CENTOGENE COVID 19 RT-PCR Testing Request Form

Please complete this form and provide patient's insurance card prior to testing.

Laboratory Personnel – FOR OFFICE USE ONLY		
Today's Date:	Location Name:	
Clinician Name:	Phone:	
Patient Information: COMPLETED BY PATIENT OR PARENT/GUARDIAN		
First Name:	Last Name:	Phone:
Address:		
City:	Zip Code:	County:
State:		
Date of Birth:	Age:	Sex: □ Male □ Female
Email (Print clearly):	Ethnicity:	Primary Language:
Does the patient live or work in a congregate setting (e.g., School, long term care facility, group home)?		
□ **YES** □ **NO**	Facility Name:	Occupation:

CLINICAL INFORMATION: COMPLETED BY PATIENT	
Date of symptom onset: □ None Any Recent Hospitalizations?	OPTIONAL: Does the patient have any underlying conditions? _____
Symptoms Observed: □ Fever □ Tiredness □ Dry Cough □ Body Ache □ Nasal Congestion □ Runny nose □ Loss of smell □ Diarrhea □ Loss of Appetite	□ None □ Unknown □ Diabetes □ Hypertension □ Cardiac Disease □ Other □ Immunocompromised □ Pregnant □ Chronic Lung Disease □ Chronic Liver Disease □ Chronic Kidney Disease □ Dialysis

LABORATORY TESTING – COMPLETED BY PATIENT	
Has the patient been tested for influenza? □ **YES** □ **NO** Result: □ **Positive** □ **Negative** Test Type: □ **Rapid** □ **PCR**	Any close contact with a laboratory confirmed COVID-19 case? □ **Yes** □**No** □**Unknown** If yes, type of contact? Household, Workplace, other

Has the patient been tested for any other viral respiratory illness?	□ **YES** □ **NO** **If yes, result:**
COVID 19 TESTING – COMPLETED BY PATIENT	
Has the patient been tested for COVID-19? □ **YES** □ **NO** Result: □ **Positive** □ **Negative** Test Type: □ **Rapid** □ **PCR**	

I hereby acknowledge and give full and complete consent for testing and request: RT-PCR Test □

I hereby acknowledge full and complete consent to and make request for a SARS-Cov2 qPCR. I am physically able to have this oral pharyngeal swab. I hereby request and authorize CENTOGENE to test this sample for me or the person named above for whom I am the legal guardian. I hereby release CENTOGENE. its principals, directors, members, employees, affiliates, suppliers, providers, subcontractors, successors, agents, their respective insurance carriers, and the location sponsoring this clinic/program, its principals, directors, employees, affiliates, successors, or agents from any and all liability, injury or damage whatsoever arising from, or in any way connected with, this SARS-CoV-2 qPCR or the administration of same including, but not limited to, acts of negligence. I authorize my medical information herein, including tests results, to be shared with my physician/insurance/employer. CENTOGENE will use and disclose your personal and health information to treat you, to receive payment for the care we provide, to public health agencies as required, and for our other health care operations which generally include those activities we perform to improve quality care. We have prepared a detailed **NOTICE OF PRIVACY AND CONFIDENTIALITY PRACTICES** to help you better understand our policies in regard to your personal health information. I acknowledge that I have received a copy of the Notice of Privacy and Confidentiality Practices. **Please provide a copy of this form to your physician and/or healthcare provider for your medical records.** This test is for informational purposes only and to be discussed with your health care professional. CENTOGENE, is not providing you with medical advice nor are they responsible for any outcome in your care or treatment. Please keep in mind that a positive result does not mean you are immune or cannot become re-infected. This test was developed, and its performance characteristics determined by CENTOGENE. This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization (EUA). This test has been validated in accordance with the FDA's Guidance Document (Policy for Diagnostics Testing in Laboratories Certified to Perform High Complexity Testing under CLIA prior to Emergency Use Authorization for Coronavirus Disease-2019 during the Public Health Emergency) issued on July 1st, 2020. FDA independent review of this validation is pending. 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.

_____ _____

Patient or Guardian Signature Date