## TOP

Laboratory accuracy.
Living room ease.

## LEFT FLAP

REF 810-00012 Rev B
LOT CSKIT-082022-1
SN DR3c-082022-1
Use by: 2023-02-20

730-00039 Rev A

730-00033 Rev C

## FRONT

aptitude®

# Metrix® COVID-19 Test

Your health, in your hands

PCR accuracy in 30 minutes
Saliva or nasal swab - your choice

FDA Authorized
For use under Emergency Use Authorization (EUA) only
Requires Metrix Reader or Metrix Reader (Gen 2), sold separately
For in vitro diagnostic use

## BOTTOM

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 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. § 360bbb3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

Aptitude Medical Systems, Inc.
125 Cremona Dr., Ste. 100
Goleta, CA 93117



Electronic instructions for use are available at aptitudemetrix.com
Do not use if package is damaged
Single use item. Do not reuse.
86°F / 59°F Store at 59-86 °F (15-30 °C)

IVD

8 60009 45121 6

## RIGHT FLAP

Your health, in your hands

## BACK

888.934.2253
support@aptitudemetrix.com
aptitudemetrix.com

Metrix is a PCR-accurate molecular test to diagnose or screen for COVID-19 infection.

For use with or without symptoms or other reasons to suspect COVID-19. For self-testing by individuals aged 14+, and by adult users testing other individuals aged 2+.

Single use only. No prescription required. Read instructions before using. Keep out of reach of children. Do not ingest any part of this test kit.

**Accurate**
Lab-quality results at home

**Fast**
Results in 30 minutes

**Easy**
You choose - saliva or swab

This single test kit contains: instructions, collector, cap, sensor, and nasal swab for optional use.

 **Single Test Configuration**

# FRONT

aptitude®

## Metrix® COVID-19 Test

Contains 25 Metrix COVID-19 Tests
Requires Metrix Reader or Metrix Reader (Gen 2), sold separately

# BACK

888.934.2253
support@aptitudemetrix.com
aptitudemetrix.com

Metrix is a PCR-accurate molecular test to diagnose or screen for COVID-19 infection.

For use with or without symptoms or other reasons to suspect COVID-19. For self-testing by individuals aged 14+, and by adult users testing other individuals aged 2+.

Single use only. No prescription required. Read instructions before using. Keep out of reach of children. Do not ingest any part of this test kit.

**Accurate**
Lab-quality results at home

**Fast**
Results in 30 minutes

**Easy**
You choose - saliva or swab

This kit contains components required to run 25 COVID-19 tests: instructions (1), collectors (25), caps (25), sensors (25), and nasal swabs (25) for optional use.

# SIDES (L/R)

A New Generation of Diagnostics

# TOP

aptitude®

## Metrix® COVID-19 Test

Your health, in your hands

PCR accuracy in 30 minutes
Saliva or nasal swab - your choice

FDA Authorized
For use under Emergency Use Authorization (EUA) only
Requires Metrix Reader or Metrix Reader (Gen 2), sold separately
For in vitro diagnostic use

# BOTTOM

730-00044 Rev B

REF 810-00044 Rev A
LOT POC-082022-1
SN CSS5-082022-1
Use by: 2023-08-20

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 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. § 360bbb3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

Aptitude Medical Systems, Inc.
125 Cremona Dr. Ste. 100
Goleta, CA 93117



Electronic instructions for use are available at aptitudemetrix.com

Do not use if package is damaged

Single use item. Do not reuse.

Store at 59-86 °F (15-30 °C)

Σ 25

IVD

8 60009 45122 3

**UG-00005 Ver 2, p. 2 of 10**

**25 Test Configuration**

# Start Here

Carefully read all instructions before beginning.

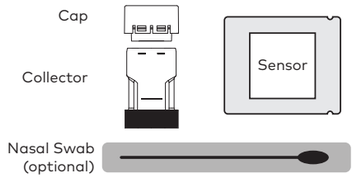
No food/drink (including water)/oral hygiene products for 30 minutes before test.

Complete the entire procedure without delay between steps.

Metrix Reader or Metrix Reader (Gen 2) required, sold separately.

