color

COVID-19 test

Sample Report

PATIENT Jane Doe

DOB: May 25, 1977
Sex: Female

ID: 123456

ORDERING PHYSICIAN

Dr. Jenny Jones
Sample Medical Group
123 Main St.
Sample, CA

SPECIMEN

Type: Nasopharyngeal
Barcode: 223 234234 2343
Collected: Mar 24, 2020
Received: Mar 24, 2020

Report date: March 25, 2020

POSITIVE for SARS-CoV-2.

This means that SARS-CoV-2 (the virus that causes COVID-19) was detected in the patient's sample collected on March 24, 2020.

DETAILS

TEST	RESULTS
SARS-CoV-2 amplification test	POSITIVE (Detected)

To learn more about the technical details of this test, visit color.com/covid19-details.

HELPFUL INFORMATION FOR PATIENT

The CDC has provided useful information on how to care for yourself at home and how others in your household may protect themselves. The CDC has also provided information on when to seek medical attention. Key points are outlined below for your reference, and you can find this information at https://www.cdc.gov/coronavirus/2019-ncov.

As the CDC instructs, before seeking medical care at an office, clinic, or hospital, please alert healthcare providers to the results of this test. However, do not delay seeking care if you are experiencing a medical emergency.

If you are a healthcare professional, first responder, or frontline worker with questions about returning to work, consider contacting your place of employment or local health department in regards to discontinuation of self-isolation, as they may differ from the CDC's guidelines.

CDC: WHEN TO SEEK MEDICAL CARE

- According to the CDC, most people will have mild to moderate illness that may be safely managed at home.
- However, some people with COVID-19 will develop more severe symptoms, and the CDC recommends that individuals experiencing the following symptoms get medical attention immediately:

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CDC: WHEN TO SEEK MEDICAL CARE
(Continued)

- Trouble breathing
- Persistent pain or pressure in the chest
- New confusion or inability to arouse
- Bluish lips or face

- When contacting emergency health services or proceeding to a hospital or clinic, alert the responders at the time of your call, or the facility ahead of your arrival, of your positive test result. **Do not delay care if you are experiencing severe symptoms.**
- Learn more at https://www.cdc.gov/coronavirus/2019-ncov.

CDC: FOR PEOPLE WHO ARE SICK

- The CDC recommends that if you are or might be sick with COVID-19 stay home except to get medical care, and avoid using public transportation if you must leave your home. Even if you have no symptoms, you can pass the infection on to others.
- Separate yourself from other people in your home as much as possible.
- Wear a facemask if you are around other people. If a facemask is unavailable, you may use a bandana or scarf to cover your mouth and nose. If you feel too short of breath to wear a mask, alert those around you so that they may wear a mask.
- Cover your coughs & sneezes with a tissue, clean your hands often with soap and water for at least 20 seconds, and avoid sharing household items (for example, utensils, towels, glasses) as much as possible.
- Clean all 'high-touch' surfaces in your home frequently.
- **Please work with your provider to determine appropriate next steps, including when to end self-isolation.**
- Learn more at https://www.cdc.gov/coronavirus/2019-ncov.

This test is only for use under the Food and Drug Administration's Emergency Use Authorization, but is not FDA cleared or approved. This laboratory developed test has been validated in accordance with CLIA and the FDA's Guidance Document (Policy for Diagnostic Tests for Coronavirus Disease-2019 during the Public Health Emergency Immediately in Effect Guidance for Clinical Laboratories, Commercial Manufacturers, and Food and Drug Administration Staff Document) issued on March 16, 2020. As per the EUA application process, validation data has been submitted to the FDA. FDA's independent review of this validation data is pending. This test is only authorized by FDA for the duration of time the declaration that circumstances exist justifying the authorization of the emergency use of laboratory developed 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.

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