

Johns Hopkins Genomics DNA Diagnostic Laboratory

General Requisition Form - Page 1

Ref	ferrer	Informat	tioı

Physician:	UPIN/NPI:				
Genetic Counselor:	Email:				
Institution:					
Address:					
Phone:	Fax:				
Additional reports to:					
Address:					
Phone:	Fax:				
*Mandatory Signature I have confirmed that the patient has been consented for the testing	Signature:				
ordered and that two matching identifiers are present on each page of this requisition.	Date:				
Patient Information (*two of these identifiers MUST also appear on the sample tube)					
Legal Name* (Last, First):					
Preferred Name (Last):	(First):				
Date of birth* (mm/dd/yy): Sex assigned	l at birth: Gender:				
Patient ID/MRN*:					
Patient Address:					
Billing Information (contact Billing Coordinator at 443-287-2486 prior to submitting)					
Billing contact:					
Phone: Fax:	Email:				
Referring Center Maryland Medicaid	Self-pay Patient Insurance Medicare				
Shipping Address: 1812 Ashland Ave, Sample Intake Rm 245, Baltimore, MD 21205					



Signature:

Johns Hopkins Genomics DNA Diagnostic Laboratory

General Requisition Form - Page 2

..120220

Clinical Information	Sample Information		
Patient Name:	Sample Type		
	Whole Blood	Cultured fibroblasts*	
DOB (mm/dd/yy):	Extracted DNA	Cleaned chorionic villi	
Indication:	Cultured amniocytes	Cultured chorionic villi	
maication.	Saliva	Cord blood	
	Other:		
	*Fibroblasts are the preferred sar	nple type for patients with	
ICD Codes:	hematologic malignancies The DNA Diagnostic Laboratory only accepts isolated or extracted nucleic acids for which the extraction or isolation is performed in a CLIA-certified laboratory or a laboratory meeting equivalent (or more stringent) requirements as determined by the College of American Pathologists (CAP) and/or the Centers for Medicare and Medicaid Services (CMS).		
Please attach detailed medical records. Ancestry: Caucasian Hispanic			
Northern European Central/South American	Collection Date:		
Western European Caribbean Eastern European Asian Middle Eastern Pacific Islander	Collection Site (JH Only):		
African American Native American Other	Has the patient had a transfusion?	Yes No	
Toot ordered.	If yes, was it PRBC transfusion	only? Yes No	
Test ordered:	Has the patient had a bone marrow	v transplant? Yes No	
Check here if testing is pregnancy-related	If yes to any of the above, please contact the laboratory prior to sending.		
Patient Informed Consent			
I grant permission for Johns Hopkins Genomics to perform the getesting may be dependent upon the clinical information provided turn-around-time. Risks and limitations of this test may include, b (non-paternity, consanguinity), uninformative negative results, un information may be used for quality control purposes, research, a only to the providers authorized on the test requisition. I understate	to the laboratory by my physician. The labut are not limited to, disclosure of unexp nexpected findings, and lab error. De-ide and shared in public healthcare database	aboratory cannot guarantee pected family information ntified clinical or genetic es. Results will be released	
Signature:	Date:		
Provider Alternate Consent			
I, the health care provider requesting the above testing, have ex and have obtained verbal consent or an alternate written consert the patient has consented for the testing ordered and that two materials.	nt (please attach) to order the test indic	ated. I have confirmed that	

Date:



Johns Hopkins Genomics DNA Diagnostic Laboratory

General Requisition Form - Page 3

v120220

Patient Information

syndrome (DiGeorge syndrome)

Phenotypes included: telomere shortening disorders

TeloZoom

Patient Name:	DOB:	
Test Directory		
CraniofacialZoom Phenotypes included: craniofacial disorders FancZoom Phenotypes included: Fanconi anemia Low Bone DensityZoom Phenotypes included: fragile bones, osteopenia MarrowZoom Phenotypes included: thrombooytopenia MDS and souto	Lung Disease Neonatal respiratory distress NGS panel Movement Disorders Huntington disease (HTT repeat sizing) Huntington disease like 2 (JPH3 repeat sizing) Targeted Variant(s)	
Phenotypes included: thrombocytopenia, MDS and acute leukemias, short telomere disorders, Diamond-Blackfan anemia/DBA-like hypoplastic anemias, Fanconi anemia, severe congenital neutropenia, sideroblastic anemia	Gene c p	
NeuromuscularZoom Subpanels available: Myopathy Charcot-Marie-Tooth Hereditary spastic paraplegia PulmZoom Subpanels available: Mucociliary disorders	Relationship to proband: *Please attach copy of previous report Prenatal Tests Maternal cell contamination study only Targeted variant(s) Gene	
Interstitial lung disease Pulmonary vascular disease RenalZoom Subpanels available:	Gene c p	
CAKUT, ciliopathies, and tubulointerstitial diseases Disorders of ion transport, nephrolithiasis and nephrocalcinosis Glomerular disease and complement genes	*Please attach copy of previous report Clinical Exome Sequencing Please see our website for the requisition form specific to exome sequencing	
SkeletalZoom Phenotypes included: short-ribpolydactyly, skeletalciliopathies Stickler22qZoom	Please ship samples to: 1812 Ashland Ave Sample Intake Rm 245	
Phenotypes included: Stickler syndrome, 22q deletion	Baltimore, MD 21205	

For sample specific requirements

please visit our website.