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# COVID-19 serologic testing: FAQs and caveats

**T**HERE HAS BEEN an immense amount of tangential discussion regarding the potential usefulness of serologic testing for COVID-19. Serologic testing has never been routinely used for diagnosing infections with “respiratory viruses” such as influenzae, parainfluenzae, respiratory syncytial viruses, adenoviruses, or metapneumovirus, nor was it used routinely for diagnosis during the global epidemics of severe acute respiratory syndrome (SARS), Middle East respiratory syndrome (MERS), and H1N1 influenza. However, the pandemic status of COVID-19 and the shortage of nucleic acid detection kits and swabs in certain areas raise the prospect of resorting to serology as an alternative to direct testing for the virus, and it is relevant to ask how useful it may be. The Infectious Diseases Society of America has recently issued a clear statement on COVID-19 serology.<sup>1</sup>

The following addresses some common questions regarding serologic testing for COVID-19.

## **Is IgM/IgA serology reliable for diagnosing acute symptomatic COVID-19?**

Based on recent publications,<sup>2</sup> the appearance of detectable immunoglobulin M (IgM) antibodies after infection with COVID-19 is delayed, resulting in abysmal sensitivity ranging from 17% to 50% in the first 10 to 14 days after the “onset” of symptoms. Note that this is not days after exposure or infection, but rather days after the onset of clinical symptoms. Unfortunately, the results may not be clinically useful because COVID-19 often progresses very quickly within the first 7 to 10 days.<sup>3</sup> Thus, by the time of seroconversion, patients could be critically ill with septic shock

or multiorgan failure, or they could die before seroconverting.

Most hospitalized patients typically receive the diagnosis of COVID-19 by nucleic acid testing before admission or up to 24 hours after admission. Unfortunately, by the time of serologic diagnosis, the patient may have inadvertently infected innumerable contacts.

There are no carefully peer-reviewed studies regarding the specificity of IgM and IgA tests, even though numerous point-of-care and non-point-of-care enzyme immunoassays (EIAs) are commercially available. IgM serologic tests, in general, have an inherent predisposition to false-positive results. Viruses as distantly related as Dengue virus have been reported to cause false-positive IgM results in COVID-19 point-of-care serologic tests.<sup>4</sup>

COVID-19 IgA EIAs had false-positive results in 20% of samples from 2,018 patients in the United States (author’s personal communications). The potential for a rapidly progressing clinical course of COVID-19, combined with the low sensitivity of IgM testing during the first 10 days of clinical infection, makes this low specificity of IgA testing a concern, since class-switching to IgA typically occurs after the appearance of IgG.

## **Is IgG serology a reliable option for diagnosing acute or convalescent COVID-19?**

IgG seroconversion is delayed after the onset of symptoms (more than 35 days in some cases), but typically occurs in 2 to 3 weeks, at which time it can be detected if the test specificity is high. Commercially available serologic assays, typically enzyme-linked immunosorbent assays (ELISAs), require validation with a plaque-reduction neutralization test (PRNT).

In brief, PRNT requires mixing live viruses with serially diluted serum followed by

**The pandemic status of COVID-19 and the shortage of supplies raise the possibility of serologic testing as an alternative to direct testing**

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**By the time of seroconversion, the patient could be critically ill or dead**

cell cultures to view cytopathic effect. PRNT is a functional assay that requires significant expertise and a biosafety level 3 facility (not available in hospitals), and it is not amenable to automation; however, it is necessary when any new assay is being validated. Ideally, this test should be done by manufacturers prior to US Food and Drug Administration submission; if this is a lab-developed test, the onus is on the lab to ensure PRNT is done on-site or in collaboration with a reference lab that has PRNT capability. Additionally, PRNT needs to be done head-to-head against other known coronaviruses, particularly those that are commonly acquired in the community (eg, 229E, OC43, NL63, HKU1), which have always been detected using nucleic acid amplification tests. Thus far, none of the published studies or commercially available kits have documentation of such validation.

That said, PRNT has its limitations. Previous exposure to common coronaviruses may lead to an early and high-titer humoral immune response to SARS-CoV-2. As time elapses, however, the humoral response probably becomes more specific to SARS-CoV-2. Studies have shown greater than 90% seroprevalence of common coronaviruses in the United States. Interestingly, Wölfel et al<sup>2</sup> report finding a significant degree of serologic cross-reactivity between SARS-CoV-2 and common coronaviruses. Further, IgG responses were much stronger and appeared earlier than IgM responses. It seems that exposure to SARS-CoV-2 triggers previous memory response to all common coronaviruses. Based on the current information, it is not clear which target provides the best specificity, but specificity should increase over time as the immune response becomes more fine-tuned. This, however, will be well beyond the recovery time and, thus, of no use for diagnostic purposes.