# Contents

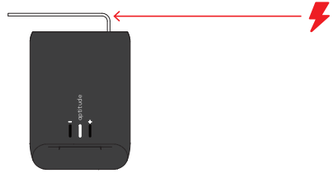
Materials required to run one test:




Scan QR code for interactive instructions and a video demonstration

## 01 Power Up

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



## 02 Collect Sample

Choose one preferred sample method:

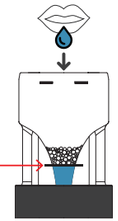
Saliva or Nasal Swab

### 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

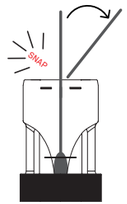
Insert the nasal swab into your nostril until the tip is fully inside. Stop when you meet resistance (about 1 inch for adults, ½ inch for children).


Roll the swab against the inside of your nostril 5 times.

REPEAT WITH OTHER NOSTRIL

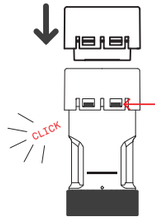
Firmly insert the swab into collector until it cannot go any further.

Snap off and discard the swab handle.




## 03 Cap Sample

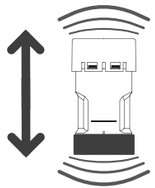
Put the cap on the collector and press down firmly until the cap clicks into place.




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

## 04 Shake to Mix

Shake the collector very hard for 20 seconds to mix.



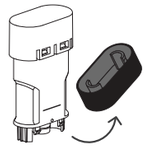
MAKE SURE THAT YOUR SAMPLE IS MIXED WELL.

## 05 Attach to Sensor

Open sensor pouch and place sensor on a flat surface.



Remove the black plastic cover from the bottom of the collector.



Firmly insert the collector into the sensor until it clicks and fluid enters the sensor.

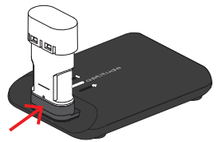
PRESS FIRMLY TO LOCK IN PLACE


## 06 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 further.

The test will begin automatically and will take 30 minutes.



Flashing White

Test in progress

Flashing Red

Test error, see reverse for troubleshooting

## 07 Read Your Results

Use the following visual signifiers to note the results of the test:

Solid green = COVID Negative
SARS-CoV-2 was not present.

Solid red = COVID Positive
SARS-CoV-2 was present.

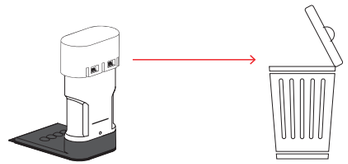
Solid purple = Invalid
Repeat test with a new kit.

IF PROOF OF TEST IS NEEDED, TAKE A PHOTO OF YOUR RESULT. THE RESULT WILL DISPLAY UNTIL THE SENSOR IS REMOVED.

## 08 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.



# 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 preclude 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 co-infection 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/publichealth/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.

aptitude

Electronic instructions for use available at aptitudemetrix.com



Aptitude Medical Systems, Inc.
125 Cremona Dr. Ste. 100
Goleta, CA 93117

# 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 reinsert 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.

## Hardware Failure

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

## No Power

Check all electrical connections. The reader is not receiving power.

If troubleshooting fails to resolve any problem, please contact support. In the event that your Metrix Reader needs to be disposed of, please place in electronic waste.

# Risks/Benefits

- Potential risks of this test include: (1) Possible discomfort during sample collection, (2) Possible incorrect test results.
- Potential benefits of this test include: (1) The results, along with other information, can help your healthcare provider make informed recommendations about your care, (2) The results of this test may help limit the spread of COVID-19 to your family and others in your community.



# Intended Use

The Metrix COVID-19 Test (Metrix) is a single-use molecular in vitro diagnostic test for the qualitative detection of nucleic acid from SARS-CoV-2, the virus that causes COVID-19. This test is authorized for non-prescription home use with anterior nasal (nares) swab and saliva specimens, self-collected from any individual aged 14 years or older, or adult-collected from any individual aged 2 years or older, including individuals without symptoms or other epidemiological reasons to suspect COVID-19.

This test utilizes nucleic acid amplification technology, similar to PCR, for the detection of SARS-CoV-2. SARS-CoV-2 viral RNA is generally detectable in anterior nasal (nares) swab and saliva samples during the acute phase of infection.

Positive results indicate the presence of viral RNA, but clinical correlation with past medical history and other diagnostic information is necessary to determine infection status. Positive results do not rule out bacterial infection or co-infection with other viruses. The agent detected may not be the definite cause of disease. Individuals who test positive with the Metrix COVID-19 Test should self-isolate and seek follow-up care with their physician or healthcare provider as additional testing may be necessary.

