

## DIVISION OF MOLECULAR & GENOMIC PATHOLOGY

**MGP Laboratory Shipping Address:**  
Molecular & Genomic Pathology Laboratory  
3477 Euler Way, Room 7012  
Pittsburgh, PA 15213  
**Phone:** (412) 864-6140  
**FAX:** (412) 864-6151

<i>(To Be Completed by MGP Staff)</i>				
<b>Received Date</b>		<b>Case #</b>		
<b>Genetic Test Requisition</b>				
<b>PATIENT IDENTIFICATION</b>				<b>* Attach patient insurance card</b>
<b>Last Name</b>		<b>First Name</b>	<b>M.I.</b>	<b>SSN/MRN</b>
<b>Birthdate</b>	<b>Sex</b> <input type="checkbox"/> M <input type="checkbox"/> F	<b>Diagnosis</b>	<b>ICD-10 Code(s)</b>	<b>Surgical Path/Cytology #</b>
<b>CLIENT INFORMATION</b>				
<b>Requesting Institution/ Physician</b>				
<b>Requesting Physician Address</b>				
<b>Phone Number</b>			<b>Fax Number</b>	
<b>BILLING INFORMATION</b>				
<b>Person/Institution Responsible For Payment</b>				
<b>Billing Address</b>				
<b>Phone Number</b>			<b>Fax Number</b>	
<b>SPECIMEN INFORMATION</b>				
<b>Collection Date:</b>			<b>Collection Time:</b>	
<input type="checkbox"/> <b>Peripheral Blood</b>			<input type="checkbox"/> <b>Other (Please Call Lab):</b>	
<b>TESTS</b>				
Please attach a clinical summary, family history and pedigree. For family studies, one specimen from each family member to be tested should be submitted (please list any known mutation that exist in family, if applicable)				
<b>Hereditary Pancreatitis Panel: Sequencing</b> (Please select below)				
<input type="checkbox"/> <b>Whole Panel</b> <input type="checkbox"/> <b>PRSS1 Mutation Analysis</b> (A16V, K23R, N29I, and R122H variants) <input type="checkbox"/> <b>SPINK Mutation Analysis</b> (N34S variant)				
<b>Other Genetic Testing</b>				
<input type="checkbox"/> <b>Fragile X Syndrome:</b> PCR Sizing (FMR1 Gene; 5' UTR)				
<input type="checkbox"/> <b>Hemochromatosis:</b> Sequencing (HFE Gene: C282Y and H63D variants)				
<input type="checkbox"/> <b>Huntington Disease:</b> PCR Sizing (HTT Gene; exon 1): Informed consent required for pre symptomatic and symptomatic testing*				
<input type="checkbox"/> <b>vonHippel-Lindau Disease:</b> Whole Gene Sequencing (VHL Gene)				
<input type="checkbox"/> <b>Other</b> (Please Specify):				
<input type="checkbox"/> <b>Is there a known mutation in the family?</b> Indicate gene and mutation _____				

\*Additional Required Documents can be found at: <http://mgp.upmc.com>

Note: It is the responsibility of the patient's physician to obtain proper informed consent for genetic testing.

## ***Specimen Instructions and Shipping Instructions***

### **Peripheral blood**

- ◆ 2-5 ml of fresh peripheral blood collected in EDTA (purple top) tube or ACD (yellow top) tube.
- ◆ Blood should be refrigerated until shipment at 4°C.
- ◆ Shipment is at ambient temperature by overnight delivery in a properly labeled shipping container for biohazard substances.