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Patterns of negative seroconversion in ongoing surveys of SARS-CoV-2 antibodies among workers in New York's largest healthcare system

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ABSTRACT

Objectives Given the importance of continued COVID-19 surveillance, our objective was to present findings from a short follow-up survey of workforce SARS-CoV-2 antibody testing in previously seropositive participants and describe associations between work locations and negative seroconversion.

Methods We conducted a follow-up cross-sectional survey on previously seropositive healthcare workers, using questionnaires and serology testing. Eligible employees previously consented to be contacted were invited by email to participate in a survey and laboratory blood draws. SAS V.9.4 was used to describe employee characteristics and seroconversion status. Binomial regression models were used to calculate unadjusted and adjusted prevalence ratios (PRs) of seronegativity. The multivariable analyses included age, gender, race/ethnicity, region of residence, work location, prior diagnosis/PCR results and days between antibody tests. Unadjusted and adjusted PRs 95% CIs and p values were reported.

Results Of the 3990 employees emailed in the follow-up, 1631 completed an exposure survey and generated a blood-draw requisition form. Average time between serology testing was 4 months. Of the 955 employees with complete serology results, 79.1% were female, 53.4% were white and 46.4% resided in Long Island; 176 participants seroconverted to negative. In multivariable regression analyses adjusted for gender, race/ethnicity and region of residence, younger employees (<20–30 years), intensive care unit workers and those with no/negative prior PCR results were more likely to have negative seroconversion.

Conclusions and relevance Patterns of negative seroconversion showed significant differences by sociodemographic and workplace characteristics. These results contribute information to workplace serosurveillance.

INTRODUCTION

Prior to the availability of vaccines for SARS-CoV-2 that causes COVID-19, exposure assessments through surveillance included antibody testing that allowed workplaces to understand prevalence patterns and potential exposure.^{1,2} As with many healthcare systems across the country, the largest one in New York implemented various COVID-19 policies that augmented infection control to increase the number of heating ventilation and air-conditioning

Key messages

What is already known about this subject?

- Antibodies for COVID-19 decline over time. However, population patterns of negative seroconversion are not well understood.

What are the new findings?

- The study discovered sociodemographic and workplace patterns of negative seroconversion among healthcare workers. In a follow-up survey of 955 previously seropositive New York healthcare workers, 176 retested negative over a 4-month study period. High negative seroconversion occurred among those with who had a negative or no prior PCR test, intensive care unit (ICU) workers (compared with non-ICU workers) and younger employees (20–29 years compared with other age groups).

How might this impact on policy or clinical practice in the foreseeable future?

- This is a large follow-up of seropositive healthcare workers, paired with a survey of work characteristics and self-reported exposures. Future directions include a review of vaccine uptake and seroconversion with molecular differentiation that includes an understanding of sociodemographic and workplace patterning of exposure over time.

inspections on floors and units, as well as adjusting specifications for aerosolising procedures and personal protective equipment (PPE) during patient handling.^{1–6} In spring 2020 (wave 1), our healthcare system offered free SARS-CoV-2 antibody testing to all 70812 employees.² Results of the testing have been previously reported for dates between 20 April 2020 and 23 June 2020. In brief, of the final consented participants (40 329), 13.7% (5523) tested positive for IgG antibodies.² The findings were comparable to New York State antibody screenings at the time, which revealed 12.3% of the general population were seropositive.^{7–9} The New York State Governor's report at the time further showed 10.5% of the New York City Police Department and 17.1% of New York City Fire Department Emergency Medical Service (FDNY/EMS) members had tested positive for antibodies.^{2,9}

Our healthcare system's wave 1 results showed that 93.7% of employees who were diagnosed positive through reverse transcription PCR tests were also positive for antibodies.² Further, a self-reported index of suspicion of exposure to COVID-19 correlated well with antibody measures. This suggested that workers without confirmed PCR tests were attuned to degrees of exposure (online supplemental figure S1) and that the suspicion index was a robust measure of exposure perception.² In the fall of 2020 (September and October) (wave 2), the healthcare system extended serology testing to 3990 employees, who were antibody positive in wave 1 and who agreed to future testing. This wave 2 serology testing introduced an opportunity to understand prevalence patterns in the workplace and target protective measures, as needed. Here, we present initial findings from a short exposure survey and the second round (wave 2, September and October 2020) of workforce testing for SARS-CoV-2 antibodies in participants previously identified as seropositive (wave 1, April and May 2020). Antibodies for COVID-19 decline over time. However, population patterns of negative seroconversion are not well understood. Given the importance of continued COVID-19 surveillance, our objective was to present findings from a short follow-up survey of workforce SARS-CoV-2 antibody testing in previously seropositive participants and describe associations between work locations and negative seroconversion.