Negative results from saliva samples are presumptive and should be confirmed by molecular testing of an alternative sample type if clinically indicated. Negative results do not preclude SARS-CoV-2 infection and should not be used as the sole basis for treatment or management decisions for the individual, including infection control decisions. Negative results should be considered in the context of an individual's recent exposures, history, and the presence of clinical signs and symptoms consistent with COVID-19. Individuals who test negative and continue to experience COVID-19-like symptoms of fever, cough, and/or shortness of breath may still have SARS-CoV-2 infection and should seek follow-up care with their physician or healthcare provider.

Individuals should report all results obtained with this product to their healthcare provider and the Aptitude secure web portal. This Aptitude secure web portal will report all test results received from individuals who use the authorized product to relevant public health authorities in accordance with local, state, and federal requirements using appropriate LOINC and SNOMED codes, as defined by the Laboratory In Vitro Diagnostics (LIVD) Test Code Mapping for SARS-CoV-2 Tests provided by the CDC.

The Metrix COVID-19 Test is only for use under the Food and Drug Administration's Emergency Use Authorization.

# Warnings/Precautions

- For in vitro diagnostic use. Single use only. Do not use if kit is visibly damaged.
- 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.
- For more information on Emergency Use Authorization, please visit: https://www.fda.gov-/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization
- Store between 59 °F (15 °C) and 86 °F (30 °C).
- Do not ingest. Keep away from children. Contains Triton X-100 (0.1%), which can be harmful if swallowed or cause skin irritation or serious eye damage. If the blue liquid contacts skin or eyes, flush with copious amounts of water. If irritation persists, call Poison Control at 1.800.222.1222.
- Once assembled, do not attempt to diassesmble or open the cap/collector/sensor assembly.
- This device was not validated for use in color vision impaired individuals, including individuals with red/green vision impairment.
- The test is best used in a room with adequate lighting away from glare and the Metrix Reader and Metrix Reader (Gen 2) should be used on a level surface without movement.
- If a power failure occurs or if the Metrix Reader or Metrix Reader (Gen 2) is unplugged while the sensor is inserted, the test result is invalid. You should retest using a new test.

501-00000 Rev B

# Metrix® COVID-19 Test Instructions

IVD

For use under Emergency Use Authorization (EUA) only

## TOP EXTERIOR

aptitude™

# Metrix™ Reader

FDA authorized

For use with the Metrix COVID-19 Test (not included).
For in vitro diagnostic use. For use under Emergency Use Authorization (EUA) only.

## FRONT EXTERIOR

Your health,
in your hands

## BACK EXTERIOR

aptitudemetrix.com
888.934.2253
support@aptitudemetrix.com

## BOTTOM EXTERIOR

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

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

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

Aptitude Medical Systems
125 Cremona Dr. Suite 100
Goleta, CA 93117

59°F 85°F



Electronic Instructions For Use are Available at AptitudeMetrix.com

REF 820-00002 Rev B
SN LN-1234-12345678-01
LOT RKIT-082022-1

730-00040 Rev A

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. § 360bbb3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

IVD

8 60009 45120 9

730-00037 Rev B

## TOP INSIDE FLAP

Meet Metrix.
The 24/7 laboratory
that lives on your countertop.

# Metrix™ Reader

Quick Start Guide

IVD

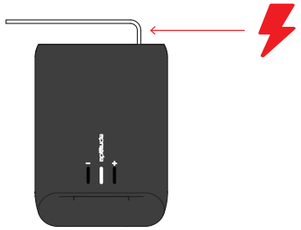
For use with the Metrix COVID-19 Test kit (available separately).
For use under Emergency Use Authorization (EUA) only.

# 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

Indicates flashing light

### 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.

## Troubleshooting

### Test Error

Remove sensor and firmly press down on collector. Firmly reinsert 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 test kit.

### Hardware Failure

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

Indicates flashing light

### No Power

Check all electrical connections. The reader is not receiving 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:

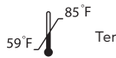
1.888.934.2253
support@aptitudemetrix.com

Or visit: AptitudeMetrix.com.

