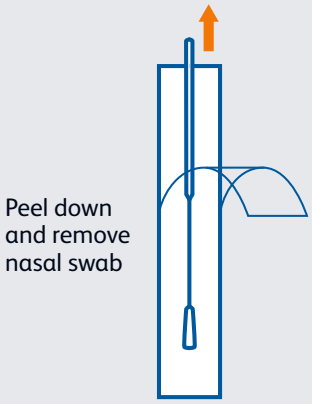
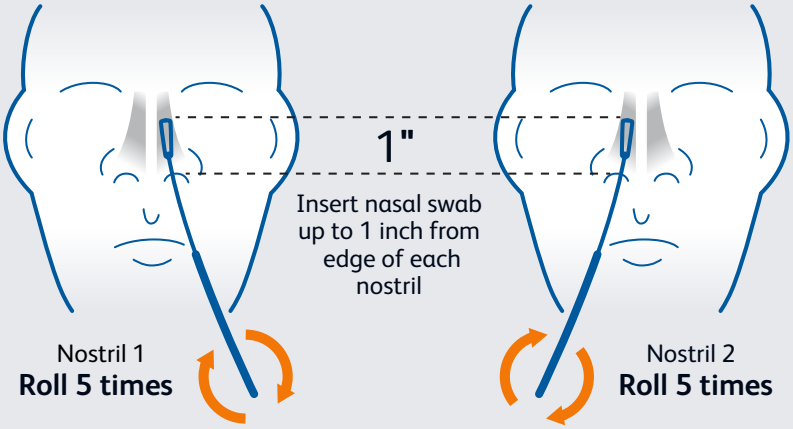
BD Veritor™ Plus System

Rapid point-of-care COVID-19 testing*

1 Sample collection



Peel down and remove nasal swab



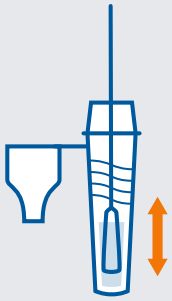
Insert nasal swab up to 1 inch from edge of each nostril

Nostril 1
Roll 5 times

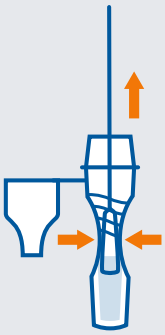
Nostril 2
Roll 5 times

2 Sample preparation

Remove cap from tube and insert nasal swab



Plunge swab up and down 15 seconds



Pinch tube slightly to extract liquid while removing swab

Firmly place cap on tube

Dos and don'ts of sample collection

- Do test sample immediately.
- Use only swabs and reagents provided with the kit.
- Refer to: Interim Guidelines for Collecting, Handling and Testing Clinical Specimens from persons for COVID-19 at https://www.cdc.gov/coronavirus/2019-ncov/lab/guidelines-clinical-specimens.html

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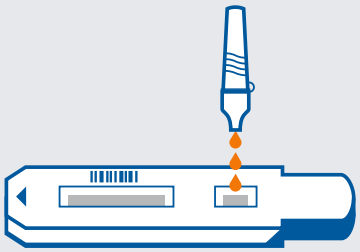
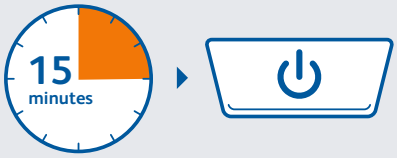
BD

3 Sample testing

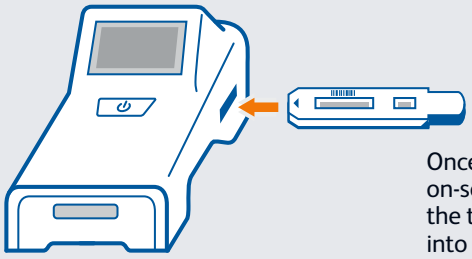
Analyze Now mode



Add 3 drops to the test device sample well



Wait 15 minutes and then press the power button

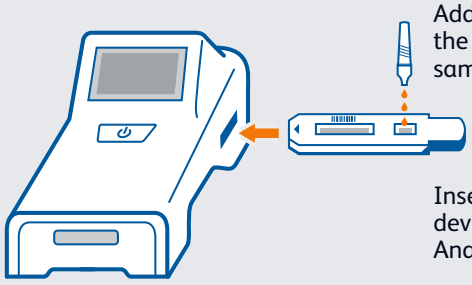


Once prompted on-screen, insert the test device into the Analyzer

Walk Away mode

Press the power button.

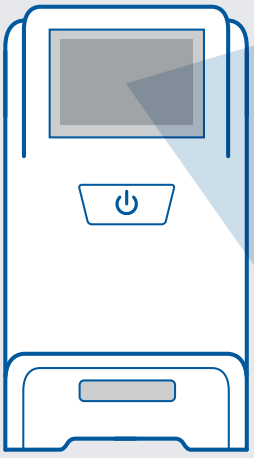
When prompted, double-click for Walk Away mode



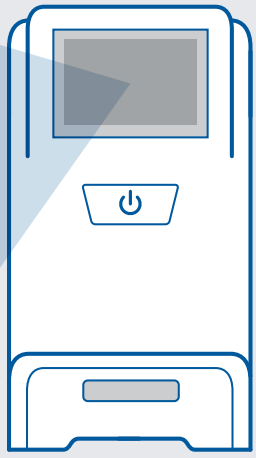
Add 3 drops to the test device sample well

Insert the test device into the Analyzer

Results are displayed within seconds



Results are displayed after 15 minutes



CoV2: –
(Negative)

or

CoV2: +
(Positive)

or

Invalid

For combined COVID-19 & Flu A+B testing, results for Flu A and Flu B will also be displayed on-screen with SARS-CoV-2 results.

BD Veritor™ System for Rapid Detection of SARS-CoV-2 & Flu A+B Instructions for Use, package insert. Franklin Lakes, NJ: Becton, Dickinson and Company.

Emergency Use Authorization Information for the BD Veritor™ SARS-CoV-2 and SARS-CoV-2 & Flu A+B assays:

- These products have not been FDA cleared or approved; but have been authorized by FDA under EUA for use by authorized laboratories
- The BD Veritor™ System for Rapid Detection of SARS-CoV-2 has been authorized only for the detection of proteins from SARS-CoV-2, not for any other viruses or pathogens; the BD Veritor™ System for Rapid Detection of SARS-CoV-2 & Flu A+B has been authorized only for the detection of proteins from SARS-CoV-2, influenza A and influenza B, not for any other viruses or pathogens; and,
- These products are 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.