



Coronavirus Disease (COVID-19) Virus Testing

Coronavirus Disease 2019, or COVID-19, is a novel respiratory disease that has been characterized as a pandemic by the World Health Organization. Advanta Analytical Laboratories has developed a rapid polymerase chain reaction (PCR) diagnostic test to identify the virus that causes COVID-19 (referred to as SARS-CoV-2).

Overview - Coronavirus Panel

- SARS-2 coronavirus (COVID-19)
- SARS coronavirus (SARS)
- Human coronavirus (229E)
- Human coronavirus (HKUI)
- Human coronavirus (NL63)
- Human coronavirus (OC43)

TAT

Results are typically available within 24 to 48 hours of your sample being received by the lab.

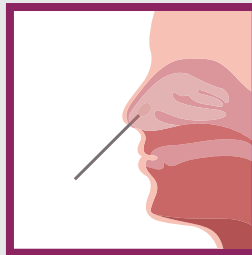
How To Order: PLEASE NOTE!

If you require a sample collection kit, please email us at advanta@aalabs.com. Visit www.aalabs.com/Covid19 for forms and sample collection instructions.

Sample Type

Please see "Specimen Collection Instructions" for accepted sample types and instructions on sample collection and ordering processes.

Nasopharyngeal



Oropharyngeal



Sputum



Disclaimer: The Coronavirus Diagnostic Panel includes the COVID19 test, a modification of the CDC's Emergency Use Authorization (EUA) approved assays. The COVID-19 (N1,N2, and RP genes) test has been validated in accordance with the FDA's Guidance Document (Policy for Diagnostics Testing in Laboratories Certified to Perform High Complexity Testing under CLIA prior to EUA for Coronavirus Disease-2019 during the Public Health Emergency) issued on February 29th, 2020 (updated March 31, 2020). The COVID-19 test in the panel is only authorized for the duration of time the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection of SARS-CoV-2 virus and/or diagnosis of COVID-19 infection under section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.

THE HUMAN ADVANTAGE IN LABORATORY TESTING

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