COVID-19 IgG Antibody & RT-PCR Test Request Form

Please complete this form and provide a copy of insurance card and identification for at the time of collection.

Laboratory Personnel – FOR OFFICE USE ONLY		
Today's Date:	Location Name:	
Clinician Name:	Phone:	
Patient Information: COMPLETED BY PATIENT		
First Name:	Last Name:	Phone:
Address:		
City:	Zip Code:	County:
State:		
Date of Birth:	Age:	Sex: □ Male □ Female
Email:		
Additional Information **required** for testing:		
Does the patient live or work in a congregate setting (e.g., long-term care facility, shelter, group home, prison, jail)		
□ **YES** □ **NO**	Facility Name: Employee Occupation:	
Does the patient receive dialysis? □ **YES** □ **NO**		

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

LABORATORY TESTING – COMPLETED BY PATIENT	
Has the patient been tested for influenza? Result: □ **Positive** □ **Negative** Test Type: □ **Rapid** □ **PCR**	□ **YES** □ **NO**
Has the patient been tested for any other viral respiratory illness? Result:	□ **YES** □ **NO**

COVID 19 TESTING – COMPLETED BY PATIENT	
Has the patient been tested for COVID-19? Result: □ **Positive** □ **Negative** Test Type: □ **Rapid** □ **PCR**	□ **YES** □ **NO**

I hereby acknowledge full and complete consent to and make request for a SARS-Cov2 IgG Antibody test. I am physically able to have this blood draw and have never had an adverse reaction to any phlebotomy services. I hereby request and authorize PMH Laboratory, Inc. designated subcontractor who is an independent nurse/ healthcare staffing agency, not directly affiliated with PMH Laboratory, Inc., to collect this sample for me or the person named above for whom I am the legal guardian. I hereby release The PMH Laboratory, Inc. 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 IgG Antibody Test 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. The PMH Laboratory, Inc., 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. I agree to remain in the general area for at least 5 minutes after collection of samples. **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. The PMH Laboratory, Inc., 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 PMH Laboratory, Inc. 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 April 20, 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 SIGNATURE:_____DATE:_____

5862 Edinger Ave ◊ Huntington Beach ◊ CA ◊ 92649 ◊ (562) 592-2890 Office ◊ (909) 803-9790 Fax
CLIA 05D2137011 ◊ STATE CLF351210 ◊ COLA 028736 Rahil R. Khan, MD – Laboratory Director Rahil R. Khan, MD – Laboratory Director