

Patient Report

Appointment record**Location****CPR number:** 3001007202**Full name:** V C**Date of issue:** 2021-06-21, 15:40**Nykøbing Falster, 4800****Fjordvej 15****General comments & Additional information****Clinical info:** POSITIVE**Ordered Items****SARS-CoV-2, PCR**

TESTS	RESULT	FLAG	UNITS	REFERENCE INTERVAL	LAB
SARS-CoV-2, PCR	POSITIVE	Abnormal		Not Detected	01

Patients who have a positive COVID-19 test result may now have treatment options. Treatment options are available for patients with mild to moderate symptoms and for hospitalized patients. Visit our website at <https://www.labcorp.com/COVID19> for resources and information. This nucleic acid amplification test was developed and its performance characteristics determined by LabCorp Laboratories. Nucleic acid amplification tests include RT-PCR and TMA. This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization (EUA). This test is only authorized for the duration of time the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection of SARS-CoV-2 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. When diagnostic testing is negative, the possibility of a false negative result should be considered in the context of a patient's recent exposures and the presence of clinical signs and symptoms consistent with COVID-19. An individual without symptoms of COVID-19 and who is not shedding SARS-CoV-2 virus would expect to have a negative (not detected) result in this assay.