

2119 E. 93rd / L15 Cleveland, OH 44106 216.444.5755 or 800.628.6816

HEMATOPATHOLOGY REQUISITION

<<FORM_ID>>

PATIENT INFORMATION (PLEASE PR	INT IN BLACK INK)						CLIENT INFORM	ATION
Last Name	Fir	st		MI				
Address	Bir	th Date (MM/I	DD/YYYY)	Sex 🗆	I M □ F			
City		#	International Patie					
			iliterriational Fatte	:IIL! LJ 1				
State	Zip Ho	me Phone						
Hospital/Physician Office Patient ID #	Ac	cession #				ORDERING	PHYSICIAN CONTACT	
MEDICAL NECESSITY NOTICE: When ordering to law to order tests) should only order tests that						Physician Name)	
INSURANCE BILLING INFORMATI	ON (PLEASE ATTACH CARD (OR PRINT IN BI	LACK INK)			Physician NPI#	:	
BILL TO: ☐ Client/Institution ☐ N	Nedicare 🗆 Insurance (Cor	nplete insuran	ce information below) \Box] Patient		Physician Phon	9	
PATIENT STATUS: □ Inpatient □	Outpatient 🗆 Non-Hospit	al Patient H	ospital discharge date: MM	/	/ _{YYY}			
WORKERS COMP: ☐ Yes ☐ No	DOI:					Physician Email		
PRIMARY: ☐ Medicare ☐ Medicaid (Ohio only) 🗆 Other Ins		Self	☐ Spouse	e 🗆 Child		to phone number: () _	
Subscriber Last Name	Fir	st		MI		☐ Fax report t	0: ()	
Beneficiary / Member #	Gr	oup #					NFORMATION number of tubes, vials, slides, tissu	e blocks provided.
Claims Address	Cit	у	State		Zip	(MM/DD/YYYY)		
SECONDARY: □ No □ Yes (if yes,	please attach) ABN:	☐ Yes	□ No				Specimer ow Biopsy*: Core C	n ID (#)s: lot
DIAGNOSIS CODE (REQUIRED) ICD-10 C	odes 1	2	3.	·		*Must Provide	e CBC/WBC Differential and Pathology Re	port
CLINICAL INFORMATION						□ Peripheral I	w Aspirate:Green top(s) Pur Blood: Green top(s) Pur	
						☐ Smears:	Air dried Fixed	Stained (type of stain)
						☐ Fluids:	CSF Pleural	
						☐ Fresh Tissu		
						☐ Paraffin blo Decalcified specia	cks: Tissue block(s) C nens will be rejected.	Cell block(s)
						☐ Slides:	Unstained Stain	ed
COMPREHENSIVE SERVICES The as needed to assist in evaluation. REQUIRE green top tubes and one 4 ml lavender top f	D: Copy of most recent WBC/C			Includ	des clinical his	tory review, morph	rrow Diagnostic Analysis (Hema ology/microscopy, flow cytometry, c May include FISH and/or molecular t	ytogenetic analysis & summary
INDIVIDUAL DIAGNOSTIC TESTS		1	MOLECULAR TESTIN	IG				
FLOW CYTOMETRY	FISH for Acute Myeloid Leuke	mia	CC-SIGN® NEXT-GENERAL		ML/MDS	n.c	MYD88 L265P Mutation Analysis	JAK2 V617F Mutation Detection
☐ Leukemia/Lymphoma Panel <i>RLLLIP</i> ☐ Lymphoma Panel for Tissue/	☐ Blood <i>FAMLPN</i> ☐ Bone Marrow <i>FAMLPM</i>		SEQUENCING PANELS PLEASE SELECT ONLY ONE	L	FLT3 ITD/D8 Mutation F3		☐ Blood <i>MYD88</i> ☐ Bone Marrow <i>MYD88M</i>	☐ Blood <i>JAK2</i> ☐ Bone Marrow <i>JAK2M</i>
Fluid RLLYMP	FISH for B Lymphoblastic Lei	ıkemia	NGS PANEL* Acute Leukemia NGS Pane		LL ⊐ IGVH Seguen	eing ICI/H	T-Cell Clonality Using Biomed-2	KIT D816V Mutation Detection
PNH, High Sensitivity, FLAER, Peripheral Blood Only PNHPNL	☐ Blood <i>FSHBLL</i> ☐ Bone Marrow <i>FSBLLM</i>		63-gene (DNA) and 107-ge (RNA)	ene _	IEMOGLOBINO	· ·	PCR Primers ☐ Blood <i>TCBMD</i>	☐ Blood <i>K816PB</i> ☐ Bone Marrow <i>K816BM</i>
CHROMOSOME ANALYSIS	FISH for Chronic Lymphocytic	Laukamia	☐ Marrow HDMNGS		☐ Alpha Thalas	semia Gene	☐ Bone Marrow <i>TCBMDM</i>	BCR/ABL1
Chromosome Analysis, Bone Marrow	☐ Blood <i>CLLFSH</i>	Louncilla	☐ Peripheral Blood <i>HDP</i>		Deletions AT	HALS	□ Other <i>TCBMDO</i>	BCR/ABL1 p210 and p190
☐ Bone Marrow <i>CHRBMH</i>	☐ Bone Marrow <i>CLLFSM</i>		Other HDONGS	R	-Cell Clonality	Using	T-Cell Receptor Beta Biomed-2 PCR	Diagnostic PCR ☐ Blood BCRPB1
☐ Reflex to AML FISH CHRAML☐ Reflex to MDS FISH CHRMDS	FISH for Myeloproliferative N ☐ Blood <i>MPNFSH</i>	eoplasms	Chronic Myeloid NGS Pane 63-gene (DNA)		IOMED-2 PCR		☐ Blood <i>TCRB</i>	☐ Bone Marrow BCRBM1
Reflex to Microarray BMCHF	☐ Bone Marrow MPNFSM		☐ Marrow MYNGSM		□ Blood <i>BCBN</i> □ Bone Marrov		☐ Bone Marrow <i>TCRBM</i> ☐ Other <i>TCRBO</i>	<i>BCR/ABL1</i> p210
Chromosome Analysis	☐ FISH for Myeloma Panel	FSHPCM	□ Peripheral Blood MYN□ Other MYNGSO		□ Other BCBN		TCR-G (PCR)	Quantitative PCR
☐ Leukemic Blood <i>CHRBLL</i>	IgH/CCND1		Hematologic Neoplasm Fus		mmunoglobulin Ising Biomed-2		☐ Blood <i>TGAMMA</i>	☐ Blood <i>P210PB</i> ☐ Bone Marrow <i>P210BM</i>
☐ Solid Tumor CHRSOL☐ Neoplastic Tissue CHRNPT	☐ Blood <i>FSHMCL</i> ☐ Bone Marrow <i>FSMCLM</i>		NGS Panel, 107-gene (RNA	A)** [□ Blood <i>IGHP</i>	CR	☐ Bone Marrow <i>TCRGM</i> ☐ Other <i>TCRGO</i>	BCR/ABL1 p190
·	PML/RARA		☐ Marrow <i>HFMNGS</i>☐ Peripheral Blood <i>HFP</i>		☐ Bone Marrov nmunoglobulin		MYELOPROLIFERATIVE	Quantitative PCR
FLUORESCENCE <i>IN SITU</i> Hybridization (FISH)	☐ Blood APLFSH		□ Other <i>HFONGS</i>	u	sing Biomed-2	PCR Primers	NEOPLASMS	☐ Blood <i>P190PB</i> ☐ Bone Marrow <i>P190BM</i>
(SEE BACK FOR OTHER FISH PROBES)	☐ Bone Marrow <i>APLFBM</i>				□ Blood <i>IGKP</i> 0 □ Bone Marrov		MPN NGS Panel, 3-Gene (DNA)* – includes <i>JAK2, CALR, MPL</i>	☐ <i>BCR/ABL1</i> Kinase Domain
FISH for Aggressive B-Cell Lymphoma ☐ Blood <i>FABCFP</i>	Other		*NGS panels overlap; please				☐ Marrow <i>MPNM</i>	Mutation Analysis <i>KINASE</i>
☐ Bone Marrow <i>FABCBM</i>	□ Other		at the top of Page 2 for inte **For selected indications on	nded use.			☐ Peripheral Blood <i>MPNP</i>	

