



Genetic Test Requisition Form

Internal Use Only - Accession #: _____

Informed consent **MUST** be signed by the Patient, Parent/Legal Guardian or Legal Next of Kin. Please contact PGMINQUIRY@cumc.columbia.edu for consent forms.

PATIENT INFORMATION:			ORDERING PHYSICIAN INFORMATION:		
LAST NAME:	FIRST NAME:	M.I.:	LAST NAME:	FIRST NAME:	M.I.:
DATE OF BIRTH:	MRN:	GENDER: <input type="checkbox"/> MALE <input type="checkbox"/> FEMALE	INSTITUTION:	NPI #:	
ADDRESS:			ADDRESS:		
CITY, STATE & ZIP:			CITY, STATE & ZIP:		
HOME PHONE:		WORK PHONE:	TELEPHONE NUMBER:		FAX NUMBER:
INSURANCE INFORMATION:			EMAIL ADDRESS:		
NAME OF POLICY HOLDER:		DATE OF BIRTH:	SIGNATURE:		
RELATIONSHIP TO PATIENT: <input type="checkbox"/> SELF <input type="checkbox"/> PARENT <input type="checkbox"/> SPOUSE <input type="checkbox"/> CHILD <input type="checkbox"/> OTHER (specify):			DATE:		
NAME & ADDRESS OF INSURANCE COMPANY:			GENETIC COUNSELOR NAME:		EMAIL ADDRESS:
POLICY NUMBER:	GROUP NUMBER:		INSTITUTIONAL BILLING: Do you have a PGM Billing Account?		
SECONDARY INSURANCE CARRIER:			<input type="checkbox"/> Yes P.O. # _____		
POLICY NUMBER:	GROUP NUMBER:		<input type="checkbox"/> No (Email PGMINQUIRY@cumc.columbia.edu to establish an account)		
MEDICARE PATIENTS ONLY: Check here to confirm that an Advance Beneficiary Notice (ABN) was signed by the Patient: <input type="checkbox"/>					
CREDIT CARD: I have provided my credit card information to the Pathology Billing Office (call 212-305-7399 to provide card information). <input type="checkbox"/>					
PREAUTHORIZATION: If health insurance preauthorization is required, check here if preauthorization is pending: <input type="checkbox"/>					

SAMPLE INFORMATION:
Amniotic Fluid Chorionic Villi Products of Conception Peripheral Blood Buccal Swab Muscle DNA (contact laboratory) Other (contact laboratory)
Collection Date: _____
ICD-10 CODES <i>List Below</i>

TEST ORDERED (FILL IN COMPLETELY):	
MOLECULAR TESTING	MITOCHONDRIAL DISEASES
<input type="checkbox"/> Huntington Disease (HTT) - CAG Repeat Expansion	<input type="checkbox"/> mtDNA Whole Genome Sequencing
<input type="checkbox"/> C9orf72 GGGGCC Repeat Expansion	<input type="checkbox"/> Southern Blot for Mitochondrial DNA Rearrangements
<input type="checkbox"/> Fragile X (FMR1) CGG Repeat Expansion	<input type="checkbox"/> Mitochondrial DNA Depletion
<input type="checkbox"/> Spinal Muscular Atrophy - SMN Copy Number	CHROMOSOMAL MICROARRAY
<input type="checkbox"/> Thrombophilia Risk Panel <input type="checkbox"/> APOE Genotyping	<input type="checkbox"/> SNP Oligonucleotide Microarray Analysis (SOMA)
• Factor V Leiden	Cytoscan HD* <i>Select Test Type</i>
• Prothrombin 20210G>A Mutations	<input type="checkbox"/> TARGETED <input type="checkbox"/> WHOLE GENOME *Maternal cell contamination (MCC) studies will be performed on all fetal samples
	Method subject to change at discretion of Pathologist

Collection Requirements
General Information: No special patient preparation is required. Specimens should be obtained and labeled as per standard hospital protocols including, but not limited to, labeling with two unique patient identifiers (i.e. Name and MRN or D.O.B.). Pathology specimens not from CUMC should have accompanying pathology reports to ensure identity.
Assay Specific Specimen Requirements: Huntington, C9orf72, Fragile X, Spinal Muscular Atrophy, Thrombophilia, APOE: Peripheral blood: (1) 3-5 mL Lavender-top EDTA tube; room temperature or refrigerated, or buccal swab (APOE) Mitochondrial Testing: Muscle tissue: flash frozen, 100-150 mg, approximately 0.1 cm in diameter and 0.5 cm in length, on Dry Ice OR Peripheral blood: (1) 3-5 mL Lavender-top EDTA; room temperature or refrigerated
SOMA: Peripheral blood; (1) 3-5 mL Lavender-top EDTA tube; room temperature or refrigerated. For other prenatal specimen requirements, please contact the laboratory. DNA: must be extracted in a CLIA-certified laboratory. Please contact the laboratory for assay-specific DNA requirements.