

**PATIENT INFORMATION:**

**Patient, Test**  
DOB: 5/13/2021  
Gender/Age: M/8 Days  
SS#: UN:918119519

**SPECIMEN INFORMATION:**

Accession #: **TS21-00084**  
Procedure Date: **5/13/2021**  
Date Received: 5/13/2021  
Reported On: 5/21/2021

**PHYSICIAN INFORMATION:**

**Test Physician**  
Test Practice  
300 Columbus Circle, Suite A, Edison, NJ 08837  
866.909.PATH, Fax:908-272-1478

**SARS-CoV-2 (COVID-19) (RT-PCR)**

**Result:** Not Detected

Reference Interval: Not Detected

**BILLING CODES:**

CPT: 87651  
ICD-10: B95.0

**DISCLAIMER:****SARS-CoV-2 (COVID-19) (RT-PCR)**

The LuminexNxTAG@CoV Expanded Panel assay has been authorized by the U.S. Food and Drug Administration (FDA) under Emergency Use Authorization (EUA). Results of this assay must be considered in conjunction with the clinical history, epidemiological data and other data available to the clinician evaluating the patient. False negative results may be obtained due to improperly collected, transported, or handled swab samples, due to the presence of sequence variants in the targets of the assay, amplification inhibitors in samples, or inadequate numbers of organism(s) for amplification. False positive results may occur due to cross-contamination by target organisms, their nucleic acids or amplified product, or from non-specific signals in the assay.

