



Our lab – at the tip of your finger

INTRODUCING

**ADEXUSDx<sup>®</sup> COVID-19**

**Total Antibody Rapid Test**

Featuring Touch-to-Test Technology

*Sample is applied directly to the tip of the test*

The ADEXUSDx<sup>®</sup> COVID-19 Test is a rapid serology test that detects total antibodies to SARS-CoV-2. It provides in-field, lab accurate results within 15 minutes, using as little as a drop of blood.

### **SIMPLE**

- **NO** buffers or external reagents needed
- **NO** lab equipment
- **NO** in demand ancillary items
- **NO** refrigeration

### **ACCURATE**

- Highly sensitive and specific
- Detects total antibody (IgG, IgM and IgA)
- Used to identify adaptive immune response (indicating recent or prior infection)

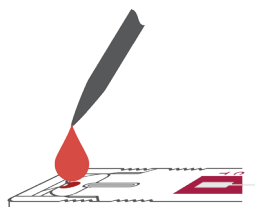


This product has not been FDA cleared or approved but has been authorized for emergency use by FDA under an EUA for use by authorized laboratories; This product has been authorized only for detecting the presence of total antibodies to SARS-CoV-2, not for any other viruses or pathogens; The emergency 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.

# Administering Test\*

## LABORATORY PIPETTE METHOD

1. Collect 40 µL of sample using a laboratory pipette device.



2. Apply sample to the test.

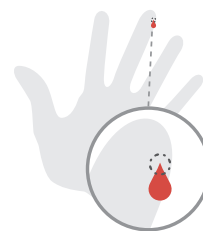
3. Set timer for 15 minutes and read result.



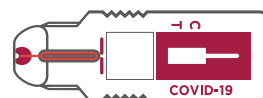
## FINGER STICK METHOD

For use under Emergency Use Authorization only

1. Perform finger stick.



2. Touch the drop of blood to the tip of the test.

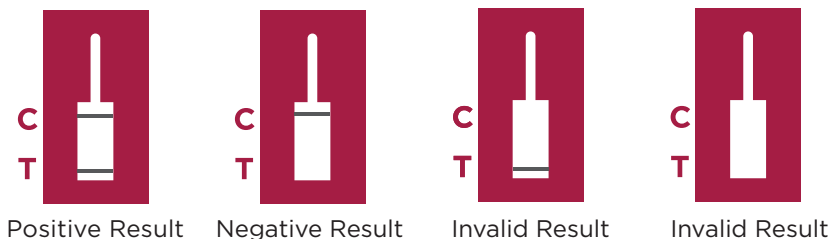


3. Set timer for 15 minutes and read result.



## INTERPRETATION OF RESULTS

A positive result has TWO LINES, one at the “C” (Control Line) and one at the “T” (Test Line). The intensity of the lines may vary. Look carefully as lines might be faint.



\* For complete instructions, refer to the Product Insert before administering the test.

Visit [C19Development.com](http://C19Development.com) for more information.

Description	Quantity	Catalog Number
ADEXUSDx® COVID-19 Test, Box	25	8075-25
ADEXUSDx® COVID-19 Test w. Lancets, Box	25	8075-25-FS
ADEXUSDx® COVID-19 Test, Box	50	8075-50
ADEXUSDx® COVID-19 Test w. Lancets, Box	50	8075-50-FS
ADEXUSDx® COVID-19 Test, Case	1000	8075-1000
ADEXUSDx® COVID-19 Test w. Lancets, Case	1000	8075-1000-FS
ADEXUSDx® COVID-19 Control Set, Box	1	475.002



Testing of venous whole blood (dipotassium EDTA), plasma (dipotassium EDTA), and serum specimens is limited to laboratories certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 U.S.C 263a, that meet requirements to perform moderate or high complexity tests. Testing of fingerstick whole blood specimens is limited to laboratories certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 U.S.C. §263a, that meet the requirements to perform high, moderate, or waived complexity tests. Testing of fingerstick whole blood specimens is authorized for use at the Point of Care (POC), i.e., in patient care settings operating under a CLIA Certificate of Waiver, Certificate of Compliance, or Certificate of Accreditation.