

HEMATOPATHOLOGY**PERIPHERAL BLOOD AND/OR BONE MARROW REQUISITION**

Intermountain Laboratory Services
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**Patient Information** (affix label, patient stamp or complete info):

Patient Name: _____

Patient Date of Birth: _____

Medical Record Number or SSN: _____

If non-hospital patient, provide demographic sheet (or complete following info)

Patient's Address: _____ Phone #: _____

Insurance Name: _____ Policy #: _____

Insurance Policy holder's name: _____

Physician:		Collection Date:	
Coordinator (if applicable):		☐ STAT (Same day pathologist review) Physician pager/cell (required):	
Clinic/Facility:	Phone:	Fax:	Additional Fax:

Clinical History/Diagnosis:Recent chemotherapy? ☐ Yes ☐ No Type and day: _____Recent growth factors? ☐ Yes ☐ No Type and last dose: _____Post BMT? ☐ Yes ☐ No BMT Date: _____ Sex of donor: ☐ M ☐ F

- ☐ **Peripheral Blood Evaluation.** Pathologist review of PB only and ancillary tests as selected.
- ☐ **Bone Marrow Evaluation.** Pathologist review of PB and BM and ancillary tests as selected and/or as added by pathologist for diagnosis/prognosis.
- ☐ **Ancillary Testing Only.** Pathologist review previously completed.

Physician Signature: _____ Date: _____ Time: _____

Specimens submitted by collecting personnel:**Peripheral Blood:** Slides _____ NaHep _____ LavEDTA _____

Submit: 2 PB slides and CBC or LavEDTA for CBC/auto diff testing

Bone Marrow: Site: ☐ RPIC ☐ LPIC ☐ Patient ID and procedure site verified

Asp slides _____ Touch slides _____ Clot _____ Core _____ NaHep _____ LavEDTA _____ Other _____

Submit: Aspirate smear slides (4-6)	Touch imprint slides (2-4)
Aspirate clot (1 mL) in AZF or formalin	Core (≥1.5 cm) in AZF or formalin
Aspirates in NaHep (2 tubes, 3 mL each)	If dry tap, 2 extra cores in saline
Aspirate in LavEDTA (1 tube, 3 mL)	If acute leukemia suspected, extra aspirate LavEDTA (3 mL)

FLOW CYTOMETRY TESTING (NaHep [preferred] or LavEDTA, unless indicated):

- ☐ Hold for Leukemia/Lymphoma Phenotyping (FLOWSP), 1-2 mL PB or BM
- ☐ Leukemia/lymphoma Phenotyping (FLOWLL), 1-2 mL PB or BM
- ☐ PNH, High sensitivity, RBC and WBC - (PNHWBC) - PB Only
- ☐ PNH, High sensitivity, WBC - (PNHWBC) - PB Only
- ☐ T-Cell Clonality by Flow Cytometry Analysis of TCR V-Beta (ARUP 0093199) NaHep -PB Only
- ☐ ZAP-70 Analysis by Flow Cytometry (ARUP 0092392) LavEDTA, LiHep, or NaHep - PB Only

MICROBIOLOGY TESTING, BM Aspirate (1-5 mL in sterile tube)

- ☐ Blood Culture (BCX)
- ☐ Fungal Culture (FBCX)
- ☐ AFB Culture (AFBBLX)

Sending Laboratory Use Only

CL7001-F1 (4/2017)

Encounter/billing number (hospital patient): _____

Collecting/routing lab: ☐ AF ☐ CA ☐ DX ☐ IMC ☐ LD ☐ LO ☐ MK ☐ UV ☐ Other _____If sending to CL: For PB Eval only, order **PATHR** and selected tests. For BM Eval, order **BMTRK** and transport in BM kit.**CYTOGENETIC TESTING (5 mL PB or 3 mL BM, Na Hep):**

- ☐ Chromosome Analysis ☐ BM (**CRANBM**) ☐ PB (**CALB**) [☐ High Priority]
- ☐ Chromosome Analysis with Reflex to Genomic Microarray ☐ BM (**ARUP 2007130**) ☐ PB (**ARUP 2007131**)
- ☐ Cytogenetics Grow, Process and Hold ☐ BM (**CGPBM**) ☐ PB (**CGPPB**)
- ☐ Cytogenomic SNP Microarray-Oncology (**ARUP 2006325**)
- ☐ Multiple Myeloma Process and HOLD (**MMPHA**)

