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SARS-CoV-2/COVID-19 Test Report

Patient:	P200969 / Glaks Diego Vega	**Sample ID:**	GLM- 226018
DOB:	06-01-2004	**Specimen:**	Nasal swab
Gender:	M	**Collection Date:**	08-28-2021 20:05:00
Job ID:	JOB-195497	**Received Date:**	08-28-2021 22:05:45
Physician:	Valerie Rusko	**Report date:**	08-29-2021
Practice:	AFC Urgent Care South Plainfield	**Correction:**	
Copy to:		**Travel Information:**	

Clinical History: Z11.52-Encounter for screening for COVID-19

Test Result

Not Detected

Interpretation

If **Positive/Detected**: indicative of active infection but do not rule out bacterial infection or other viral infections.

If **Negative/Not Detected**: does not preclude the COVID-19 infection and should not be used as the sole basis for treatment or other patient management decisions. Negative results must be combined with clinical observations, patient history, and epidemiological information.

Test Information

Testing was performed using the Thermo Fisher TaqMan 2019-nCoV Assay Kit v1, TaqMan 2019-nCoV Control Kit v1 Test, TaqPath COVID-19 Combo Kit and/ or Panther Fusion® SARS-CoV-2 Assay Kit by real-time (RT) PCR. Positive results do not rule out co-infection with organisms not included in the Thermo Fisher TaqMan 2019-nCoV Assay Kit v1 and TaqMan 2019-nCoV Control Kit v1, TaqPath COVID-19 Combo Kit and/ or Panther Fusions® SARS-CoV-2 Assay Kit by real-time (RT) PCR. The agent detected may not be the definite cause of this disease. This assay is for in vitro diagnostic use under the FDA Emergency Use Authorisation (EUA). This test is only authorized for the duration of time of the declaration and for the detection of SARS-CoV-2 RNA virus and/or diagnosis of COVID-19 infection under section 564 (b)(1) of the act, 21 U.S.C.360bbb•3(b)(1) unless the authorization is terminated or revoked sooner.

Laboratory Director: Anis Rangwala
Laboratory Certification: CLIA 31D1103468