In addition to cross-reactivity with common coronaviruses, false-positive results are seen using serum with elevated antinuclear antibody titers. Elevated titers are relatively common in patients over age 50, which happens to overlap with the median age for COVID-19 diagnosis. False-positive results have also been documented with serum from patients with influenza or influenza vaccine re-

cipients. Flu vaccine recipients constitute a large population—45% of the adult US population, according to the US Centers for Disease Control and Prevention (CDC)—who may have overlapping signs and symptoms of influenza and COVID-19.

On the IgG side, false-positives using both EIA and point-of-care testing kits also have been observed in serum samples from patients with herpes simplex virus type 1, human metapneumovirus, enterovirus, parvovirus B19, and sera-positive rheumatoid factor, among others. Finally, even if IgG is to be used with a highly specific ELISA for diagnosing acute COVID-19 infection, one still has to wait several weeks to see a minimum 4-fold rise in antibody levels. This would be too late to be of clinical use. And testing requires a minimum of 2 blood draws (acute and convalescent), exposing sick patients to even more healthcare environments.

### **Is IgG serology reliable for evaluating infectivity and clinical immunity to reinfection with COVID-19?**

No one knows. Patients with a positive IgG result may still be sick and can shed the virus through their respiratory secretions or stools. SARS-CoV-2 is an enveloped RNA virus belonging to the Coronaviridae family, which includes common coronaviruses such as 229E, OC43, NL63, HKU1, and several that infect animals. Upper respiratory samples can remain positive for viral RNA for a few weeks after onset, when patients are supposed to have IgG antibodies. Viral shedding in stool has been reported for up to 47 days, which speaks against authentic neutralizing capacity of tissue-transudated IgG and secretory IgA antibodies.<sup>5</sup> SARS-CoV (a SARS-CoV-2 sister virus) has been grown in cultures from upper respiratory samples in 54% of cases at 2 weeks after symptom onset and in 16% of cases at 3 weeks after symptom onset, despite documented seroconversion in more than 92% of patients assessed by PRNT that detected “neutralizing antibodies.”<sup>6</sup> Thus, having circulating neutralizing antibodies may not ensure lack of infectivity. This has yet to be shown in SARS-CoV-2.

As of this writing, the CDC has not established guidelines for occupational health isola-

tion disposition based on serologic testing, other than using 2 consecutive negative nucleic-acid amplification tests at least 24 hours apart.<sup>7</sup>

Regarding COVID-19, the correlate of protection is not known, although these levels have been established for many other viral diseases. For example, the correlate of protection for hepatitis B is a surface antibody level at or very close to 10 mIU/mL, and this measure is routinely used for occupational health purposes. For COVID-19, the correlate of protection has to be established in large, well-designed randomized controlled trials, which have not been conducted. Therefore, determination of “immune status” of individuals, including healthcare workers, to SARS-CoV-2 cannot be established at this time using serology. To further confound matters, all individuals can be infected and become sick with common coronaviruses in the community in almost every season and sometimes several times during a season. This suggests that immunity to some coronaviruses is short-lived, and lingering IgG antibodies from previous seasons does *not* mean an individual is necessarily immune to infection with the same coronaviruses. Furthermore, cell-mediated immunity (typically mediated through CD8+ memory T cells) also plays a role.

### Is IgG serology reliable for screening a COVID-19-convalescent donor?

The discussions in the previous 2 items provide a segue to answer this question. First, we do not know if EIA results correlate well with PRNT (ie, ELISA antibodies vs neutralizing antibodies). And if they do correlate well, then second, we do not know if the so-called neutralizing antibodies are neutralizing enough to confer immunity.

Shen et al<sup>8</sup> gave critically ill patients infusions of 400 mL of convalescent plasmas collected from donors with clinically resolved COVID-19. Interestingly, the critically ill recipients' pretransfusion neutralization titers were approximately only 1 dilution different than those of the donors (pretransfusion neutralizing antibody geometric titers of 192 and 80, in donors and patients, respectively). Further, Duan et al<sup>9</sup> found that severely ill patients had neutralization titers as high as 1:640 before receiving transfusions of convalescent

plasma. Healthy and COVID-19-resolved donors had titers higher than 640.

These results raise the question as to why patients who already had mounted neutralizing antibody titers were still critically ill. This could be explained by the phenomenon called *antibody-dependent enhancement*, in which viruses can gain access to Fc gamma receptor-expressing cells via antibody-recognizing receptors as opposed to viral receptors and proliferate or trigger those cells to respond with a vigorous and potentially harmful cytokine release (cytokine storm). More recently, Wölfel et al<sup>2</sup> grew SARS-CoV-2 in upper and lower respiratory samples from onset until day 8 but not beyond that. This suggests that transfusion of convalescence-phase plasma may not have a role beyond day 8 after onset. This is important, as passive immunotherapy is typically considered in critically ill patients who are typically well beyond this time point.

