

# 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](https://aptitudemetrix.com/reader)



### 01 Power Up

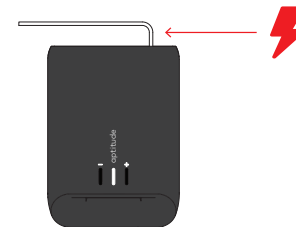
Connect Reader to power supply. The center light will turn solid (not flashing) when ready.

### 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.



### 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.





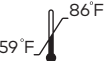






For support, please contact us at:

888.934.2253  
[support@aptitudemetrix.com](mailto:support@aptitudemetrix.com)  
[aptitudemetrix.com](https://aptitudemetrix.com)



Indicates flashing light

Legend of Symbols

	For in vitro diagnostic use		Do not use if packaging is damaged
	Direct current (DC) voltage		Manufacturer of device
	Storage temperature limitations of the product		Date of manufacture
	Keep dry		Manufacturer's catalog number
	Please consult the instruction manual		Certification that the electromagnetic interference from the device is under limits approved by the Federal Communications Commission.
	Dispose of in electronic waste		

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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