

oup/Organization:				Location City:	
☐ Employee or Member of Group			□ Family	□ Re-Test	
	COVII	D_19 Datid	ent Test Request	Form	
Diagram and the the	<u> </u>		-		
Please complete this				d and identification at the time of collection.	
Consider Collection Date:	Patient Infor	mation: (•	tient or Guardian	
Specimen Collection Date:			Clinician Name	Clinician Name (if applicable):	
First Name:			Last Name:	Last Name:	
Address:					
City:		State:	Zip Code:	County:	
Email (Print Clearly):					
Phone Number:					
Date of Birth:			Age:	Sex: Non- Male Female	
				Binary	
	k in a congregate settir	ng (e.g., lo	ng-term care facilit	cy, shelter, group home, prison) 🗆 YES 🗆 NO	
Patient Clinical Information					
Date of symptom onset:					
Symptoms Observed:	□ None		Does the patier	nt have any underlying conditions?	
Fever		□ None			
□ Tiredness	□ Runny nose		□ Unknown	□ Pregnant	
□ Dry Cough	□ Loss of smell		□ Diabetes	□ Chronic Lung Disease	
□ Body Ache □ Diarrhea			, ,	☐ Hypertension ☐ Chronic Liver Disease	
□ Nasal Congestion □ Loss of Appetite LABORATORY TESTING − Completed by Patient □ Loss of Appetite			☐ Cardiac Disea☐ Other	□ Cardiac Disease □ Chronic Kidney Disease □ Other	
Has the patient received Influe	nza Vaccine?	□ No			
Has the patient received COVI	D-19 Vaccine?	□ No Vac	ccination Company N	Name:	
COVID 19 TESTING – Completed by Patient					
Has the patient been tested for COVID-19?			Result:	□ Positive □ Negative	
<u> </u>	reby acknowledge and	give full a	nd complete cons	ent for testing and request:	
		_	gG Antibody Test	□ SARS-Cov2 IgM Antibody Test	
			•	ngeal Swab (Nasal) Oropharyngeal Swab (Throat	
		•		physically able to have this nasal swab/blood draw and have never had a	
	, .			ubcontractor who is an independent nurse/ healthcare staffing agency, no I am the legal guardian. I hereby release PMH Laboratory, Inc. its principals	
		•		tive insurance carriers, and the location sponsoring this clinic/program, it	
	_			atsoever arising from, or in any way connected with, this SARS-CoV-2 qPCI	
nd/or 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 m shysician/insurance/employer/school/organization or group. PMH Laboratory, Inc., will use and disclose your personal and health information to treat you, to receive payment for the care w					
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					
etailed NOTICE OF PRIVACY AND CONFIDENTIALITY PRACTICES to help you better understand our policies regarding your personal health information. I acknowledge that I have received opy 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 you					
hysician and/or healthcare provider for your medical records. This test is for informational purposes only and to be discussed with your health care professional. PMH Laboratory, Inc., is required to the professional discussed with your health care professional. PMH Laboratory, Inc., is required to the professional discussed with your health care professional. PMH Laboratory, Inc., is required to the professional discussed with your health care professional. PMH Laboratory, Inc., is required to the professional discussed with your health care professional. PMH Laboratory, Inc., is required to the professional discussed with your health care professional. PMH Laboratory, Inc., is required to the professional discussed with your health care professional discussed with your heal					
roviding 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 cann ecome 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 bee					
uthorized 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 Laborato					
ertified 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. 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					
in vitro diagnostic tests for detection	•			on 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization i	
rminated or revoked sooner.					
atient/Guardian Signature:				DATE:	
ratient/Guardian Signature:					