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 Tests[2]** | **Symptomatic Use** | **Asymptomatic Use[3]** | **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 | |
| QuickVue At-Home OTC COVID-19 Test | X | | X | |
| BinaxNOW COVID-19 Antigen Self Test | X | | X | |
| InteliSwab COVID-19 Rapid Test | X | | X | |
| CareStart COVID-19 Antigen Home Test | X | | X | |
| InteliSwab COVID-19 Rapid Test Rx | X | | | X |
| BinaxNOW COVID-19 Ag Card Home Test | X | | | X |
| QuickVue At-Home COVID-19 Test | X | | | X |
| | | | | |
| **Molecular Tests[4]** | | | | |
| 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.