METHODS

On 8 September 2020 (with reminders on 23 September 2020), 3990 emails were sent to eligible employees, who had initially tested positive for SARS-CoV-2 antibodies between April and June 2020 (wave 1) and who had consented to future serology testing. Some participants had received prior PCR testing.² For this wave 2 study, blood draw requisitions and lab results were collected from 8 September 2020 to 5 October 2020. Antibody testing was validated at our healthcare system laboratories using a combination of ELISA, chemiluminescent microparticle immunoassays and immunometric assays.^{2 10} These are instrument-based, qualitative IgG SARS-CoV-2 immunoassays with up to 97% sensitivity.^{2 10} The antibody test is not a test for active infection but may identify previous exposure to the virus within at least the past 1–2 weeks.^{2 10} Additional information regarding the testing panels is available as online supplemental table S1.

Data collected from wave 2 testing were gathered, prepared and analysed for continued presence of SARS-CoV-2 antibodies. We used SAS V.9.4 software (SAS Institute Inc.) to conduct binomial regression analyses to calculate unadjusted and adjusted prevalence ratios (PRs) of seronegativity and frequency distributions for sociodemographic characteristics, work location, time between testing, county of residence, prior diagnosis/PCR, recent travel and level of suspicion of re-exposure to virus: 'Do you believe you may have been exposed again to COVID-19?' (1–9; 1=no; 9=yes definitely; 7–9=high suspicion of exposure). 'If so, did you feel ill?' (Yes/No). 'Do you usually take public transportation to work?' (Yes/No). 'In the past month, have you traveled outside of the NY metropolitan region and northeast?' (Yes/No). Several industrial hygiene/infection control practices were noted, to include PPE policies (universal masking was required with N95 minimum for staff and procedural mask minimum for patients from as early as 7 March 2020; visitors were restricted and also required masking with temperature checks), ventilation requirements that adhered to American Society of Heating, Refrigerating and Air-Conditioning Engineers recommendations with minimum air changes per hour (ACH) equal to 12–15 for

intensive care units (ICUs) that required minimum efficiency reporting values (13–14) to ventilate the rooms and positive pressure breathing apparatus (level B PPE). Patient waiting areas had a minimum recommendation of 4–6 ACH.^{5 6}

RESULTS

Of the 3990 wave 2 email invitations, 1631 employees responded to the survey and obtained requisitions, rendering a response rate of 40.9%; of those, 955 (58.6%) received wave 2 serology results and 779 (81.6%) retested positive (table 1). Average months between testing were 4.3 (SD=0.5), median=4.0 and IQR=1. Average age of non-participants in wave 2 was similar to that of participants. Blacks, Hispanics and males were less likely to participate in wave 2 (online supplemental table S2).

Overall negative seroconversion proportion was 176/955 (18.4% (95% CI=16.0% to 21.0%)). In regression models that included age, gender, race/ethnicity, residential county, days between testing and prior PCR test, those working in areas other than the ICU were less likely to seroconvert to negative (PR=0.58, 95% CI=0.36 to 0.93, p=0.02), compared with ICU workers. Those who were previously seropositive and with a negative PCR result (PR=2.22, 95% CI=1.38 to 3.59, p=0.001) or no PCR test (PR=2.56, 95% CI=1.83 to 3.58, p<0.0001) were 2½ times more likely to seroconvert to negative than those with a positive PCR test (table 1). Residential and work counties, recent travel, transportation type and suspicion of re-exposure were not significantly associated with negative seroconversion.

DISCUSSION

Our results present patterns of negative seroconversion in a large, single study follow-up of previously seropositive healthcare workers. As expected, SARS-CoV-2 antibodies decreased over time.^{4 10–12} Almost one-fifth of previously seropositive employees experienced negative seroconversion over a 4-month period.

