

Bordetella Pertussis/Parapertussis

■ Flu A/B and RSV

# Respiratory Molecular Diagnostic Requisition $\square$ (Not to be billed

	All lesis on this form	nay require a signed ABN	to insurance)	
Employer Ir	nformation	Patient Information		
Company Name		Patient Name		
Company Name		<del>_</del>	(First Name, Last Name)	
Employer Name			State Zip	
Employer Phone		Cell #	ia automated system with telephone calls or	
			ur lab order. You may opt out of receiving	
X DR. Anthony Kearney/18	351340756	SS#		
Physician name/NPI		Date of Birth/Gender: □ Male □ Female		
Collection Information	tion	Drug Allergy(ies): □NK <u>A</u>		
Collection Date//		•		
	(Please attach copies of both	cards, front and back)		
PRIMARY  ☐ Medicare ☐ Medicaid	☐ Insurance ☐ Self Pa	ay □Ordering Physician	☐ Billing Information Attached	
			oup#	
SECONDARY				
☐ Medicare ☐ Medic	aid □ Insurance □	Self Pay □Ordering Phy	ysician (no ins. info needed)	
ICD-10 CODE(S)		verse side for references.)		
	·	•		
■ Respiratory Pathog	gen Panel plus COVID-1	19	SPECIMEN LABEL INSTRUCTIONS:  1. Complete required information above	
■ Respiratory Pathog	ien Panel		<ul><li>(highlighted areas and test request).</li><li>2. Remove labels and place one bar</li></ul>	
	jon r andr		coded label VERTICALLY on each specimen vial (not on the lid).	
Viral Targets	Dhina ima /Fatana ima	A dana a dim ta	3. Please discard any unused labels.	
Influenza A H1	Rhinovirus/Enterovirus	Adenovirus	The unique barcode identifies the patient with this requisition.	
Influenza A H1 Influenza A H3	Parainfluenza virus 1 Parainfluenza virus 2	Coronavirus HKU1 Coronavirus NL63	Please ensure the patient name,	
Influenza B	Parainfluenza virus 3	Coronavirus 229E	test request and specimen source is indicated so that both the label	
Respiratory Syncytial Virus		Coronavirus OC43	and registration match. Two patient	
Respiratory Syncytial Virus		Human Bocavirus	identifiers are required on each specimen submitted.	
Bacterial Targets			*ADVANCE BENEFICIARY NOTICE INSTRUCTIONS	
Chlamydophila pneumonia	e Mycoplasma pneumoniae	Legionella pneumoniae	All tests on this form are subject to	
	_		coverage limitations by Medicare and may require that an Advance	
COVID - 19 Only			Beneficiary Notice (ABN) be signed by the patient prior to obtaining the	
Group A Strep			specimen. When ordered tests are likely to be denied by Medicare.	

please complete a separate ABN with the patient's signature and date, submitting it with this requisition.

NOTE: For the convenience of the ordering physicians, the below ICD-10 codes are listed. Physicians are not required to use these codes but should report the diagnostic codes that best describes the reason for performing the test.

RPP DIAGNOSIS ICD-10 CODES			
MARK (✔)	ICD-10	DESCRIPTION	
	Z20.828	Exposure to COVID19	
	B97.29	Other Viral Pneumonia	
	J20.8	Acute Bronchitis	
	B97.29	Acute bronchitis due to other specified organisms	
	NOS – J22	Lower Respiratory Infection	
	B97.29	Unspecified acute lower respiratory infection	
	J80, B97.29	Acute Respiratory Distress Syndrome (ARDS)	
	J12.89	Viral Pneumonia	
	R05	Cough	
	R06.02	Shortness of Breath	
	R50.9	Fever	



#### INFORMED CONSENT TO SPECIMEN COLLECTION AND LAB TESTING

Please carefully read and sign the following Informed Consent:

I voluntarily agree to this testing for COVID-19:

- A. I authorize RCA Laboratory Services, LLC d/b/a GENETWORx or its subcontractor ("GENETWORx") to conduct collection and testing for COVID-19 through a nasal swab.
- B. I authorize my test results to be disclosed to the county, state, or to any other governmental entity as may be required by law.
- C. I acknowledge that a positive test result is an indication that I must self-isolate and/or wear a mask or face covering as directed in an effort to avoid infecting others.
- D. I understand that GENETWORx is not acting as my medical provider, this testing does not replace treatment by my medical provider, and I assume complete and full responsibility to take appropriate action with regards to my test results. I agree I will seek medical advice and treatment from my medical provider if I have questions or concerns, or if my condition worsens.
- E. I understand that, as with any medical test, there is the potential for a false positive or false negative COVID-19 test result. I, the undersigned, have been informed about the test purpose, procedures, possible benefits and risks, and I have received a copy of this Informed Consent. I have been given the opportunity to ask questions before I sign, and I have been told that I can ask additional questions at any time.

(Name) (Signature) (Date)

The signature of a parent or authorized guardian is required for individuals under age 18:

(Name) (Signature) (Date)



### **AUTHORIZATION FOR RELEASE OF INFORMATION**

I authorize RCA Laboratory Services, LLC ("GENETWORx") to release my COVID-19 test results to Keeneland Association Inc ("Provider")

### INFORMATION TO BE RELEASED

I understand that the information released will include any of the following Protected Health Information, as available:

• COVID-19 test results, including to detect the presence of COVID-19.

## CONDITIONS OF AUTHORIZATION

I understand that GENETWORx is providing this COVID-19 screening at Keeneland Association Inc request and for purposes of disclosing the results to ("Provider") and, therefore, if I refuse to sign this authorization, then I will not be eligible to receive the COVID-19 screening.

I have read the above and a Assocation, Inc:	nuthorize the release of my Prot	ected Health Information to	Keeneland
(Name)	(Signature)	(Date)	