FISH Testing: ☐ Upfront ☐ Reflex (perform only if chromosome analysis is normal)

- ☐ Acute Myeloid Leukemia Panel (**FSHAML**)
- ☐ Chromosome, FISH -Interphase (**FISHI**) Specify probe(s): _____ [☐ STAT, avail for t(9;22)]
- ☐ Chronic Lymphocytic Leukemia Panel (**FISHCL**)
- ☐ Eosinophilia Panel (**ARUP 2002378**)
- ☐ Lymphoma (Aggressive) Panel (**AGGLYM**)
- ☐ MDS Panel (**MDSPAN**)
- ☐ Multiple Myeloma Panel (**MMF**)
- ☐ Myeloproliferative Disorders Panel (**ARUP 2002360**)
- ☐ PML RARa Translocation by FISH (**FISHP**) [☐ STAT]

MOLECULAR TESTING (5 mL PB or 3 mL BM, LavEDTA):

- AML:**
 - ☐ CEBF-MYH11 inv (16) Quant (**INV16**)
 - ☐ CEBPA Mutation Detection (**CEBPA**)
 - ☐ FLT3 Mutation Detection by PCR (**FLT3A**)
 - ☐ FLT3 Signal Ratio Mutation Detection by PCR (**FLT3SR**)
 - ☐ KIT Mutations in AML by Fragment Analysis and Sequencing (**KMAML**)
 - ☐ NPM1 Mutation by PCR and Fragment Analysis, Fluid (**NPM1A**)
 - ☐ PML/RARa Translocation, t(15;17)(**PMLAT**)
 - ☐ RUNX1-RUNX1T1(AML1-ETO) t(8;21) Detection, Quant (**RUNX1**)
- LPD:**
 - ☐ B-cell Clonality Screening (IgH and Igk) by PCR (**BCCHK**)
 - ☐ BRAF V600E Mutation in Hairy Cell Leukemia by Real-Time PCR, Quant (**ARUP 2007132**)
 - ☐ IGHV Mutation Analysis by Sequencing (**ARUP 0040227**)
 - ☐ Mantle Cell Lymphoma, bcl-1/JH (11;14), Real-time PCR, Cell-based [Quant] (**Quest #14991**)
 - ☐ MYD88 L265P Mutation Detection by PCR, Quant (**ARUP 2009318**)
 - ☐ T-cell Clonality Screening by PCR (**TCCPCR**)
 - ☐ T-cell Clonality Screening by NGS (**TCCSPC**)
- MPN:**
 - ☐ BCR ABL1 Major (p210) Quant (**BCRMAJ**)
 - ☐ BCR ABL1 Minor (p190) Quant (**BCRMIN**)
 - ☐ BCR-ABL1 Mutation Analysis by NGS (**ARUP 2008420**)
 - ☐ BCR ABL1, Qualitative Reflex to Quant (**BCRA**)
 - ☐ CALR (Calreticulin) Exon 9 Mutation Analysis by PCR (**CALR**)
 - ☐ JAK2 Gene, V617F Mutation, Qual (**JAK2**)
 - ☐ JAK2 Gene, V617F Mutation, Quant (**ARUP 0040168**) - PB Only
 - ☐ JAK2 Exon 12 Mutation Analysis by PCR (**ARUP 2002357**)
 - ☐ JAK2 (V617F) Mutation with Reflex to CALR Exon 9 with Reflex to MPL515 (**ARUP 2012084**)
 - ☐ KIT (D816V) Mutation [Mastocytosis] (**ARUP 0040137**)
 - ☐ MPL 515 Mutation, Quantitative (**ARUP 2005545**)
- Viral:**
 - ☐ Cytomegalovirus (CMV) (**CMVPCR**)
 - ☐ Epstein-Barr Virus (EBV), Qual (**EBVPCR**)
 - ☐ Parvovirus B 19, by PCR, Bone Marrow (**ARUP 0060028**)
- Other:**
 - ☐ Chimerism Post Transplant (**CHIMP**)
 - ☐ DNA Extraction and Storage (**DNAEXS**)
 - ☐ Myeloid Malignancies Mutation Panel by NGS (**ARUP 2011117**)
 - ☐ Myeloid Malignancies Somatic Mutation and Copy Number Analysis Panel (**ARUP 2012182**)
 - ☐ Ph-like B-ALL Multiplex RT-PCR (**ChildLab MOL74**)
 - ☐ RNA Extraction and Storage (**RNAEXS**)