Antibody-dependent enhancement has been shown in coronaviruses, which may potentially lead to more severe subsequent coronaviral diseases. Although this may have implications for vaccine design (similar to those of Dengue vaccine), it may also lead to potential adverse outcomes for convalescent plasma therapy. At this juncture, we do not have any evidence that plasma from patients who have recovered offers clear clinical benefit, as it showed mixed results for SARS or MERS.<sup>10</sup> Further, SARS-CoV and SARS-CoV-2 can cause syncytium formation among lung epithelial cells, thereby paving the way for cell-to-cell transmission of the virions. In this way, virions may be protected from antibody neutralization.

Using a serologic test with poor or unknown performance characteristics to “green-light” distributing blood products (plasma) is not really an undertaking for hospital labs. The US Food and Drug Administration (FDA),<sup>11</sup> however, recommends neutralizing antibody titers of at least 1:160, but a titer of 1:80 may be considered acceptable if an alternative matched unit is not available. The FDA also recommends that convalescent plasma be considered only for patients with severe or immediately life-threatening COVID-19. The

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**At this point, using serology to screen COVID-19 convalescent donors is fraught with risk**

FDA further clarifies that, although promising, convalescent plasma has not yet been shown to be effective in every disease studied. It is therefore important to determine through clinical trials, before routinely administering convalescent plasma to patients with COVID-19, that it is safe and effective to do so.<sup>11</sup>

In short, at this point, using serology to screen COVID-19-convalescent donors is fraught with risk, not only because there is no robust science to back it up, but also because there are no FDA-approved products for donor screening.

## Is IgG serology reliable for SARS-CoV-2 serosurveys?

Maybe, but it very much depends on the specificity of the assay. Serology may only be good for surveillance or seroepidemiologic studies, which is a public health function or an academic project. Once fine-tuned assays are available and resources allow, impact assessments will need to be done on a large scale in collaborative studies performed using well-balanced and unbiased samples that include multiple age, sex, and geographic cohorts.

Another aspect here is to assess what percentage of infected individuals remain asymptomatic and to calculate the case-fatality rate (CFR). The former is helpful as background epidemiologic knowledge, but the CFR is even more important, although it comes at a potential cost. The cost is that taking asymptomatic cases into account when the CFR is calculated amid an outbreak, a sudden drop in the publicly announced CFR would potentially lead to loosening precautionary measures by the general public, which may lead to further spread of the infection.

Here's some math to consider. As of April 16, 2020, there were 657,720 cases in the United States. If we are at the peak of the outbreak, then by the end of the outbreak, we will have an estimated 1,315,440 cases (not a bell-shaped curve, but for ease of calculation). Also, let's say only 20% of cases become symptomatic, which is a gross underestimation because according to the *Morbidity and Mortality Weekly Report*, 87% will be symptomatic. Thus, at most, we'll have a total of 6,577,200 infected cases in the United States by the end of the outbreak. Given the 2019 US population of 328.2 million, this means a seroprevalence of

2% (roughly by midsummer, very generously calculated). An ideal IgG kit (for serosurveys) with a sensitivity of 100% and specificity of 95%, used in a context of a pretest probability of 2%, would give us a positive predictive value (PPV) of 29% (best-case scenario). Of course, this is for the whole country; some states such as New York can immensely skew the calculations.

The seroprevalence by the end of the outbreak for a state like Ohio can be estimated as 0.7% (based on published state government data as of April 16, 2020). Therefore, the PPV will be 12%. Again, all of these are calculated very generously.

Let's take a recent example from the media on coronavirus infections in Northern California.<sup>12</sup> This study by Stanford University researchers suggested a seroprevalence of 2.5% to 4.2% in Santa Clara county. According to the above calculations and based on the premise for the test performance (sensitivity 100%, specificity 95%), the PPV for the California study would be 33% to 46%, which is translated to a large false alarm. Another way to look at this is to compare their claim with peer-reviewed literature. According to this serosurvey, the actual numbers of cases is estimated to be 50 to 85 times higher than what the county has announced. This roughly means that only 1.17% to 2% of infected individuals become symptomatic.

