

Luminex Testing Options: SARS-CoV-2

SARS-CoV-2 Targeted Detection Assays

MODERATE COMPLEXITY: SAMPLE-TO-ANSWER

ARIES®

SARS-CoV-2 Assay (EUA)

- 50-10047
 - 24 tests/kit
 - Sample type: NPS in UTM*
 - Up to 12 samples per run
 - TAT: ~2 hours
 - LoD: 333 copies/mL
 - Ship: 4°C
 - Store: -15-25°C
 - Gene targets: ORF1ab and N

ARIES® Workflow



Respiratory Panels That Detect SARS-CoV-2

HIGH COMPLEXITY: HIGH-THROUGHPUT

NxTAG®

CoV Extended Panel (EUA)

- I054C0463
 - 96 tests/kit
 - Sample type: NPS in UTM*
 - Up to 96 samples per run
 - TAT: ~4 hours (including extraction)
 - LoD: 5,000 copies/mL
 - Ship: 2-30° C
 - Store:
 - Plates 2-8°C
 - MS2 -25-8°C
 - Foil seals 15-30°
 - Gene targets: ORF1ab, N, and E

NxTAG® Workflow (Post-Extraction)

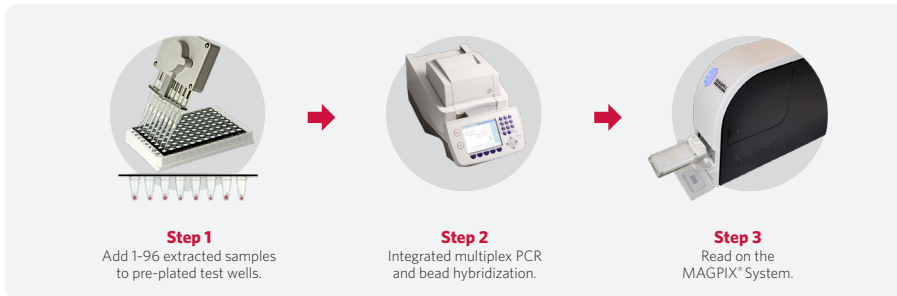


NxTAG®

RPP + SARS-CoV-2 (EUA)

- I056C0468
 - 19 viral pathogens, including SARS-CoV-2
 - 2 bacterial pathogens
 - 96 tests/kit
 - Sample type: Nasopharyngeal swabs
 - Up to 96 samples per run
 - TAT: ~4 hours (including extraction)
 - LoD (SARS-CoV-2): 500 copies/mL
 - Ship: 2-30° C
 - Store:
 - Plates 2-8°C
 - MS2 -25-8°C
 - Foil seals 15-30°
 - Gene targets (SARS-CoV-2): ORF1ab and M

NxTAG® Workflow (Post-Extraction)



*In UTM or equivalent. For the ARIES® Assay, the use of Amies transport medium or PBS is not recommended.

Gene targets for the detection of the SARS-CoV-2 virus vary by assay to ensure optimal multiplex product performance and may aid in Luminex's ability to support SARS-CoV-2 testing in the event of future viral genetic drift.

Luminex
complexity simplified.

For more information, please visit luminexcorp.com/covid19-testing-solutions

EUA - In Vitro Diagnostic Use Under Emergency Use Authorization. This test has not been FDA cleared or approved. This test has been authorized by the FDA under an EUA for use by authorized laboratories.
 The NxTAG® CoV Extended Panel and the ARIES® SARS-CoV-2 Assay have not been FDA cleared or approved. They have been authorized by the FDA under an EUA for use by authorized laboratories. This test has been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens. These tests are 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.
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