



COVID-19 saliva testing

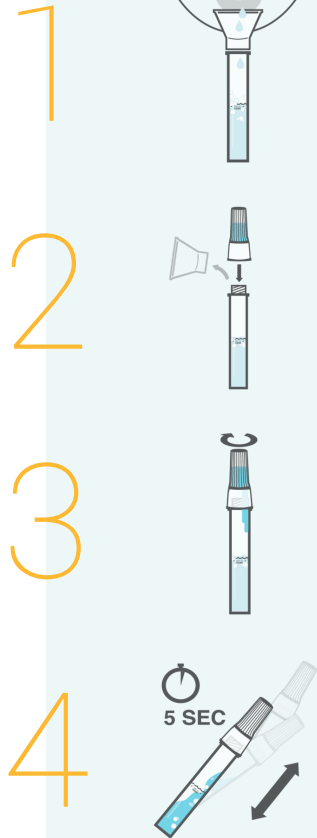
IBX has gained the First Emergency Use Authorization from the FDA for the use of Saliva Collection as part of its COVID-19 molecular diagnostics test

A Game Changer

Infinity Biologix, formerly Rutgers' RUCDR Infinite Biologics, has launched a test for the SARS-CoV-2 coronavirus and is using its automation experience and infrastructure to test as many as tens of thousands of samples daily.

Why saliva testing?

- Research is showing that saliva as a sample collection method, is as sensitive as nasopharyngeal swabs in detecting SARS-CoV-2 in COVID-19 patients
- It has the potential to reduce the risk and challenges associated with nasopharyngeal swabs, which require trained personnel and may show variability of collection
- Partnering with key suppliers enables IBX to scale quickly and use fully automated processing, extraction and testing methodologies for fast results without compromising accuracy or precision
- It enables health care workers to self-collect a sample for return and testing at our high complexity laboratory and subsequently release themselves from quarantine, safely returning back to work
- It allows for self-collection of samples at home, which are then shipped to the Infinity BiologiX laboratory for testing.



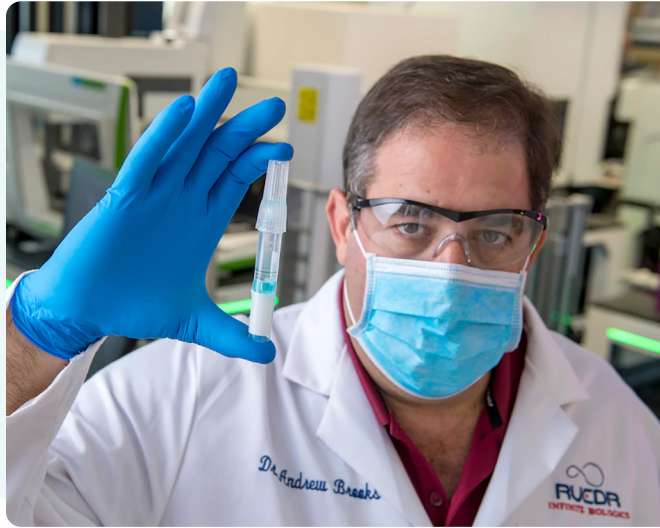
You can find full Instructions For Use on our website:

ibx.bio/covid-19/IFU



We are processing tens of thousands of samples daily with a capacity of fifty thousand per day and an ability to ramp our capacity within weeks.

Dr Andrew Brooks
Chief Executive Officer



Scalability

Partnering with key suppliers enables IBX to scale quickly and use fully automated processing, extraction and testing methodologies for fast results without compromising Accuracy or Precision.



It's definitely going to be a game-changer

Dr. Christina Tan
New Jersey State Epidemiologist

For more information our website

ibx.bio



- This product has not been FDA cleared or approved;
- This product has been authorized by FDA under an EUA for use by the authorized laboratory;
- This product has been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens; and
- 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 authorization is terminated or revoked sooner



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