

# LABCORP AT HOME

## COVID-19 TEST COLLECTION SERVICE



## LabCorp At Home Service Offers Convenient and Easy Access for Patient Care



LabCorp At Home for COVID-19 PCR testing can be ordered through any standard LabCorp account using an interfaced EMR system or LabCorp Link™. Providers will be prompted to enter the patient's home address and contact information. A test kit is sent overnight to the patient's home for self-collection. The patient performs a simple nasal wipe using a foam swab and places the kit in a prepaid FedEx envelope for routing to LabCorp.

Results are delivered directly to the provider's electronic medical record (EMR), making the process seamless for the ordering provider and convenient for patients. The result gives providers the information needed to make the appropriate clinical decision.

### LabCorp At Home Benefits

#### Health Care Providers

- Trusted testing using an FDA Emergency Use Authorized RT-PCR swab collection
- Seamless order through existing LabCorp account via interfaced EMR or LabCorp Link
- Turnkey delivery of specimen collection kit to patients at their home
- Simple collection and shipment of patient sample

#### For Patients

- Easy-to-follow collection and return shipping instructions
- Non-invasive nasal swab (short swab) collection
- No need to leave home for testing
- Results delivered in LabCorp Patient app (when registered)

**Kit includes:**  
Patient instructions, gentle foam swabs, a collection tube, biohazard bag and shipping envelope.



This home collection kit has not been FDA cleared or approved. This home collection kit has been authorized by the FDA under an EUA. This home collection kit has been authorized only for the home collection and maintenance of nasal swab specimens as an aid in detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. This home collection kit in combination with the authorized test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.



[www.LabCorp.com](http://www.LabCorp.com)

To set up LabCorp At Home ordering for your LabCorp account, please contact your local account representative.