NEXT-GENERATION SEQUENCING (NGS) PANELS & INTENDED USE

Please order only one NGS panel at a time. For a full list of interrogated genes, please refer to the Test Directory on clevelandcliniclabs.com.

Panel Name	Order Code, (Specimen Type)	Intended Use	Gene Number (Variants Type)	Variants Detected
Acute Leukemia	HDMNGS (M) HDPNGS (PB) HDONGS (0)	Known or suspected Acute Leukemias (e.g. Acute Lymphoblastic Leukemia — ALL, Acute Myeloid Leukemia — AML, Acute Promyelocytic Leukemia — APL)	63 (DNA) 107 (RNA)	DNA: SNVs, Insertions, Deletions RNA: Gene fusions, Structural Variants
SUBPANELS				
Chronic Myeloid	MYNGSM (M) MYNGSP (PB) MYNGSO (0)	Known or suspected Chronic Myeloid Neoplasms (e.g. MPNs, MDS)	56 (DNA)	DNA: SNVs, Insertions, Deletions
MPN	MPNM (M) MPNP (PB)	Known or suspected Myeloproliferative Neoplasms (MPNs) — includes <i>JAK2, CALR, MPL</i>	3 (DNA)	DNA: SNVs, Insertions, Deletions
Hematologic Neoplasm Fusion	HFMNGS (M) HFPNGS (PB) HFONGS (O)	Fusion-Only Targeted Analysis (e.g. myeloid and lymphoid neoplasms with eosinophilia and abnormalities of PDGFRA, PDGFRB or FGFR1; BCR-ABL1-negative CML); or, ALLs with insufficient specimen for combined DNA & RNA NGS testing	107 (RNA)	RNA: Gene fusions, Structural Variants

