

# Power of Sensitivity

In a recent report published by the FDA, the Hologic assays are the **most sensitive** fully-automated, high-throughput molecular SARS-CoV-2 assays available.<sup>1</sup>



**Aptima® SARS-CoV-2 Assay™**

**PANTHER FUSION® SARS-CoV-2 Assay™**

The Panther® platform offers scalability, the potential to grow and full automation required to respond in these unprecedented times.

AVAILABLE TODAY



PANTHER®



ADD ON PANTHER FUSION®



ADD ON PANTHER PLUS



ADD ON PANTHER LINK



COMING SOON



ADD ON PANTHER TRAX®

Reference: 1. U.S. Food and Drug Administration. SARS-CoV-2 Reference Panel Comparative Data. Published September 15, 2020. Accessed September 21, 2020. <https://www.fda.gov/medical-devices/coronavirus-covid-19-and-medical-devices/sars-cov-2-reference-panel-comparative-data>

™ The Aptima and Panther Fusion SARS-CoV-2 assays.

\* These tests have not been FDA cleared or approved.

† These tests have been authorized by FDA under an EUA for use by authorized laboratories.

‡ These tests have been authorized only for the detection of nucleic acid from SARS-CoV-2, not for any other viruses or pathogens; and

§ 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.

¶ The Panther Fusion SARS-CoV-2 assay is available for EUA use in the United States, Australia, New Zealand, and Canada. EUA does not apply to the European Union. The Aptima SARS-CoV-2 assay is available for EUA use in the United States.

‡ In development and not for sale.

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**PANTHER®**