Those working in ICUs were more likely to seroconvert (to negative) than employees in other work areas. Though not possible to confirm with this study, it is plausible that ICU locations have decreased potential for reinfection (exposure) because of effective infection control practices at work (increased ventilation checks and adherence to optimal standards, appropriate administrative policies and targeted PPE), low infection among coworkers, coupled with low prevalence of active community infection (at the time of wave 2 data collection, New York State had new daily infection rates below 1%).⁹ It is unclear why those over the age of 30 years were less likely to seroconvert to negative and thus, seemed to retain measurable SARS-CoV-2 antibodies for longer than younger workers. Similarly, those with confirmed disease through past PCR positive results were less likely to seroconvert to negative. The PCR testing for this cohort was conducted early in the pandemic (March/April 2020) when most tests were reserved for sicker, hospitalised individuals. It is conceivable that asymptomatic and those with milder symptoms were less likely to have received a PCR test.

There are limitations to these findings. This study sample was voluntary in waves 1 and 2 and results may not be generalisable to the population of all our seropositive healthcare workers. With wave 2 testing, there were patterns of non-response that may have influenced results. Determining the exact timeline between infection, antibody generation and antibody reduction was not possible because some employees were not diagnostically tested (PCR) at the time of infection. It is also possible that those who were more likely to seroconvert (to negative) during

Table 1 Characteristics of those with follow-up IgG antibody testing with regression results for seroconversion to negative

Characteristic	Survey with serology results (N=955)						Multivariable** PR (95% CI), p value
	Seropositive (N=779)			Seronegative (N=176)			
	Category	n	%	n	%	Bivariate** PR (95% CI), p value	
Age (years)	20-29	101	73.7	36	26.3	1.00 (REFERENCE)	1.00 (REFERENCE)
	30-39	175	82.6	37	17.5	0.66 (0.43 to 0.99), p=0.045	0.69 (0.45 to 1.04), p=0.08
	40-49	159	82.4	34	17.6	0.68 (0.45 to 1.04), p=0.07	0.74 (0.49 to 1.11), p=0.14
	50-59	209	82.3	45	17.7	0.65 (0.44 to 0.97), p=0.03	0.77 (0.52 to 1.14), p=0.18
	60-69	124	85.0	22	15.1	0.57 (0.36 to 0.92), p=0.02*	0.60 (0.37 to 0.97), p=0.04*
	70+	11	84.6	2	15.4		
Gender†	Female	620	82.1	135	17.9	1.00 (REFERENCE)	1.00 (REFERENCE)
	Male	154	79.4	40	20.6	1.15 (0.84 to 1.59), p=0.39	1.18 (0.85 to 1.63), p=0.32
Race/ethnicity†	White	413	81.0	97	19.0	1.00 (REFERENCE)	1.00 (REFERENCE)
	Black	91	81.3	21	18.8	0.98 (0.64 to 1.50), p=0.92	1.06 (0.69 to 1.63), p=0.78
	Hispanic	125	83.9	24	16.1	0.84 (0.56 to 1.27), p=0.40	0.86 (0.57 to 1.29), p=0.47
	Asian/Pacific Islander	123	80.4	30	19.6	0.95 (0.65 to 1.39), p=0.80*	0.97 (0.65 to 1.44), p=0.87*
	American Indian	3	100.0	0	0		
	Two or more races	5	83.3	1	16.7		
	Non-specified/missing	19	86.4	3	13.6		
	Full-time day	521	82.8	108	17.2	1.00 (REFERENCE)*	
	Full-time other	110	76.4	34	23.6		
	Part-time day	95	76.6	29	23.4	1.06 (0.74 to 1.51), p=0.75*	
Part-time other	35	94.6	2	5.4			
Work location	Missing	18	85.7	3	14.3		
	ICU	35	71.4	14	28.6	1.00 (REFERENCE)	1.00 (REFERENCE)
Work county#	ED	72	82.76	15	17.2	0.62 (0.33 to 1.17), p=0.14	0.61 (0.33 to 1.13), p=0.12
	Hospital units (non-ICU)	209	80.4	51	19.6	0.69 (0.41 to 1.15), p=0.15	0.7 (0.43 to 1.14), p=0.15
	Other	456	82.9	94	17.1	0.60 (0.37 to 0.97), p=0.04	0.58 (0.36 to 0.93), p=0.02
	Missing	7	77.8	2	22.2		
	Nassau	176	80.0	44	20.0	1.00 (REFERENCE)	
	Manhattan	84	78.5	23	21.5	1.09 (0.69 to 1.70), p=0.72	
Residential county#	Queens	93	82.3	20	17.7	0.88 (0.55 to 1.43), p=0.62	
	Richmond	88	82.2	19	17.8	0.86 (0.52 to 1.41), p=0.54	
	Suffolk	85	83.3	17	16.7	0.85 (0.50 to 1.46), p=0.56	
	Westchester	26	74.3	9	25.7	1.31 (0.70 to 2.44), p=0.39	
	Other/unknown	227	83.8	44	16.2	0.78 (0.52 to 1.14), p=0.21	
Residential county#	Nassau	219	79.6	56	20.4	1.00 (REFERENCE)	
	Bronx	21	80.8	5	19.2	0.98 (0.43 to 2.22), p=0.95	
	Kings	36	81.8	8	18.2	0.92 (0.47 to 1.81), p=0.81	
	New York	26	74.3	9	25.7	1.23 (0.64 to 2.36), p=0.53	
	Queens	141	82.9	29	17.1	0.87 (0.58 to 1.31), p=0.51	