(M) = Marrow

 $(PB) = Peripheral\ Blood$

(0) = Other: FFPE, Clot sections

 ${\sf SNV} = {\sf Single \ Nucleotide \ Variant(s)}$

FLUORESCENCE IN SITU HYBRIDIZATION (FISH) PANELS

Panel Name	Probes	Blood Order Code	Bone Marrow Order Code	Tissue Order Code
FISH for Acute Myeloid Leukemia	t(15;17), t(8;21), inv(16), KMT2A (MLL)	FAMLPN	FAMLPM	
FISH for Aggressive B-Cell Lymphoma	BCL2 (18q21), BCL6 (3q27), c-MYC (8q24) and (8;14)(q24;q32)	FABCFP	FABCBM	FABCEL
Anaplastic Large Cell Lymphoma Panel	TP63, DUSP22/IRF4			DUIRFH
FISH for B Lymphoblastic Leukemia (B-ALL)	t(9;22), KMT2A (MLL), t(12;21), 4/10 cen, CRLF2, CRLF2/IGH	FSHBLL	FSBLLM	
FISH for Chronic Lymphocytic Leukemia	17p13 (TP53), 11q22 (ATM), 12 cen, 13q34 (D13S319, LAMP1), 11;14 (IgH/CCND1)	CLLFSH	CLLFSM	
FISH for Myelodysplastic Syndrome	5q31 (EGR1), 7q31 (D7S486), 8 cen, 20q12 (D20S108)	FSHMDS	FSMDSM	
FISH for Myeloma*	13q, 17p, 1p/1q, CEP9/CEP15, IgH, IgH/CCND1 (11;14) If an IgH translocation is present that is not IgH/CCND1, then will reflex to IgH/MMSET (4;14) and IgH/MAF (14;16).		FSHPCM	
FISH for Myeloproliferative Neoplasms	t(9;22), 4q12 (PDGFRA), 5q33 (PDGFRB), 8p12 (FGFR1)	MPNFSH	MPNFSM	

^{*}Plasma cells only

FISH PROBES (For a full list of FISH probes, please refer to the Test Directory on clevelandcliniclabs.com)

Test Name	Gene(s)	Location	Blood Order Code	Bone Marrow Order Code	Tissue Order Code
FISH for BCL2	BCL2	18q21	BCL2FH	BCL2FM	BCL2FT
FISH for BCL6	BCL6	3q27	BCL6FH	BCL6FM	BCL6FT
FISH for BCR/ABL1	BCR/ABL1	t(9;22)	BCRFSH	BCRFSM	
FISH for BIRC3/MALT1	BIRC3/MALT1 (API2/MALT1)	t(11;18)			T1118
FISH for CBFB/MYH11	CBFB	inv(16)	INV16F	INV16M	
FISH for CCND1	CCND1	11q13	CCND1F	CCND1M	CCND1T
FISH for CRLF2 Rearrangement	CRLF2	Xp22.33/Yp11.32	CRLF2B	CRLF2M	
FISH for IGH::CRLF2 Rearrangement	CRLF2, IGH	Xp22.33/Yp11.32 and 14q32.33	CRIGHB	CRIGHM	
FISH for del (5q)	EGR1	5q31	5QFSH	5QFSBM	
FISH for del (7q)	D7S486	7q31	FISH7Q	FSH7QM	
FISH for DUSP22/IRF4	DUSP22/IRF4	6p25.3			DUIRFH
FISH for ETV6/RUNX1	ETV6/RUNX1 (TEL/AML1)	t(12;21)	1221FH	1221FM	
FISH for FGFR1	FGFR1	8p12	FGFR1F	FGFR1M	
FISH for IGH/BCL2	IGH/BCL2	t(14;18)	FSHFCL	FSFCLM	T1418
FISH for IGH/CCND1	IGH/CCND1	t(11;14)	FSHMCL	FSMCLM	T1114
FISH for IGH/MYC/CEP8	IGH/MYC	t(8;14)	814FSH	814FSM	T814
FISH for MALT1	MALT1	18q21			MALT1
FISH for MLL	MLL	11q23	MLLFSH	MLLFBM	
FISH for MYC(8q24)	MYC	8q24	MYCFSH	MYCFSM	MYC
FISH for PDGFRA	PDGFRA	4q12	PDGFRA	PGFRAM	
FISH for PDGRRB	PDGRRB	5q33	PDGFRB	PDGFBM	
FISH for PML/RARA	PML/RARA	t(15;17)	APLFSH	APLFBM	
FISH for RARA	RARA	17q21	RARFSH	RARFSM	
FISH for 8;21 Translocation for AML	RUNX1/RUNX1T1 (AML1/ETO)	t(8;21)	AMLFSH	AMLFBM	
FISH for Trisomy 4, 10	Trisomy 4/10	4 cen, 10 cen	FHT410	FT410M	
FISH for Trisomy 8, del(20q)	Trisomy 8/D20S108	8 cen, 20q12	20Q8FH	20Q8BM	
FISH for TP63	TP63	3q28			TP63FH