

HEMATOPATHOLOGY Patient Information Sheet

CLIENT INFORMATION	PATIENT INFORMATION		
	Last name	First name	M.I.
	Date of Birth	Age	Sex
Treating Physician (please print: first last):	Physician's contact number:		Medical Record #

SPECIMEN INFORMATION
<p style="text-align: center;">Collection Date mm ____ / dd ____ / yyyy ____ Collection Time ____ <input type="checkbox"/> AM <input type="checkbox"/> PM</p> <p><input type="checkbox"/> Peripheral Blood (attach CBC result): Green Top(s) ____ Purple Top(s) ____ Other ____ <input type="checkbox"/> Fresh Tissue(Biopsy or FNA) in <input type="checkbox"/> RPMI <input type="checkbox"/> Other ____ (Site) ____</p> <p><input type="checkbox"/> Bone Marrow (attach CBC result): Green Top(s) ____ Purple Top(s) ____ Core Biopsy ____ Clot ____ <input type="checkbox"/> Fluid: CSF ____ Pleural ____ Other: ____</p>

CLINICAL INFORMATION
<p>**** This section must be complete or report will be delayed ****</p>
<p>Known Diagnosis: _____</p> <p>Pertinent history: <input type="checkbox"/> Pancytopenia <input type="checkbox"/> Thrombocytopenia <input type="checkbox"/> Leukopenia <input type="checkbox"/> Anemia <input type="checkbox"/> Lymphocytosis <input type="checkbox"/> Abnormal lymphocytes <input type="checkbox"/> Monocytosis <input type="checkbox"/> Atypical cells/blasts <input type="checkbox"/> Eosinophilia <input type="checkbox"/> Monoclonal gammopathy <input type="checkbox"/> Plasmacytosis <input type="checkbox"/> Lymphadenopathy <input type="checkbox"/> Extranodal mass <input type="checkbox"/> Splenomegaly <input type="checkbox"/> Other _____</p> <p>Diagnosis under consideration (check all that apply): <input type="checkbox"/> non-Hodgkin Lymphoma <input type="checkbox"/> Hodgkin Lymphoma <input type="checkbox"/> Acute Leukemia <input type="checkbox"/> Chronic Lymphoproliferative Disorder <input type="checkbox"/> Myelodysplastic Disorder <input type="checkbox"/> Myeloproliferative Neoplasms <input type="checkbox"/> Multiple Myeloma <input type="checkbox"/> Other _____</p> <p>Status: <input type="checkbox"/> New diagnosis <input type="checkbox"/> Follow up <input type="checkbox"/> Minimal residual disease <input type="checkbox"/> Relapse <input type="checkbox"/> BM Transplant ICD-10 Code (Required) _____</p>

