Name: Glen Cooper | DOB: 6/3/1974 | MRN: 5E29F4A5-2CED-4C09-BA42-E07FE76A1411 | PCP: No PCP, NP

SARS-CoV-2 RNA, QL, RT PCR (COVID-19) - Details

SARS-COV-2 RNA, QL, RT PCR (COVID-19)

Results

Status: **Final result (Collected: 8/25/2020 11:17 AM)**

Hello,

You have a **NEGATIVE** Covid-19 Test Result.

Component	Value	Flag	Ref Range	Units	Status
SARS-CoV-2 RNA (COVID-19), QUALITATIVE NAAT	Not Detected		Not Detected		Final
Comment:					

Component	Value	Flag	Ref Range	Units	Status

A Not Detected (negative) test result for this test means that SARS-CoV-2 RNA was not present in the specimen above the limit of detection.
A negative result does not rule out the possibility of COVID-19 and should not be used as the sole basis for treatment or patient management decisions. If COVID-19 is still suspected, based on exposure history together with other clinical findings, re-testing should be considered in consultation with public health authorities. 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.

This patient specimen was tested using an FDA EUA pooling method.
Negative results from pooled testing should not be treated as definitive. If the patient's clinical signs and symptoms are inconsistent with a negative result or results are necessary for patient management, then the patient should be considered for individual testing. Specimens with low viral loads may not be detected in sample pools due to the decreased sensitivity of pooled testing.
Please review the "Fact Sheets" and FDA authorized labeling available for health care providers and patients using the following websites: https://www.questdiagnostics.com/home/Covid-19/HCP/QuestLDTP/fact-sheet; https://www.questdiagnostics.com/home/Covid-19/Patients/QuestLDTP/fact-sheet.html
This test has been authorized by the FDA under an Emergency Use Authorization (EUA) for use by authorized laboratories.

Due to the current public health emergency, Quest Diagnostics is receiving a high volume of samples from a wide variety of swabs and media for COVID-19 testing. In order to serve patients during this public health crisis, samples from appropriate clinical sources are being tested. Negative test results derived from specimens received in non-commercially manufactured viral collection and transport media, or in media and sample collection kits not yet authorized by FDA for COVID-19 testing should be cautiously evaluated and the patient potentially subjected to extra precautions such as additional clinical monitoring, including collection of an additional specimen.

Methodology: Nucleic Acid Amplification Test (NAAT) includes PCR or TMA

Additional information about COVID-19 can be found at the Quest Diagnostics website: www.QuestDiagnostics.com/Covid19