

THE INTELISWAB™ COVID-19 RAPID DIAGNOSTIC TEST



The remarkably simple COVID-19 test in 30 minutes. Now authorized for OTC use, prescription home use and POC use.

Product Features

- **Emergency Use Authorization (EUA)** for over-the-counter use, prescription home use and point-of-care use
- **Fast Results**—Provides visual results in approximately 30 minutes
- **Simple Sample Collection**—Unique design incorporates a built-in swab into the test stick
- **Easy Process**—No confusing steps, batteries or mailing to labs
- **Accurate**—Overall test performance: Positive Percent Agreement 84%, Negative Percent Agreement 98
- **Practical**—9 month shelf life initially on all tests and no need for ancillary supplies
- **Designed and Developed in the U.S.A.**



Order Information

Configuration	Inteliswab™ COVID-19 Rapid Test	Inteliswab™ COVID-19 Rapid Test Rx	Inteliswab™ COVID-19 Rapid Test Pro
	Over-the-Counter	Prescription Home Test	Point-of-Care
Catalog Number	1001-0616	1001-0620	1001-0614
Test Devices	24 x 2-test kits	24 x 1-test kits	25 tests

This product has not been FDA cleared or approved; but has been authorized by FDA under an EUA.

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

Learn More Today



CORREAL LOGISTICS

YOUR SINGLE SOURCE

TEST KITS PARTNER

CORREA LOGISTICS Offers:

- ✓ Unmatched Range of Covid-19 Testing Options
- ✓ Shipping within 24 hours

OTHER COVID-19 TESTING OPTIONS

Mesa Biotech™ COVID-19 RAPID MOLECULAR TEST

- The Accula™ System is a PCR-based test for Flu A/B and COVID-19
- 30-minute test operating at the speed of antigen testing
- FDA Emergency Use Authorization (EUA)



Status™ COVID-19/FLU A&B ANTIGEN TEST

- Market leading accuracy
 - COVID-19 — Sensitivity 93.9%, Specificity 100%
 - Flu A — Sensitivity 91.4%, Specificity 95.7%
 - Flu B — Sensitivity 87.6%, Specificity 95.9%
- 15-minute test
- FDA Emergency Use Authorization (EUA)



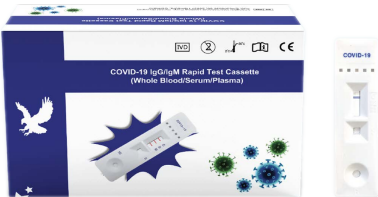
CareStart™ COVID-19 RAPID ANTIGEN TEST

- Large scale COVID-19 screening with anterior nares testing
- 10-minute test
- FDA Emergency Use Authorization (EUA)



Healgen™ COVID-19 RAPID TEST SEROLOGY KIT

- Multi-variable analysis of immunoglobulin IgG & IgM
- 10-minute test
- FDA Emergency Use Authorization (EUA)



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CORREA
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