TEST MENU		
<p>Flow Cytometry - EDTA</p> <p><input type="checkbox"/> Global <input type="checkbox"/> Tech-Only</p> <p><input type="checkbox"/> Lymphoma/Lymphocytosis Panel FLWCY (For CLL, MM, and NHL studies)</p> <p><input type="checkbox"/> Leukemia/Lymphoma Comprehensive Panel FLWCY (For ALL, AML, CML and MDS studies)</p> <p>-----Reflex panels if indicated-----</p> <p><input type="checkbox"/> Plasma Cell <input type="checkbox"/> Hairy Cell <input type="checkbox"/> Acute Leukemia Intracellular Markers: (nTdT, cMPO, cCD3, cCD79a) <input type="checkbox"/> T-cell Receptor</p> <p><input type="checkbox"/> PNH PNHRW ARUP Laboratories</p> <hr/> <p>Cytogenetics (Chromosome Analysis) - Sodium Heparin ARUP Laboratories</p> <p><input type="checkbox"/> Peripheral Blood CHRLB <input type="checkbox"/> BM Aspirate CHABM</p>	<p>Fluorescence in situ Hybridization (FISH) - Sodium Heparin RQFSH</p> <p><input type="checkbox"/> *Process and hold <input type="checkbox"/> *Plasma Cell Enrichment - process and hold *Client Services must be called within 14 days of collection for panel selection</p> <p><input type="checkbox"/> Global <input type="checkbox"/> Tech-Only-pathologist for interp: _____</p> <p>Select a panel below: [FISH probes on panels may be ordered individually by checking the box beside test]</p> <div style="display: flex; justify-content: space-between;"> <div style="width: 48%;"> <p><input type="checkbox"/> ALL Panel (Adult)</p> <p><input type="checkbox"/> BCR/ABL t(9;22) <input type="checkbox"/> MLL Rearrangement (11q23)</p> <p><input type="checkbox"/> AML Panel</p> <p><input type="checkbox"/> Deletion 5q/Monosomy 5 <input type="checkbox"/> Deletion 7q/Monosomy 7 <input type="checkbox"/> Trisomy 8 <input type="checkbox"/> Deletion 20q <input type="checkbox"/> Inv(3) 3q26 <input type="checkbox"/> RUNX1/RUNX1T1 (AML/ETO) t(8;21) <input type="checkbox"/> PML/RARA t(15;17) (APL) <input type="checkbox"/> MYH11/CBFB; inv(16), t(16;16) <input type="checkbox"/> MLL Rearrangement (11q23)</p> <p><input type="checkbox"/> CLL/SLL Panel</p> <p><input type="checkbox"/> Deletion 11q (ATM) <input type="checkbox"/> Deletion 13q/Monosomy 13 <input type="checkbox"/> Deletion 17p (TP53) <input type="checkbox"/> Trisomy 12</p> <p><input type="checkbox"/> MDS Panel</p> <p><input type="checkbox"/> Deletion 5q/Monosomy 5 <input type="checkbox"/> Deletion 7q/Monosomy 7 <input type="checkbox"/> Trisomy 8 <input type="checkbox"/> Deletion 20q <input type="checkbox"/> MLL Rearrangement (11q23)</p> </div> <div style="width: 48%;"> <p><input type="checkbox"/> CML Panel:</p> <p><input type="checkbox"/> BCR/ABL t(9;22)</p> <p><input type="checkbox"/> Myeloma Panel</p> <p><input type="checkbox"/> Deletion 1p/1q Gain <input type="checkbox"/> Deletion 13q/Monosomy 13 <input type="checkbox"/> Trisomy 3, 5, 9 <input type="checkbox"/> Deletion 17p (TP53) <input type="checkbox"/> IGH Rearrangement (14q32) Reflex to <input type="checkbox"/> IGH/CCND1, t(11;14) <input type="checkbox"/> IGH/FGFR3, t(4;14) <input type="checkbox"/> IGH/MAF, t(14,16)</p> <p><input type="checkbox"/> NHL Panel</p> <p><input type="checkbox"/> ALK Rearrangement (2p23) <input type="checkbox"/> BCL6 Rearrangement (3q27) <input type="checkbox"/> MALT1 Rearrangement (18q21) <input type="checkbox"/> MYC Rearrangement (8q24) <input type="checkbox"/> IGH Rearrangement (14q32) <input type="checkbox"/> IGH/BCL2, t(14;18) <input type="checkbox"/> IGH/CCND1, t(11;14) <input type="checkbox"/> IGH/MYC, t(8;14).</p> <p><input type="checkbox"/> Eosinophilia Panel ARUP 2002378 PDGFR-α, (FIP1L1), PDGFR-β, FGFR1, and CBFB</p> </div> </div>	<p>Molecular Genetics - EDTA ARUP Laboratories www.aruplab.com ***UnitedHealthcare patients require prior authorization***</p> <p><input type="checkbox"/> BCR/ABL1 QL, Reflex to QT, major & minor BCRRX</p> <p><input type="checkbox"/> JAK2 V617F Mutation QL 0051245</p> <p><input type="checkbox"/> JAK2 V617F Mutation QL, Reflex to Exon 12 (Polycythemia Vera) JAK2R</p> <p><input type="checkbox"/> JAK2 V617F Mutation QL, Reflex CALR, MPL (ET, PMF) JACAR</p> <p><input type="checkbox"/> B-Cell Clonality by PCR (IgH/IgK) BCPCR</p> <p><input type="checkbox"/> T-Cell Clonality by PCR TCPCR</p> <p><input type="checkbox"/> FLT3 Mutation by PCR 2014683</p> <p><input type="checkbox"/> CEBPA Mutation 2004247</p> <p><input type="checkbox"/> NPM1 Mutation by PCR Qt 3000066</p> <p><input type="checkbox"/> WT1 Mutation by Sequencing 2005766</p> <p><input type="checkbox"/> IDH1 & IDH2 Mutation Analysis, exon 4 2006444</p> <p><input type="checkbox"/> Myeloid Malignancies Mutation Panel by NGS 2011117</p> <p><input type="checkbox"/> MYD88 L265P Mutation Detection by PCR, Quantitative 2009318</p> <p><input type="checkbox"/> Other _____</p>

LIS LABELS ONLY