## Legend of Symbols

IVD — For vitro diagnostic use

Direct current (DC) voltage




Temperature limitations of the product

Keep dry

Please consult the instruction manual

Dispose of in electronic waste

Do not use if packaging is damaged

Manufacturer of device

Date of manufacture

REF — Manufacturer's catalog number

Certification that the electromagnetic interference from the device is under limits approved by the Federal Communications Commission.



## Warnings and Precautions

- Do not use components that are visibly damaged.
- The Metrix 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

aptitude™

# TOP EXTERIOR

aptitude®

Metrix® Reader

Gen 2

FDA authorized

For use with specific Metrix tests under EUA, sold separately

For use under Emergency Use Authorization (EUA) only

# BACK EXTERIOR

888.934.2253

support@aptitudemetrix.com

aptitudemetrix.com

# TOP INSIDE FLAP

Meet Metrix.
The 24/7 laboratory
that lives on your countertop.

↓

# BOTTOM EXTERIOR

Kit contains: one reusable Metrix Reader (Gen 2), quick start guide, power adapter, and USB-C power cable.

This product is for use with specific Metrix tests under EUA.
Go to www.aptitudemetrix.com for a list of all compatible devices.

Read instructions before using.
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

86°F (30°C)
59°F (15°C)
Store at 59-86 °F (15-30 °C)

Aptitude Medical Systems, Inc.
125 Cremona Dr. Ste. 100
Goleta, CA 93117



For use with specific Metrix tests under Emergency Use Authorization (EUA) only. This product has not been FDA cleared or approved, but has been authorized by FDA under an EUA. When used in combination with the Metrix COVID-19 Test: This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. When used in combination with the Metrix COVID-19/Flu Test: This product has been authorized only for the detection and differentiation of nucleic acid from SARS-CoV-2, influenza A, and influenza B, 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.

IVD

REF 820-00009 Rev A

SN LN-1234-12345678-01

LOT RKIT-082022-1

730-00040 Rev A

730-00073 Rev A

# FRONT EXTERIOR

Your health,
in your hands

# Metrix® Reader

Quick Start Guide

Gen 2

IVD

For use with specific Metrix tests under EUA, sold separately

For use under Emergency Use Authorization (EUA) only

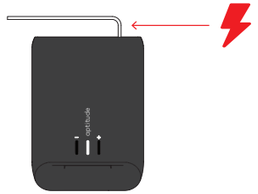
# Start Here

For more information about how to use the Metrix Reader, please scan the QR code with your mobile device or visit:

aptitudemetrix.com/reader

## 01 Power Up

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



## 02 Collect and Run Your Sample

Open your Metrix Test kit (available separately) if you have not done so already. The instructions within the kit will guide you through how to collect and run your sample.

## 03 Read Your Results

Please refer to the instructions included in your Metrix Test kit to interpret your test results.



# Troubleshooting

### 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.

### Invalid Result

The test is complete and the result is invalid. Repeat with a new Metrix Test kit.

Indicates flashing light

### Test Error

Remove Sensor and firmly press down on Collector. Firmly reinsert 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 Test kit. Ensure you are using the correct Metrix Reader for the specific EUA Metrix test under use.

### Hardware Failure

There is an error with the Reader. Disconnect and reconnect the power.

If troubleshooting fails to resolve any problem, contact support.

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

For support, please contact us at:

888.934.2253
support@aptitudemetrix.com
aptitudemetrix.com

## Legend of Symbols

IVD — For in vitro diagnostic use

Direct current (DC) voltage

Storage temperature limitations of the product

Keep dry

Please consult the instruction manual

Dispose of in electronic waste

Do not use if packaging is damaged

Manufacturer of device

Date of manufacture

REF — Manufacturer's catalog number

Certification that the electromagnetic interference from the device is under limits approved by the Federal Communications Commission.

## Warnings/Precautions

- Do not use components that are visibly damaged.
- The Metrix 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.
- Store the reader in a secure location and do not use if the Reader shows signs of damage or tampering.
- 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.
- For use with specific Metrix tests under Emergency Use Authorization (EUA) only.
- For use under EUA only.
- This product has not been FDA cleared or approved, but has been authorized by FDA under an EUA.
- When used in combination with the Metrix COVID-19 Test: This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens.
- When used in combination with the Metrix COVID-19/Flu Test: This product has been authorized only for the detection and differentiation of nucleic acid from SARS-CoV-2, influenza A, and influenza B, 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.



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