FAQ: COVID-19 Home Test Kits

Frequently Asked	Kroger Health	Optum
Questions	COVID-19 Test Home Collection	At-home COVID-19 Test Kit
Can any employer order these home testing kits?	Kit Yes, any employer can order as there are no restrictions on employer eligibility.	Yes, however there is a 500-kit minimum order. It should be noted that for multi-site customers this program is not available for consumers residing in NY, NJ, and RI (as of 8/12/20).
When will home kits arrive after an employer order has been placed?	When a test is needed, the individual test kit is delivered directly to the employees home next day via overnight delivery.	Participant must first complete the registration process online. Test kits are then delivered to the individual's home.
Are the return envelopes pre-paid and pre-addressed for overnight delivery to a lab for processing?	Yes. All shipping logistics are pre-paid and pre-addressed for overnight delivery back to the lab for processing.	Yes. The self-collected participant samples are returned to lab with the overnight shipper provided with each test kit.
Can the return envelopes be sent by USPS, or is it required to be sent by Fed-Ex or UPS?	 Currently, our arrangement to ensure overnight delivery is via FedEx. Patients will receive a link listing the nearest drop box and location options. The drop box allows for zero contact delivery. 	Kits are delivered to participants via USPS. The pre-paid return label / envelope is sent back to the lab via United Parcel Service (UPS).
Which lab/labs serve as the sample processing partner?	Gravity Diagnostics in Covington, KY is the laboratory partner.	N/A
What is the processing and turnaround time to receive test results?	Processing takes 24-48 hours once the specimen has reached the lab. A turnaround time of 3-4 days can be expected depending on the diligence of patients/employers in taking and returning test kits in a timely manner.	Processing takes 24-48 hours once the sample has reached the lab. Turnaround time for results is 3-5 days from order date, including overnight shipping.



Have the processing labs been experiencing any delays in turnaround time?	There have not been any delays to date. Currently, 80% of samples are processed within the first 24 hours of the turnaround time frame, and laboratory capacity is being actively expanded. Future issues with turnaround time are not anticipated.	N/A
How are test results provided?	Overall test results are delivered to the employer via email. Individual Negative test results are provided to the employee via email. Individual Positive results are provided to the employee via phone call from a licensed clinician.	 Results are provided online on HIPAA-compliant digital platform. For those with positive results: Consult with a board-certified physician regarding next steps is provided. Positive results are automatically reported to the appropriate mandated local and federal reporting agencies.
How are these home test kits priced?	1 – 10,000 tests: \$119/test 10,001 – 50,000 tests: \$109/test 50,001+ tests: \$99/test	 Total cost for each test including lab work is \$119.90: \$59.00 plus a \$10.90 admin fee billed upon shipment of kit \$50.00 lab processing fee billed when a kit is returned for processing
Are the costs of these kits covered by commercial insurers in Indiana such as Anthem and UnitedHealthcare?	Currently the tests are available via B2B contract only and are not covered by or billable to insurance. The \$119/test cost would be invoiced to the employer monthly based on test utilization.	N/A



What if an employer	The EDA ELLA encompasses the entire	
	The FDA EUA encompasses the entire	
wants to order 1000	testing process (not just the kit) so we	_
tests and just pass	are not currently able to bulk ships tests	N/A
them out to	to be passed out by an employer. To	
employees, versus	mitigate employer obstacles, we can	
direct mailing to	assign vouchers for employers to hand	
employees? Will test	out to employees so they can be sent a	
kits be directly sent to	test immediately upon receipt of a	
employers, schools,	voucher.	
and people who want		
a test kit at their home		
to have a test just in		
case?		
Who can I contact for	Jeremy Richardson	Paula Conner
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Abbott's new rapid test:

Abbott BinaxNOW COVID-19 Ag Card

Emergency Use Authorization	The FDA issued an EUA for on August 26 th for the use of BinaxNOW COVID-19 Ag Card in the qualitative detection of the nucleocapsid protein antigen from SARS-CoV-2 in direct nasal swabs from people suspected of COVID-19 by their healthcare provider within the first seven days of symptom onset.
What is the cost of this test?	It is \$5 per test.
What are the capabilities of this test?	This new rapid test has shown a high sensitivity of 97.1% and specificity of 98.5% in clinical studies according to Abbott.*
How is it used?	A healthcare professional takes a nasal swab sample from the patient with a prescription. The swab is then inserted into the test card.



Does this test require the sample to be sent to a lab?	 No. The test card displays results in a similar manner to a home pregnancy test. If the patient is negative, the display will show one band at the 'control' line. If the patient is positive, the display will show two bands at the 'control' and 'sample' line.
How long does this test take to display results?	This test provides results in 15 minutes.
How can patients get proof of their COVID-19 test results?	 Abbott offers an optional free mobile app, the NAVICA app. This allows patients to display their results, obtained through a healthcare provider, to access facilities or activities that require proof of testing. The NAVICA app is an optional tool that allows people to store, access, and display their results with organizations that accept the results. The app is supported by Apple and Android digital wallets.
Is the NAVICA app used for contact tracing?	 No. The app is not used for contact tracing, and only collects the patient's first name, last name, email address, phone number, zip code, date of birth, and test results. Regardless of whether patients use the app, healthcare providers in all settings will be required to report positive test results to the CDC and other public health authorities.
Where can I find out more about the BinaxNOW COVID- 19 Ag Card?	 You can read more about the procedures, result interpretation, and performance characteristics of the test from an FDA publication found here: BinaxNOW COVID-19 Ag CARD. You can also view the following videos about the test: Abbott video WGN News video

* Higher test **sensitivity** means fewer false negative results, which means less disease cases are missed. A higher **specificity** means the test is better able to designate an individual who does not have a disease as negative. A highly specific test means that there are few false positive results.