Autumn Diagnostics

REDEFINING SIMPLE

Point of Care COVID-19 Antigen Testing

FDA Emergency Use Authorized

MADE IN USA

SCov-2 Ag *Detect*™ Rapid Test

Patent Design

- Direct nasal swab to cassette format (Patent pending)

Easy to Use

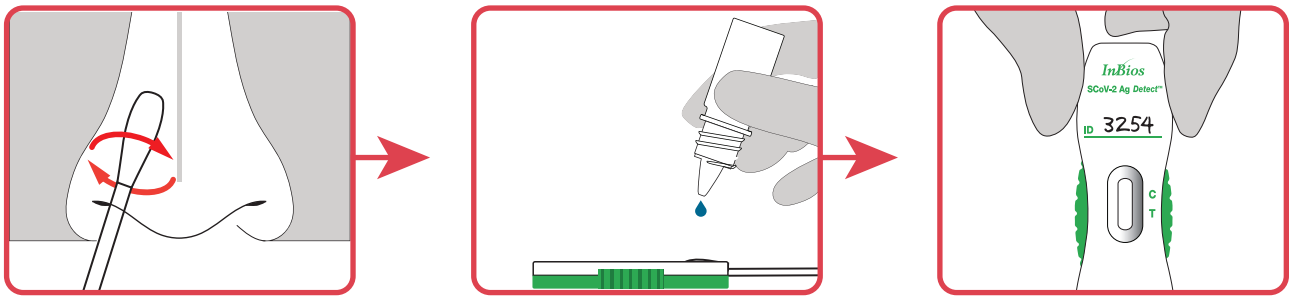
- No transport media step or instrumentation needed

Versatile

- For symptomatic and asymptomatic serial testing
- Can be performed in a variety of settings, from physician offices to school health clinics

Fast & Accurate

- Results in 20 minutes



Swab | Test | Results

See Instructions for Use for complete details.

Exclusive Distributer of the POC InBios SCoV-2 Ag Detect™ in the USA.

Autumn Diagnostics LLC
702 E. MAIN STREET
Jackson, OH US 45640

+1 866 2019503
orders@autumndiagnostic.com

autumndiagnostic.com

SCov-2 Ag *Detect*™ Rapid Test

KEY FEATURES / PERFORMANCE

- Point of Care: Can be performed in a variety of settings, from physician offices to school health clinics, with symptomatic or asymptomatic individuals.
- Identifies acute infection in symptomatic patients with high accuracy
- Uses direct nasal samples without transport media.
- Includes everything required to perform test – no instrumentation needed.
- Includes 50 tests, swabs for sample collection, and positive and negative controls.
- Simple to use and requires minimal training.
- Room temperature storage.
- Fast results in ~20 minutes.
- **100% manufactured in the USA.**

PACKAGING:

Catalog No.	Format	Quantity/Kit	Time to Result	Sample type	Storage	Shelf life
SP-INB-RC	ICT(cassette)	50	~20 minutes	Direct nasal swabs	Room Temperature(15-30°C)	13 months
SP-INB-20	ICT(cassette)	20	~20 minutes	Direct nasal swabs	Room Temperature(15-30°C)	13 months

IMPORTANT INFORMATION:

This product has not been FDA cleared or approved, but has been authorized by FDA under an EUA for use by authorized laboratories.
This product has been authorized only for the detection of proteins from SARS-CoV-2, not for any other viruses or pathogens; and,
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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