OTHER TESTING:

- ☐ Specify lab, test name and test code: _____
- ☐ Study protocol: _____

Indication		Probe Target	Gene(s)/Unique Sequence
Acute Lymphocytic Leukemia Panel (ALL)	Adult ARUP 2002647	8q24	MYC
		t(9;22)(q34;q11.2)	BCR-ABL1
		11q23	KMT2A (MLL)
		14q32	IGH
		19p13	TCF3 (E2A)
	Pediatric ARUP 2002719	+4, +10	CEP4, CEP10
		t(9;22)(q34;q11.2)	BCR-ABL1
		11q23	KMT2A (MLL)
	t(12;21)(p13;q22)	ETV6-RUNX1 (TEL-AML1)	
Acute Myelogenous Leukemia Panel (AML): FSHAML		t(15;17)(q24;q21)	PML-RARA
		t(8;21)(q22;q22)	RUNX1T1-RUNX1 (ETO-AML1)
		inv(16)(p13.3q22)	CBFB
		11q23	KMT2A (MLL)
		inv(3) or t(3;3)	RPN1/MECOM (EVI1)
		del(5)(q31)	EGR1
		del(7)(q31)/-7	D7S486
Acute Myelogenous Leukemia with Myelodysplastic Syndrome or Therapy-Related AML Panel ARUP 2002653		del(5)(q31)	EGR1
		del(7)(q31)/-7	D7S486
		11q23	KMT2A (MLL)
Chronic Lymphocytic Leukemia Panel (CLL) FISHCL		del(11)(q22.3)	ATM
		+12	D12Z3
		del(13)(q14.3)	D13S319
		del(17)(p13.1)	TP53(p53)
Chronic Myelogenous Leukemia Panel (CML): FISHI		t(9;22)(q34;q11.2)	BCR-ABL1,ASS1
Eosinophilia Panel ARUP 2002378		4q12	PDGFRA-CHIC2-FIP1L1
		5q33.1	PDGFRB
		8p12	FGFR1
		inv(16)	CBFB
Lymphoma (Aggressive) Panel AGGLYM		3q27	BCL6
		8q24	MYC
		t(14;18)(q32;q21)	IGH-BCL2
Lymphoma FISHI	Burkitt	8q24	MYC
	Diffuse large cell	3q27	BCL6
	Follicular	t(14;18)(q32;q21)	IGH-BDL2
	IgH rearrangement	14q32	IGH
	Mantle cell	t(11;14)(q13;q32)	IGH-CCND1
	MALT	18q21	MALT1

Indication	Probe Target	Gene(s)/Unique Sequence
Multiple Myeloma Panel MMF	1q21	CKS1B
	+9	ASS1
	t(11;14)(q13;q32)	IGH-CCND1
	14q32	IGH rearrangement
	+15	PML
	del(17)(p13.1)	TP53(p53)
	NOTE: If IGH is positive, additional testing include:	
	t(4;14)(p16;q32)	IGH-FGFR3
Myelodysplastic Syndrome Panel (MPS) MDSPAN	t(14;16)(q32;q23.1)	IGH-MAF
	del(5)(q31)	EGR1
	del(7)(q31)/-7	D7S486
	+8	CEP8
Myeloproliferative Disorder Panel (MPD) ARUP 2002360	del(20)(q12)	D20S108
	4q12	PDGFRA-CHIC2-FIP1L1
	5q33.1	PDGFRB
	8p12	FGFR1
	T99;22)(q34;q11.2)	BCR-AB;1