continued

Table 1 continued

Characteristic	Survey with serology results (N=955)					
	Seropositive (N=779)			Seronegative (N=176)		
	Category	n	%	n	%	
	Richmond	86	86.9	13	13.1	
	Suffolk	141	83.9	27	16.1	
	Westchester	50	79.4	13	20.6	
	Other/unknown	59	78.7	16	21.3	
	New York City	310	82.9	64	17.1	
	Long Island	360	81.3	83	18.7	
	Other	109	79.0	29	21.0	
	Positive	391	89.9	44	10.1	
	Negative	61	75.3	20	24.7	
	No PCR test	327	74.7	111	25.3	
	Equivocal	0	0	1	100	
	Low	463	83.1	94	16.9	
	Medium	203	79.0	54	21.0	
	High	113	80.1	28	19.9	
	No	728	81.3	167.0	18.7	
	Yes	51	85.0	9.0	15.0	
	No	718	81.4	164	18.6	
	Yes	60	83.3	12	16.7	
	Missing	1	100	0	0	
	No	681	81.6	154	18.4	
	Yes	94	81.7	21	18.3	
	Missing	4	80.0	1	20.0	
	Days between antibody testing	Mean=140 SD=10 Minimum=105 Maximum=174		Mean=139 SD=12 Minimum=62 Maximum=158		

*In regression models, the following categories were combined: Age=60+ years of age; race/ethnicity (Asian/Pacific Islander, American Indian, two or more races); shift=full time versus part-time.

†Administrative data collected from self-reports in prespecified fixed categories.

‡New York City=Bronx, Kings, New York, Queens, and Richmond; Long Island=Nassau and Suffolk; other=Westchester and other/unknown.

§Do you believe you may have been exposed again to COVID-19? (1=3=low; 4=6=medium; 7=9=high suspicion of re-exposure).

¶If so, did you feel ill? (Yes/No).

**In the past month, have you traveled outside of the NY metropolitan region and northeast? (Yes/No).

††Do you usually take public transportation to work? (Yes/No).

‡‡Clopper-Pearson 95% CIs. SAS GLIMMIX procedure modelling the probability of being seronegative (includes key sociodemographic and work characteristic variables), using a Newton-Raphson optimisation and type III test for fixed effects, missing values were excluded from the analyses; bolded prevalence ratios (PRs) indicate statistically significant associations at p<0.05. A PR greater than 1 is compared with the reference group; similarly, a PR less than 1.00 indicates that the group was less likely to seroconvert to negative.

ED, emergency department; ICU, intensive care unit; n, subsample count; N, sample count.

this follow-up were infected earlier than those who retained antibody seropositivity.

Nevertheless, these results are informative and document patterns of negative seroconversion in a sample of healthcare workers. This study also introduces exposure questions and biomarkers that will be expanded in future surveys. Sophisticated biological/molecular testing is particularly important as we begin to integrate vaccination results and patterns of infection.

Globally, frontline workers were among the hardest hit group early in the SARS-CoV-2 pandemic.^{3,4} In one of the largest cross-continental cohort studies that encompassed the UK and USA risk of reporting COVID-19 exposure or infection was highest among frontline healthcare workers.³ Data from New York showed variability by institute and county that may indicate community spread as the driver of these rates.² In all studies, minority and socially disadvantaged groups were most affected and had among the highest prevalence of SARS-CoV-2 antibodies.^{1,2} Our results provide a summary of follow-up serology testing in New York State's largest workforce. Future directions include ongoing monitoring and analyses of the effects of vaccine delivery with a concerted outreach plan for healthcare workers and the surrounding communities.

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