



Back to Patient Care program

Providing guidance in the face of uncertainty

Quest Diagnostics understands that the COVID-19 pandemic has had a significant impact on your practice and on your patients. That's why we have created the **Back to Patient Care program**—a solution to help you get your practice back online so that your patients can receive the care they need.

The impact of COVID-19 extends beyond the infection symptoms—since the start of the pandemic, a recent survey showed a 49% drop in primary care visits.¹ This deferred patient care makes it even more important to get your patients back to a regular course of care, which includes important routine testing.

Back to Patient Care program provides guidance that can help you:

- Make informed decisions about your practice and how to appropriately staff
- Get back to the office setting
- Minimize risk of exposure in your work environment
- Treat patients in-office, as appropriate

Our Back to Patient Care program can help your patients and your staff take steps towards managing their COVID-19 exposure, and get the care and treatment they need.

A suggested model to help get your patients back to care with 4 easy steps.

1 

Test staff
with SARS-CoV-2
(COVID-19) molecular*
and antibody tests

2 

Screen patients with questionnaire prior to office visit
for COVID-19
exposure and risk

3 

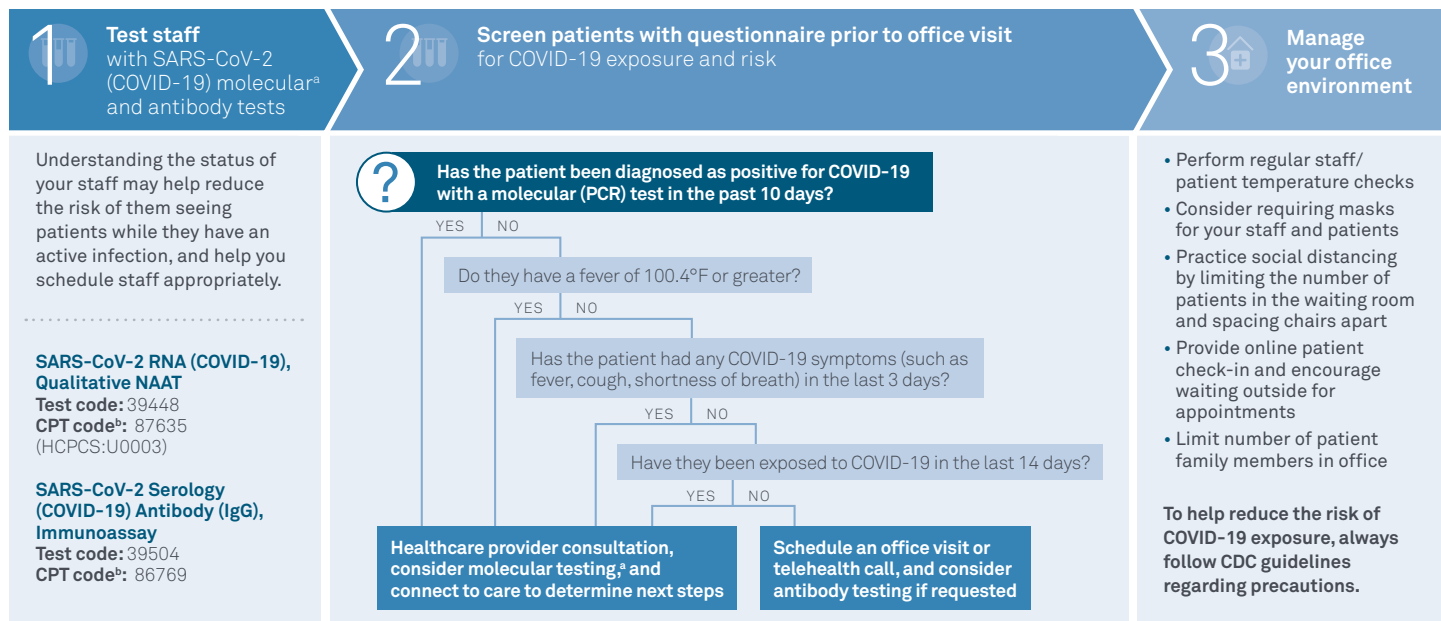
Manage your office environment

4 

Back to caring for your patients

A model for getting back to care

While every practice's model for providing care will vary, we've developed one baseline approach based on CDC guidelines for wellness, chronic, and acute care visits that HCPs may consider as they return back to a more regular schedule of patient care.² This model may change as CDC guidelines and FDA recommendations are updated.



^aSpecimen collection for molecular testing for active COVID-19 infection is not available in our Patient Service Centers.

^bCPT codes are based on American Medical Association guidelines and are for informational purposes only. CPT coding is the sole responsibility of the billing party. Please direct any questions regarding coding to the payer being billed.

Meeting your testing needs

Back to Patient Care is supported by the Quest Diagnostics network of 2,250 Patient Service Centers (PSCs). Our PSCs remain open as an essential service to put your patients' care first every day. **Our Peace of Mind program** has enhanced safety features consistent throughout the day for all patients.

Quest Diagnostics PSCs also offer COVID-19 antibody testing. Staff and patients with a physician's order can make appointments for COVID-19 antibody testing at [QuestDiagnostics.com/Appointment](https://questdiagnostics.com/Appointment).



For more information or to talk to your Quest sales representative, call **1.866.MYQUEST (1.866.697.8378)**.

Delivering Peace of Mind at our PSCs

- Social distancing
- All patients are required to wear a mask or face covering
- Safeguarding patient health with employee masks and PPE
- More frequent cleaning
- Wait by Text program

The antibody tests (sometimes known as the serology tests or IgG tests) are intended for use as an aid in identifying individuals with an adaptive immune response to SARS-CoV-2, indicating recent or prior infection. Results are for the detection of SARS-CoV-2 antibodies. IgG antibodies to SARS-CoV-2 are generally detectable in blood several days after initial infection, although the duration of time antibodies are present post-infection is not well characterized. At this time, it is unknown for how long antibodies persist following infection and if the presence of antibodies confers protective immunity. Individuals may have detectable virus present for several weeks following seroconversion. Negative results do not preclude acute SARS-CoV-2 infection. If acute infection is suspected, molecular testing for SARS-CoV-2 is necessary. The antibody test should not be used to diagnose acute SARS-CoV-2 infection. False positive results for the antibody test may occur due to cross-reactivity from pre-existing antibodies or other possible causes.

- The antibody tests and the molecular tests (together "All tests") have not been FDA cleared or approved;
- All tests have been authorized by FDA under EUAs for use by authorized laboratories;
- The antibody tests have been authorized only for the detection of IgG antibodies against SARS-CoV-2, not for any other viruses or pathogens;
- The molecular tests have been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens; and,
- All tests are only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner

References:

1. Mehrotra A, Chernew M, Linetsky D, Hatch H, Cutler D. What impact has COVID-19 had on outpatient visits? *To the Point (blog), The Commonwealth Fund*. April 23, 2020. <https://doi.org/10.26099/ds9e-jm36>
2. Centers for Disease Control and Prevention. Evaluating and testing persons for Coronavirus Disease 2019 (COVID-19). <https://www.cdc.gov/coronavirus/2019-nCoV/hcp/clinical-criteria.html>. Accessed May 4, 2020.

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