COVID-19 Test & MOLECULAR DETECTOR

WHERE SPEED, EFFICIENCY, AND ACCURACY ALIGN

FEATURES & BENEFITS

COST-EFFECTIVENESS ✓ Superior functionality and performance at an affordable price.

ACCURACY ✓ Highly accurate test results, minimizing chances of errors and enhancing diagnostic confidence. The AscencioDx COVID-19 Test targets three unique locations on the N-gene to minimize the risk of mutation escape.

EASY & CONVENIENT ✓ The CLIA-waived AscencioDx system is incredibly user-friendly and does not require extensive training, making it a suitable choice for all healthcare settings.

TIME TO RESULT ✓ Receive actionable positive test results in as little as 20 minutes.

CALL 800-676-5565
FOR MORE INFORMATION

JANT PHARMACAL CORPORATION
Laboratory & Point of Care Diagnostics

SIMPLE-TO-PERFORM TEST PROCEDURE

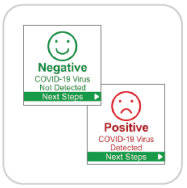
Swab

Stir

Transfer

Place

Close Lid & Wait



Get Result

Share

PRODUCT APPLICATIONS

Assisted Living

Government

Lab

Occupational Health

Pharmacy

Physician Office

Point of Care

Urgent Care

AscencioDx TECHNICAL APPROACH

Test Target Technology

- RT-LAMP molecular analysis
- Real-time SARS-CoV-2 viral RNA detection via fluorescence signals
- Novel enzyme and probe design (patent pending)
- Redundancy: detects 3 distinct regions of the SARS-CoV-2 viral genome, robust against mutation
- RNA internal control in every test flags problems due to sample mishandling

CATALOG NUMBERS

Item	Catalog #
The AscencioDx Molecular Detector - Individual	IDADX100000
The AscencioDx Molecular Detector - Case (10 Detectors)	IDADX100200
The AscencioDx Molecular Detector - Master Pack (4 cases)	IDADX101100
The AscencioDx COVID-19 Test - Master Pack (40 tests)	IDADX100400
The AscencioDx COVID-19 Starter Kit - 1 Detector & 1 Test Master Pack	IDADXSTARTER

Note: This product has not been FDA cleared or approved, but has been authorized for emergency use by FDA under an EUA for use by authorized laboratories. 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.

Authorized Distributor:
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