

Florida Department of Health Emergency Rule 64DER21-12 provides opportunities for public school students to reduce quarantine or isolation if they are asymptomatic and receive a negative diagnostic test for COVID-19.¹ If a public school student is using one of the opportunities, as described in Emergency Rule 64DER21-12, to obtain a diagnostic COVID-19 test result to shorten their isolation or quarantine period, the COVID-19 diagnostic test should have received Emergency Use Authorization (EUA) from, or full approval by, the U.S. Food and Drug Administration (FDA) and the COVID-19 test should be used in the same manner specified by FDA's authorization or approval. The table below lists current COVID-19 diagnostic tests that have received an EUA by the FDA and how they can (i.e., over the counter or at-home) and should (i.e., symptomatic and/or asymptomatic) be used.

COVID-19 Tests that have received EUA from the FDA				
Antigen Tests²	Symptomatic Use	Asymptomatic Use³	Over the Counter Tests	Prescription At-Home Tests
Ellume COVID-19 Home Test	X	X	X	
BinaxNOW COVID-19 Ag Card 2 Home Test	X	X	X	
QuickVue At-Home OTC COVID-19 Test	X	X	X	
BinaxNOW COVID-19 Antigen Self Test	X	X	X	
InteliSwab COVID-19 Rapid Test	X	X	X	
CareStart COVID-19 Antigen Home Test	X	X	X	
InteliSwab COVID-19 Rapid Test Rx	X	X		X
BinaxNOW COVID-19 Ag Card Home Test	X	X		X
QuickVue At-Home COVID-19 Test	X	X		X
Molecular Tests⁴				
Any PCR Test given by provider or sent to a commercial laboratory ⁵	X	X		
Lucira CHECK-IT COVID-19 Test Kit	X	X	X	
Cue COVID-19 Test for Home and Over the Counter (OTC) Use	X	X	X	
Lucira COVID-19 All-In-One Test Kit	X	X		X

¹ <https://www.flgov.com/wp-content/uploads/2021/08/8.6.21-DOH-Rule.pdf>

² <https://www.fda.gov/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/in-vitro-diagnostics-euas-antigen-diagnostic-tests-sars-cov-2>

³ These tests have been approved for "Screening" and can be used for asymptomatic testing; this is not to be confused with "Serial Screening."

⁴ <https://www.fda.gov/medical-devices/coronavirus-disease-2019-covid-19-emergency-use-authorizations-medical-devices/in-vitro-diagnostics-euas-molecular-diagnostic-tests-sars-cov-2>

⁵ Includes "Direct to Consumer" and "Home Collection" tests, as these tests are sent to a commercial laboratory.