Start Here
Carefully read all instructions before beginning.
No food/drink (including water)/oral hygiene products for 30 minutes before test.
Complete the entire procedure without delay between steps.
Matrix Reader required. Available separately.

Contents
Cap
Collector
Nasal Swabs (optional)

Power Up
Connect reader to power supply. The center light will turn solid white (not flashing) when ready.

Collect Sample
Choose one preferred sample method:
Saliva or Nasal Swab

Saliva Collection
Deposit a small amount of saliva into the collector.
Fluid level must not go above the black line.
Ignore bubbles.

DO NOT OVERFILL

Nasal Swab Collection
Insert the nasal swab into your nostril until the tip is fully inside. Stop when you meet resistance (about 1 inch for adults, ½ inch for children).
Roll the swab against the inside of your nostril 5 times.
Repeat with other nostril.

Shake to Mix
Shake the collector very hard for 20 seconds to mix.

Shake with method:
5x

Make sure that your sample is mixed well.

Cap Sample
Put the cap on the collector and press down firmly until the cap clicks into place.

CHECK THAT BLACK LINES ARE VISIBLE THROUGH THE OPENINGS ON THE CAP

Run the Test
CONFIRM THAT THE READER IS READY. THE CENTER LIGHT WILL BE SOLID (NOT FLASHING).

Insert sensor into reader until it cannot go any further.

The test will begin automatically and will take 30 minutes.

Attach to Sensor
Open sensor pouch and place sensor on a flat surface.

Flashing White
Test in progress

Flashing Red
Test error, see reverse for troubleshooting

Read Your Results
Use the following visual signifiers to note the results of the test:
Solid green = COVID Negative
Solid red = COVID Positive
Solid purple = Invalid
Repeat test with a new kit.

Discard the Sensor
Pull the sensor out of the reader and discard the sensor. Do not disassemble.
The reader is ready to begin a new test.

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Meaning of Results
A negative test result indicates that SARS-CoV-2, the virus that causes COVID-19, was not detected in your sample. However, it is possible for this test to give a negative result that is not correct (false negative) in some people with COVID-19. Negative results do not exclude SARS-CoV-2 infection and should not be used as the sole basis for treatment of an individual, including infection control decisions. If you have symptoms, contact a healthcare provider for additional testing.

A positive test result indicates that SARS-CoV-2, the virus that causes COVID-19, was detected in your sample. It is very likely that you have COVID-19. Positive test results do not rule out bacterial infection or coinfection with other viruses. Individuals who test positive with the Metrix COVID-19 Test should self-isolate and seek follow-up care with a healthcare professional as additional testing may be necessary.

The Metrix COVID-19 Test detects active COVID-19 infections and does not test for previous infections.

After Your Test
To report your Metrix COVID-19 Test results to public health agencies, please visit: aptitudemetrix.com/subshealth/reporting

If symptoms persist or if you are concerned about your health, please seek follow-up care from a healthcare professional.

For free support, or to obtain a physical copy of the product information card free of charge, please call us at 1-888-934-2253 or email us at support@aptitudemetrix.com.

Fact sheets and FAQs available at AptitudeMetrix.com.

For the most up to date information on COVID-19, please visit: www.cdc.gov/covid-19

Reader Statuses

Starting Up
The reader is starting up. Wait until the center light is solid white before inserting a sensor.

Ready
The reader is ready to start a test.

Test Running
The reader is running a test. Do not remove the sensor or unplug the reader.

Positive Result
The test is complete and SARS-CoV-2 was detected in the sample.

Negative Result
The test is complete and SARS-CoV-2 was not detected in the sample.

Indicates flashing light

Troubleshooting

Invalid Result
The test is complete and the result is invalid. Repeat with a new Metrix COVID-19 Test kit.

Test Error
Remove sensor and firmly press down on collector. Firmly remount sensor into reader. If error persists, discard sensor and use a new test kit.

Canceled Test
The test did not complete. Discard the sensor and run the test with a new Metrix COVID-19 Test Kit.

No Power
Check all electrical connections. The reader is not receiving power.

Hardware Failure
There is an error with the reader. Disconnect and reconnect the power.

Invalid Reader
This is a new Metrix Reader. Insert the sensor.

Test Running
The reader is running a test. Do not remove the sensor or unplug the reader.

Positive Result
The test is complete and SARS-CoV-2 was detected in the sample.

Negative Result
The test is complete and SARS-CoV-2 was not detected in the sample.

Indicates flashing light

Intended Use
The Metrix COVID-19 Test is a qualitative nucleic acid amplification test for the detection of SARS-CoV-2 RNA in nasopharyngeal aspirate (NPA) samples from individuals aged 2 years or older, including individuals without symptoms or other epidemiological reasons to suspect COVID-19. This product has been authorized by FDA under an EUA for in vitro diagnostic use by FDA under an EUA.

This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, which is the virus that causes COVID-19. The test is not authorized for detection or diagnosis of other viral or bacterial pathogens.

This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, which is the virus that causes COVID-19. This product has not been FDA cleared or approved, but has been tested to meet emergency-use criteria and FDA has determined that this product may be used to help prevent the spread of COVID-19. Use of this product has not been studied to determine if the product is effective for SARS-CoV-2 infections. This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, which is the virus that causes COVID-19. This product has not been FDA cleared or approved, but has been authorized for emergency use by FDA under an EUA.

For in vitro diagnostic use. Single use only. Do not use if kit is visibly damaged.

