilities already encounter documented disparities in accessing health care treatment and services and involvement in emergency planning programs and activities. Of considerable concern is that 23% of the sample reported experiencing new challenges to obtaining health care treatment, health care access, and/or prescriptions. A careful analysis of the open-ended responses as to why these new barriers exist is underway.

COVID-19 Information & Impact

Several unexpected findings emerged from the information and impact survey questions. Television is an important information source whether as the most important, second-most important or third-most important source, as well as the Internet. Health Care Providers and Health Systems played a significant information role for the sample, Social Media did not.

While several national polls report varying levels of endorsement for COVID-19 guidelines, the C-19 & D sample overwhelmingly reported high levels of compliance with these guidelines at 96%. This suggests that this sample are taking the risks of the coronavirus pandemic very seriously. Echoing national findings, only a small percentage

of respondents had received a COVID-19 test (7%) but there was significant uncertainty if health insurance would cover this cost. Interestingly, 88% of respondents did not need a test and only 4% of persons reported that they did not get a test because it was unavailable or a health provider would not authorize it. Importantly, general accessibility issues were not described as a problem.

Among persons that were experiencing anxiety or depression during the pandemic, 21% were not receiving emotional support or services.

Limitations

The findings in this report are subject to at least six limitations. First, the survey data are self-reported and may be subject to reporting or recall bias. Second, the responses represent a snap shot between April 17, 2020, until May 1, 2020, and the coronavirus pandemic situation may have changed since these responses were given. Third, persons without internet access were unable to access and complete the survey introducing a systematic bias in the sample. Fourth, the survey completion rate among adults ranged among items (74% - 100%), suggesting the presence of some response bias. Fifth, because of the use of a convenience sample and a very small sample size, the results are not generalizable to the overall population of persons with disabilities. Sixth, persons with multiple disabilities may not be accurately identified since we asked for a person's "main" disability.

E. Conclusion

The results of this survey provide insights into a moment in time during the coronavirus pandemic. The intricacies and changing dynamics of the pandemic establish a critical need for deeper and ongoing surveillance and policy and programmatic responses to this extraordinary health and public health challenge. The challenges and issues described in this report amply demonstrate a number of areas that need to be assessed at the population-level and longitudinally.

References

Bethell, C., Fiorillo, J., Lansky, D., Hendryx, M., & Knickman, J. (2004). Online consumer surveys as a methodology for assessing the quality of the United States health care system. Journal of Medical Internet Research, 6(1), e2. <https://doi.org/10.2196/jmir.6.1.e2>