

# FACT SHEET FOR PATIENTS

GenMark Diagnostics, Inc.

cobas eplex<sup>®</sup> respiratory pathogen panel 2 (RP2 Panel)

Updated: April 4, 2024

Coronavirus  
Disease 2019  
(COVID-19)

You are being given this Fact Sheet because your sample(s) was tested for the Coronavirus Disease 2019 (COVID-19) and/or other respiratory infections using the cobas eplex<sup>®</sup> respiratory pathogen panel 2 (RP2 Panel).

This Fact Sheet contains information to help you understand the risks and benefits of using this test for the diagnosis of COVID-19 and/or other respiratory infections caused by pathogens detected by the cobas eplex<sup>®</sup> respiratory pathogen panel 2 (RP2 Panel). After reading this Fact Sheet, if you have questions or would like to discuss the information provided, please talk to your healthcare provider.

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**For the most up to date information on COVID-19 please visit the CDC Coronavirus Disease 2019 (COVID-19) webpage:**

<https://www.cdc.gov/COVID19>

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## What is COVID-19?

COVID-19 is caused by the SARS-CoV-2 virus which is a new virus in humans causing a contagious respiratory illness. COVID-19 can present with a mild to severe illness, although some people with COVID-19 may have no symptoms at all. Older adults and people of any age who have underlying medical conditions have a higher risk of severe illness from COVID-19. Serious outcomes of COVID-19 include hospitalization and death. The SARS-CoV-2 virus can be spread to others not just while one is sick, but even before a person shows signs or symptoms of being sick (e.g., fever, coughing, difficulty breathing, etc.). A full list of symptoms of COVID-19 can be found at the following link:

<https://www.cdc.gov/coronavirus/2019-ncov/symptoms-testing/symptoms.html>.

## What is the cobas eplex<sup>®</sup> respiratory pathogen panel 2 (RP2 Panel)?

The test is designed to detect the virus that causes COVID-19 (SARS-CoV-2), in addition to 17 other pathogens that cause respiratory infections, in nasopharyngeal swab specimens.

## Why was my sample tested?

You were tested because your healthcare provider believes you may have been exposed to the virus that causes COVID-19 based on your signs and symptoms (e.g., fever, cough, difficulty breathing), and/or because:

- You live in or have recently traveled to a place where transmission of COVID-19 is known to occur; or
- You have been in close contact with an individual suspected of or confirmed to have COVID-19

Testing of the samples will help find out if you may have COVID-19 and/or other respiratory infections caused by pathogens detected by the cobas eplex<sup>®</sup> respiratory pathogen panel 2 (RP2 Panel).

## What are the known and potential risks and benefits of the test?

Potential risks include:

- Possible discomfort or other complications that can happen during sample collection.
- Possible incorrect test result (see below for more information).

Potential benefits include:

- The results, along with other information, can help your healthcare provider make informed recommendations about your care.
- The results of this test may help limit the spread of COVID-19 and/or other respiratory infections to your family and those you come in contact with.

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**Where can I go for updates and more information?** The most up-to-date information on COVID-19 is available at the CDC General webpage: <https://www.cdc.gov/COVID19>. In addition, please also contact your healthcare provider with any questions/concerns.

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## What does it mean if I have a positive test result?

If you have a positive test result, it is very likely that you have COVID-19. Therefore, it is also likely that you may be placed in isolation to avoid spreading the virus to others. You should follow CDC guidance to reduce the potential transmission of disease.

There is a smaller possibility that this test can give a positive result that is wrong (a false positive result) particularly when used in a population without many cases of COVID-19. Your healthcare provider will work with you to determine how best to care for you based on the test results along with medical history, and your symptoms.

## What does it mean if I have a positive test result for another respiratory pathogen?

If you have a positive test result for the presence of another respiratory pathogen, it is very likely that you have a respiratory infection. If you have a positive result, your healthcare provider will determine the best way to care for you based on the test results along with other factors in your medical history. There is a very small chance that this test can give a positive result that is wrong (a false positive result). Your healthcare provider will work with you to determine how best to care for you based on the test results, medical history, and your symptoms.

## What does it mean if I have a positive test result for SARS-CoV-2 and another respiratory pathogen?

It is possible for an individual to be infected with SARS-CoV-2 and/or another respiratory pathogen detected by this test at the same time. Your healthcare provider will work with you to determine how best to care for you based on these test results, your medical history, and your symptoms.

## What does it mean if I have a negative test result for SARS-CoV-2 or another respiratory pathogen?

A negative test result for any of the pathogens detected by this test means that these pathogens were not found

in your sample. For COVID-19, a negative test result for a sample collected while a person has symptoms usually means that COVID-19 did not cause your recent illness.

This test can also detect 17 other pathogens that cause respiratory infections. If you have a negative result for the other 17 pathogens, it means that those pathogens were not found in your sample

However, it is possible for this test to give a negative result that is incorrect (false negative) in some people with COVID-19 or another respiratory infection detected by this test. You might test negative if the sample was collected early during your infection. You could also be exposed to COVID-19 or another respiratory illness detected by this test after your sample was collected and then have become infected.

This means that you could possibly still have COVID-19 or another respiratory infection detected by this test even though the test result is negative. If your test is negative, your healthcare provider will consider the test result together with all other aspects of your medical history (such as symptoms, possible exposures, and geographical location of places you have recently traveled) in deciding how to care for you.

It is important that you work with your healthcare provider to help you understand the next steps you should take.

## Is this test FDA-approved or cleared?

No. This test is not yet approved or cleared by the United States FDA, but it has been issued an Emergency Use Authorization (EUA). FDA may issue an EUA when certain criteria are met, which includes that there are no adequate, approved, available alternatives. The EUA for this test is supported by the Secretary of Health and Human Service's (HHS's) declaration that circumstances exist to justify the emergency use of in vitro diagnostics for the detection and/or diagnosis of the virus that causes COVID-19. This EUA will remain in effect (meaning this test can be used) for the duration of

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the COVID-19 declaration justifying the emergency use of in vitro diagnostics, unless it is terminated or revoked by FDA (after which the test may no longer be used).

## What are the approved alternatives?

There are approved/cleared tests for some of the targeted pathogens (e.g., influenza and RSV tests). Any tests that have received full marketing status (e.g., cleared, approved), as opposed to an EUA, by FDA can be found by searching the medical device databases here: <https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/medical-device-databases>. A cleared or approved test should be used instead of a test made available under an EUA, when appropriate and available. FDA has issued EUAs for other tests that can be found at:

<https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization>

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