Warnings/Precautions
Potential risks of this test include: (1) Possible discomfort during sample collection, (2) The results of this test may help limit the spread of COVID-19 to your family and others in your community.

The emergency use of this product is authorized for the duration of the declaration by the Director, Center for Food Safety and Applied Nutrition, FDA under section 564(b) of the Federal Food, Drug, and Cosmetic Act. The declaration is terminated in its entirety by authorization to terminate.

The Metrix COVID-19 Test and Metrix Reader are for FDA Emergency Use Authorization (EUA) Only.

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For free support, or to obtain a physical copy of the product information card free of charge, please call us at 1-888-934-2253 or email us at support@aptitudemetrix.com.

Fact sheets and FAQs available at AptitudeMetrix.com.

For the most up to date information on COVID-19, please visit: www.cdc.gov/covid-19

Risks/Benefits
Potential benefits of this test include: (1) The results, along with other information, can help your healthcare provider make the most appropriate decisions for you, (2) The results of this test could limit the spread of COVID-19 from persons who test positive

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For use under Emergency Use Authorization (EUA) only.

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AptitudeMetrix.com

Electrical Instructions For use available at AptitudeMetrix.com

UG-00006 Ver. 1

UG-00005 Ver. 1

UG-00006 Ver. 1
Meet Metrix.
The 24/7 laboratory that lives on your countertop.

For in vitro diagnostic use. For use under Emergency Use Authorization (EUA) only. Metrix Reader™ is FDA authorized for use with the Metrix COVID-19 Test (not included).

Your health, in your hands.

Kit contains: one reusable Metrix reader, quick start guide, power adapter, and USB-C power cable.

This kit is for use with test-specific Metrix kits. Go to www.aptitudemetrix.com for a list of all compatible devices.

Dimensions of Metrix Reader: 105 x 76 x 20 mm
Weight of Metrix Reader: 82 g
Power: 5V DC, 3A

Electronic Instructions For Use are available at AptitudeMetrix.com

For use under Emergency Use Authorization (EUA) Only. This product has not been FDA cleared or approved, but has been authorized for emergency use by FDA under an EUA. This product has been authorized only for detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. The emergency use of this product is 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 Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb-3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

730-00037 Rev B
REF 820-00002 Rev B
LOT RKIT-082022-1
SN LN-1234-12345678-01
730-00040 Rev A

For use under Emergency Use Authorization (EUA) only. Metrix Reader™ is FDA authorized for use with the Metrix COVID-19 Test (not included).
Start Here

Scan QR code for interactive instructions and video demonstration.

Power Up
Connect reader to power supply. The center light will turn solid (not flashing) when ready.

Collect Sample
Open the Metrix COVID-19 Test kit (available separately) if you have not done so already.

The Metrix COVID-19 Test kit will guide you through how to collect and run your sample.

Reader Statuses

Starting Up
The reader is starting up. Wait until the center light is solid white before inserting a sensor.

Ready
The reader is ready to start a test.

Test Running
The reader is running a test. DO NOT remove the sensor or unplug the reader.

Positive Result
The test is complete and SARS-CoV-2 was detected in the sample.

Negative Result
The test is complete and SARS-CoV-2 was NOT detected in the sample.

Invalid Result
The test is complete and the result is invalid. Repeat with a new Metrix COVID-19 Test kit.

For use with the Metrix COVID-19 Test kit (available separately).
For use under Emergency Use Authorization (EUA) only.

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Troubleshooting

Test Error
Remove sensor and firmly press down on collector label. Replace sensor into reader. Firmly reinsert sensor into reader. If error persists, discard sensor and use a new test kit.

No Power
Check all electrical connections. The reader is not receiving power.

Canceled Test
The test did not complete. Discard the sensor and run the test with a new test kit.

Hardware Failure
There is an error with the reader. Disconnect and reconnect the power.

If troubleshooting fails to resolve any problem persist please contact support.

If your Metrix Reader needs to be disposed of, please place in electronic waste.

For support, please contact us at: 1.888.934.2253 support@aptitudemetrix.com Or visit: AptitudeMetrix.com.

Legend of Symbols
- For in vitro diagnostic use
- Do not use if packaging is damaged
- Manufacturer of device
- Direct current (DC) voltage
- Temperature limitations of the product
- Keep dry
- Please contact the instruction manual
- Dispose of in electronic waste
- Certification that the electromagnetic interference from the device is under limits approved by the Federal Communications Commission.
- For in vitro diagnostic use
- Manufacturer’s catalog number
- Manufacturer of device
- Date of manufacture
- REF
- IVD
- No Power
- Keep dry
- Direct current (DC) voltage
- Temperature limitations of the product
- Please consult the instruction manual
- Dispose of in electronic waste
- Indicates flashing light

Warnings and Precautions
Do not use components that are visibly damaged.

The Matrix Reader can be cleaned by wiping the exterior with disinfectant. Do not spray disinfectant into or onto the reader.

If a power failure occurs or if the Metrix Reader is unplugged while the Sensor is inserted, the test result is invalidated. The test should be redone with a new test kit.

Use only the provided power cable and power adapter.

This device complies with Part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) this device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

For in vitro diagnostic use:
This product has not been FDA cleared or approved, but has been authorized for emergency use by FDA under an EUA.

This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens.

The emergency use of this product is 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 Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb-3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

The Metrix COVID-19 Test and Metrix Reader are for FDA Emergency Use Authorization (EUA) only.

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