Poseidon Diagnostics Corp

CLIA: 10D2218659

Patient Name:	Jasmine Hannah	**Physician:**	Dominique Thompson
Patient Gender:	Female	**Facility:**	Fast Test Now
Date of Birth:	04/11/1994	**Date Collected:**	2023-03-13T10:01:17
Patient Phone:	3528073778	**Received:**	2023-03-14T10:53:23
Patient Address:	610 NW 7th Ave Apt 80 Pompano Beach, FL 33060	**Released:**	2023-03-14T12:49:31
		Accession:	E19683
Requisition:	POSCV211062917	**Specimen Type:**	Nasopharyngeal Swab

Result:

COVID-19 RT-PCR Diagnostic Panel	
Test	**Result**
2019-nCoV RT-PCR	Negative

The Fastplex Triplex SARS-CoV-2 Detection Kit has received Emergency Use Authorization (EUA) from U.S. Food & Drug Administration (FDA. The Fastplex Triplex SARS-CoV-2 Detection Kit (RT-digital PCR) is only authorized for use in laboratories certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 U.S.C. §263a, that meet requirements to perform high complexity tests. The United States FDA has made this test available under an emergency access mechanism called an Emergency Use Authorization (EUA). The EUA is supported by the Secretary of Health and Human Service's (HHS's) declaration that circumstances exist to justify the emergency use of in vitro diagnostics (IVDs) for the detection and/or diagnosis of the virus that causes COVID-19.

A positive test result for COVID-19 indicates that RNA from SARS-CoV-2 was detected, and the patient is infected with the virus and presumed to be contagious. Laboratory test results should always be considered in the context of clinical observations and epidemiological data in making a final diagnosis and patient management decisions. Patient management should follow current CDC guidelines. The Fastplex Triplex SARS-CoV-2 Detection Kit has been designed to minimize the likelihood of false positive test results. A negative test result for this test means that SARS- CoV-2 RNA was not present in the specimen above the limit of detection. However, a negative result does not rule out COVID-19 and should not be used as the sole basis for treatment or patient management decisions. A negative result does not exclude the possibility of COVID-19. This document shall not be reproduced except in full, without the written approval of Poseidon Diagnostics

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