On a related note, according to a CDC study from the state of Washington, 87% of coronavirus-infected individuals became symptomatic.<sup>13</sup> This finding (87%) is in sharp contrast with the results of the Stanford University serosurvey estimates. A subsequent study by Arons et al reported that 94% of residents of a skilled nursing facility with confirmed status became symptomatic.<sup>14</sup> More recently, the CDC reported that of 4,336 exposed health-care workers (median age 42) with confirmed COVID-19 diagnosis (by RNA testing), only 8% did not report any symptoms.<sup>15</sup> It should also be mentioned that according to an earlier study, also from California, about 5% of individuals with flulike illness tested positive for COVID-19 by RNA testing.<sup>16</sup>

All in all, a careful analysis of harm and cost vs benefit needs to be done prior to conducting such large-scale serosurveys, if needed at all. ■



## REFERENCES

1. **Infectious Diseases Society of America.** IDSA COVID-19 antibody testing primer. Updated April 29, 2020. Accessed May 1, 2020. <https://www.idsociety.org/globalassets/idsa/public-health/covid-19/idsa-covid-19-antibody-testing-primer.pdf>
2. **Wölfel R, Corman VM, Guggemos W, et al.** Virological assessment of hospitalized patients with COVID-2019. *Nature* 2020 April 1. doi:10.1038/s41586-020-2196-x
3. **Bhatraju PK, Ghassemieh BJ, Nichols M, et al.** Covid-19 in critically ill patients in the Seattle region - case series. *N Engl J Med* 2020 March 30. doi:10.1056/NEJMoa2004500
4. **Yan G, Lee CK, Lam LTM, et al.** Covert COVID-19 and false-positive dengue serology in Singapore. *Lancet Infect Dis* 2020 March 4. doi:10.1016/S1473-3099(20)30158-4
5. **Centers for Disease Control and Prevention.** Coronavirus Disease 2019 (COVID-19). April 16, 2020. Accessed May 1, 2020. <https://www.cdc.gov/coronavirus/2019-ncov/hcp/faq.html>
6. **Chan KH, Poon LL, Cheng VC, et al.** Detection of SARS coronavirus in patients with suspected SARS. *Emerg Infect Dis* 2004; 10(2):294–299. doi:10.3201/eid1002.030610
7. **Centers for Disease Control and Prevention.** Discontinuation of transmission-based precautions and disposition of patients with COVID-19 in healthcare settings (interim guidance). March 3, 2020. Accessed May 1, 2020. <https://www.cdc.gov/coronavirus/2019-ncov/hcp/disposition-hospitalized-patients.html>
8. **Shen C, Wang Z, Zhao F, et al.** Treatment of 5 critically ill patients with COVID-19 with convalescent plasma. *JAMA* 2020; 323(16):1582–1589. doi:10.1001/jama.2020.4783
9. **Duan K, Liu B, Li C, et al.** Effectiveness of convalescent plasma therapy in severe COVID-19 patients. *Proc Natl Acad Sci U S A* 2020; 117(17):9490–9496. doi:10.1073/pnas.2004168117
10. **Stockman LJ, Bellamy R, Garner P.** SARS: systematic review of treatment effects. *PLoS Med* 2006; 3(9):e343. doi:10.1371/journal.pmed.0030343
11. **US Food and Drug Administration.** Recommendations for investigational COVID-19 convalescent plasma. April 13, 2020. Accessed May 1, 2020. <https://www.fda.gov/vaccines-blood-biologics/investigational-new-drug-ind-or-device-exemption-ide-process-cber/investigational-covid-19-convalescent-plasma-emergency-ind>
12. **Mason M, Netburn D.** Coronavirus infections could be much more widespread than believed, California study suggests. *Los Angeles Times* April 17, 2020. Accessed May 1, 2020. <https://www.latimes.com/california/story/2020-04-17/coronavirus-antibodies-study-santa-clara-county>
13. **Kimball A, Hatfield KM, Arons M, et al.** Asymptomatic and presymptomatic SARS-CoV-2 infections in residents of a long-term care skilled nursing facility — King County, Washington, March 2020. *MMWR Morb Mortal Wkly Rep* 2020; 69(13):377–381. doi:10.15585/mmwr.mm6913e1
14. **Arons MM, Hatfield KM, Reddy SC, et al.** Presymptomatic SARS-CoV-2 infections and transmission in a skilled nursing facility. *N Engl J Med* 2020 Apr 24 [Epub ahead of print]. doi:10.1056/NEJMoa2008457
15. **CDC COVID-19 Response Team.** Characteristics of health care personnel with COVID-19 — United States, February 12–April 9, 2020. *MMWR Morb Mortal Wkly Rep* 2020; 69(15):477–481. doi:10.15585/mmwr.mm6915e6
16. **Spellberg B, Haddix M, Lee R, et al.** Community prevalence of SARS-CoV-2 among patients with influenzalike illnesses presenting to a Los Angeles Medical Center in March 2020. *JAMA* 2020 [Epub ahead of print]. doi:10.1001/jama.2020.4958

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