

PATIENT INFORMATION:	Last Name:	First Name:	MI:	SSN/MRN#
MOLECULAR & GENOMIC PATHOLOGY (MGP)				
Molecular Oncology Test(s) Requested * Check All That Apply *				
<input type="checkbox"/> Comprehensive Hematopathology Molecular Analysis as per Pathologist (based on BM evaluation) - BMA only		<input type="checkbox"/> DNA/RNA Storage		
Lymphoid Neoplasms		Myeloid Neoplasms (Bone Marrow Aspirate preferred)		
<input type="checkbox"/> TP53 NGS (mutations and copy number alterations)		<input type="checkbox"/> Myeloid NGS Panel (54 genes, including FLT3, NPM1, TP53, JAK2, MPL, CALR, CEBPA) <i>If BMA, test will be held; reviewing Pathologist will order if appropriate.</i>		
<input type="checkbox"/> B-Cell Clonality Analysis (IgH and IgK gene rearrangement testing, PCR)		<input type="checkbox"/> FLT3 Analysis includes internal tandem duplication with allelic ratio and 835/836 codon analysis, PCR		
<input type="checkbox"/> T-Cell Clonality Analysis (Beta and Gamma chain gene rearrangement, PCR)				
Minimal Residual Disease		Single Gene Testing		
<input type="checkbox"/> t(9;22) BCR-ABL1 Quantitative RT-PCR Major (M) Breakpoint		<input type="checkbox"/> CALR (Calreticulin) Mutation Analysis exon 9, Sanger Sequencing		
<input type="checkbox"/> t(9;22) BCR-ABL1 Quantitative RT-PCR Minor (m) Breakpoint		<input type="checkbox"/> JAK2 V617F Mutation Testing myeloproliferative disorders, PCR		
<input type="checkbox"/> t(15;17) PML-RARA Translocation RT-PCR intron 3 breakpoint		<input type="checkbox"/> CEBPA Gene Sequencing for mutations, Sanger Sequencing, <i>*BMA preferred</i>		
<input type="checkbox"/> t(15;17) PML-RARA Translocation RT-PCR intron/exon 6 breakpoint				
<input type="checkbox"/> NPM1, Quantitative testing (Types A, B and D)		<input type="checkbox"/> Other (please specify):		

UPMC CYTOGENETICS LABORATORY	
Disease Phase	Test Requested (MUST be completed)
<input type="checkbox"/> New Diagnosis: <input type="checkbox"/> Relapse	<input type="checkbox"/> Comprehensive Hematopathology Cytogenetic Analysis as per Pathologist <i>(includes karyotype, FISH tests and/or panel, oncology microarray testing, diagnosis specific)</i> <input type="checkbox"/> Culture and Hold <i>(Relevant diagnostic testing will be ordered by the reviewing pathologist)</i> <input type="checkbox"/> Chromosome Analysis (Karyotype) with Confirmatory FISH Testing* <input type="checkbox"/> PML/RARA FISH <i>(reflex to RARA breakapart if necessary; STAT for new diagnosis only)</i> <input type="checkbox"/> BCR/ABL1 t(9;22) FISH
	<input type="checkbox"/> Integrated B-ALL Package: B-ALL FISH panel**, Onco Array, Karyotype* <input type="checkbox"/> Onco Array <input type="checkbox"/> T-ALL Package: T-ALL FISH Panel**, Karyotype
	<input type="checkbox"/> Integrated MDS Package: Onco Array, Karyotype* <input type="checkbox"/> Integrated AML Package: Onco Array, Karyotype*, FISH** (CBFB; RUNX1T1/RUNX1; KMT2A) rearrangements
	<input type="checkbox"/> Integrated CLL Package: Onco Array (CD19+ or whole PB/BM), Karyotype*, complementary FISH testing**
	<input type="checkbox"/> MM Package (includes plasma cell separation): Karyotype, FISH (IGH, IGH/MYC); MM Microarray, Includes Reflex FISH for IGH partners (CCND1, FGFR3, CCND3, MAF, MAFB)**
<input type="checkbox"/> Remission (Post Therapy)	<input type="checkbox"/> Culture and Hold <input type="checkbox"/> Karyotype <input type="checkbox"/> MM Follow-up CD138+ FISH <input type="checkbox"/> Follow-up FISH (specify): <i>Unless specified, ONLY follow-up FISH testing for previously detected abnormal clone(s) will be performed on remission specimens once an initial FISH testing has been performed.</i>
<input type="checkbox"/> Post- Transplant	<input type="checkbox"/> XX/XY donor FISH test
<input type="checkbox"/> Other Test	<input type="checkbox"/> FISH as per Pathologist (specify):
<input type="checkbox"/> Post-Bone Marrow Transplant: Days post transplant _____ Genetic Sex of Donor: <input type="checkbox"/> Male <input type="checkbox"/> Female	
<i>*Confirmatory FISH testing for clinically relevant regions will be performed on samples with abnormal karyotypes according to the laboratory best practice and diagnostic guidelines. Laboratory reserves the rights to determine a suitable methodology for testing including unstimulated and/or stimulated short and long-term cultures, FISH and/or microarray assay preferences, and FISH probe selection.</i> <i>**Visit our website for complete probe and panel listing as well as disease-specific testing approaches (https://geneticslab.upmc.com/)</i>	

Specimen Requirements (Each laboratory requires an individual specimen)		
Clinical Flow Cytometry Laboratory	Bone Marrow (2 ml Green top)	Ship at room temp by overnight delivery in a properly labeled shipping container for biohazard substances. If bone marrow, send one non-heparinized, unstained bone marrow slide if possible.
	Peripheral Blood (5 ml Purple top)	
Molecular & Genomics Pathology Laboratory	Bone Marrow (3 ml Purple top)	Blood should be refrigerated until shipment at 4°C. Ship at room temp by overnight delivery in a properly labeled shipping container for biohazard substances
	Peripheral Blood (3 ml Purple top)	
<input type="checkbox"/> (MUST complete Molecular & Genomic Pathology Laboratory section above)		
UPMC Cytogenetics Lab (Classical, FISH, Microarray)	Bone Marrow (3 ml Green top)	Specimen should be drawn in a heparinized syringe and placed in a Green top (sodium heparin) tube. Ship at room temp. An additional 1 ml Purple top (EDTA) tube is needed for microarray orders.
	Peripheral Blood (1 ml min. Green top)	
	<input type="checkbox"/> (MUST complete UPMC Cytogenetic Laboratory section above)	

