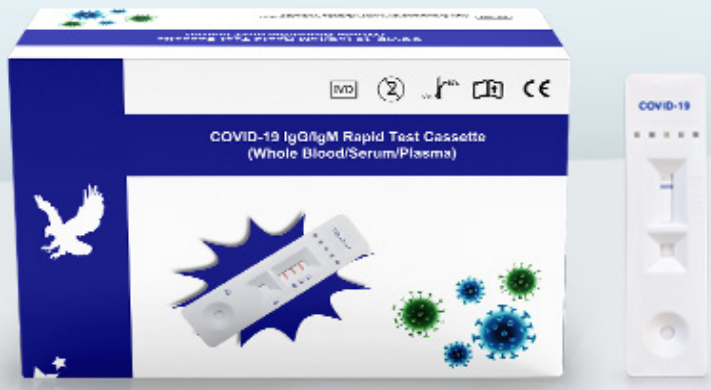
Coronavirus
(COVID-19) Rapid Test
For Professional Use



NOW AVAILABLE FOR PRE-ORDER*

Our easy-to-use COVID-19 test kit provides results in just 2-5 minutes

COST-EFFECTIVE **FAST** **ACCURATE**

HOW ACCURATE IS THE CORONAVIRUS (COVID-19) RAPID TEST?

Our test has shown in clinical evaluation to have a total agreement of **97.19%** and a kappa value of 0.94.

TEST REAGENT	REFERENCE REAGENT		TOTAL
	POSITIVE	NEGATIVE	
Positive	245	4	249
Negative	16	439	455
Total	261	443	704

Results of the first clinical evaluation, published on March 12, 2020

Clinical evaluation took place across 10 hospitals in China during February 2020 and included 704 tests. Ongoing clinical evaluation and other information is available upon request.

COVID-19 "CORONAVIRUS" IGG/IGM RAPID TEST KIT

To support public health & safety, we are offering a proven, cost-effective option to detect and help stop the spread of coronavirus.

- Rapid results in 2-5 minutes
- Detection Window IgM: Symptomatic 3-5 days, Asymptomatic 7 days
- Sample: Test can work with whole blood, plasma and serum samples.
- Storage: The kit can be stored at room temperature or refrigerated (2-30°C).
- Shelf Life: 24 months from manufacture date
- Sold in packs of 100
- Forensic/Professional Use Only**
- Pending FDA review***

AVAILABLE UPON REQUEST:

- Package Insert
- Letter of Intent
- Stability Report
- Clinical Evaluation Report

MORE INFO ABOUT OUR RAPID CORONAVIRUS (COVID-19) TEST KIT

For additional information, such as instructions, how-to videos, clinical trial info, the product insert, and CE documentation, ask your sales representative or visit:

AMPLICON

*Pre-Order: Products estimated to ship from U.S. distribution April 1st

**Professional use only: Tests should be conducted by a licensed phlebotomist, or a medical professional. Verification of use case prior to shipping is mandatory.

***Warning: This Coronavirus instant test kit has not been reviewed by the FDA and results from antibody testing should not be used as the sole basis to diagnose or exclude SARS-CoV-2 infection or to inform infection status. The FDA has allowed the sale and use of these kits in the United States prior to FDA EUA approval, for laboratory and healthcare workers at the point-of-care, during the public health emergency. More information can be found from the FDA at https://www.fda.gov/media/135659/download

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