

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

HEMATOPATHOLOGY REQUISITION

<<FORM ID>>

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PATIENT INFORMATION (PLE	ASE PRINT IN BLACK INK)			CLIEN	T INFORMATION
Last Name	F	irst M	11		
Address	E	sirth Date (MM/DD/YYYY) So	ex		
City	S	S # International Patient?	☐ Yes ☐ No		
State	Zip H	ome Phone			
Hospital/Physician Office Patient ID	# A	ccession #		ORDERING PHYSICIAN COI	NTACT
		rsement will be sought, physicians (or other indivice diagnosis or treatment of a patient, rather than fo		Physician Name	
INSURANCE BILLING INFOR				Physician NPI#	
		omplete insurance information below) D	atient	Physician Phone	
WORKERS COMP:		ital Patient Hospital discharge date:MM	DD YYYY	Physician Email	
PRIMARY: □ Medicare □ Me			Spouse Child	☐ Call results to phone number:	()
Subscriber Last Name	F	irst N	<u></u>	☐ Fax report to: () _	
Beneficiary / Member #	G	iroup #		SPECIMEN INFORMATION Please indicate number of tubes, v	ials, slides, tissue blocks provided.
Claims Address	0	ity State	Zip	Collection Date://_	Time:
SECONDARY: No Yes		. □ Yes □ No	,	Body Site: Bone Marrow Biopsy*: Core	Specimen ID (#)s:e Clot
DIAGNOSIS CODE (REQUIRED) 10	CD-10 Codes 1	2 3		*Must Provide CBC/WBC Differentia	al and Pathology Report
CLINICAL INFORMATION				□ Bone Marrow Aspirate:Green to □ Peripheral Blood: Green to	p(s) Purple top(s) Other Other
					Fixed Stained (type of stain)
					Pleural FNA Other
					or Lymph Node lock(s) Cell block(s)
				Decalcified specimens will be rejected.	dStained
COMPREHENSIVE SERVICE: as needed to assist in evaluation. Rt green top tubes and one 4 ml lavend	EQUIRED: Copy of most recent WBC/	ologist is authorized to add other testing CBC, peripheral blood smears, two 4 ml	Includes clinical hist	tory review, morphology/microscopy,	Analysis (Hematopathologist Directed) flow cytometry, cytogenetic analysis & summary nd/or molecular testing as medically necessary.
INDIVIDUAL DIAGNOSTIC T	ESTS	MOLECULAR TESTING			
FLOW CYTOMETRY Leukemia/Lymphoma Panel	FLUORESCENCE IN SITU Hybridization (FISH)	CC-SIGN® NEXT-GENERATION SEQUENCING PANELS	AML/MDS □ <i>CEBPA</i> Mu	utation <i>CEBPA</i>	☐ T-Cell Receptor Beta (TCRB) only <i>TCBMDM</i> ☐ T-Cell Receptor Gamma (TCRG) only <i>TCRGM</i>
RLLLIP	(SEE BACK FOR OTHER FISH PROBES) ☐ FISH for Aggressive B-Cell	PLEASE SELECT ONLY ONE NGS PANEL.*	□ <i>FLT3</i> ITD/I □ <i>NPM1</i> Mu	D835 Mutation <i>F3ITDM</i>	MYELOPROLIFERATIVE NEOPLASMS
☐ Lymphoma Panel for Tissue/ Fluid <i>RLLYMP</i>	Lymphoma <i>FABCFP</i>	Acute Leukemia NGS Panel, 63-gene (DNA) and 107-gene (RNA)	CLL	itation W W1	MPN NGS Panel, 3-Gene (DNA)* – includes <i>JAK2, CALR, MPL</i>
 PNH, High Sensitivity, FLAER, Peripheral Blood Only 	☐ FISH for AML Panel FAMLPN☐ FISH for B-ALL Panel FSHBLL	□ Marrow HDMNGS□ Peripheral Blood HDPNGS	☐ IGVH Sequ	iencing <i>IGVH</i>	☐ Marrow MPNM
PNHPNL	☐ FISH for CLL Panel <i>CLLFSH</i>	□ Other <i>HDONGS</i>	HEMOGLOBIN □ Alpha Thai	NOPATHY lassemia Gene Deletions ATHALS	☐ Peripheral Blood <i>MPNP</i>
CHROMOSOME ANALYSIS	☐ FISH for MDS Panel <i>FSHMDS</i>	Chronic Myeloid NGS Panel, 56-gene (DNA)	LYMPHOMA		☐ JAK2 V617F Mutation Detection JAK2
Chromosome Analysis, Bone Marrow	☐ FISH for MPN Panel MPNFSH☐ FISH for Myeloma Panel	☐ Marrow MYNGSM ☐ Peripheral Blood MYNGSP	☐ B-Cell Clo	nality (IGH and IGK Gene	☐ JAK2 Exon 12-16 Sequencing JAKNON☐ Calreticulin (CALR) Exon 9 Mutation
☐ Bone Marrow CHRBMH	FSHPCM	Other MYNGSO	☐ Immunogl	ement) <i>BCBMD</i> obulin Heavy Chain (IGH)	Analysis CALR
☐ Reflex to AML FISH CHRAML☐ Reflex to MDS FISH CHRMDS	☐ <i>BCR/ABL1</i> , t(9;22) <i>BCRFSH</i> ☐ <i>IGH/CCND1</i> , t(11;14)	Hematologic Neoplasm Fusion NGS Panel, 107-gene (RNA)**	only <i>IGHM</i> □ Immunogle	1 obulin Kappa Chain (IGK)	☐ <i>MPL</i> Mutation Analysis <i>MPL</i> ☐ <i>KIT</i> (D816V) Mutation <i>K816BM</i>
☐ Reflex to Microarray <i>BMCHF</i>	FSHMCL	☐ Marrow HFMNGS ☐ Peripheral Blood HFPNGS	only <i>IGKM</i>		BCR/ABL1
Cytogenetic Analysis Leukemic Blood CHRBLL	☐ PML/RARA, t(15;17) APLFSH	☐ Other <i>HFONGS</i>	Chronic LPD № □ Bone Mar	NGS Panel, 7-gene (DNA)*	☐ BCR/ABL1 p210, Quantitative P210PB
☐ Solid Tumor CHRSOL	☐ Other		□ Peripheral	I Blood LPPNGS	☐ BCR/ABL1 p190 RT-PCR, Quantitative P190PB
☐ Neoplastic Tissue <i>CHRNPT</i>		*NGS panels overlap; please order only <u>one</u> , Refer to the NGS Panel Table at the top of	☐ Other <i>LPC</i> ☐ MYD88 L2	0NGS 65P Mutation Detection MYD88	☐ BCR/ABL1 Kinase Domain Mutation Analysis KINASE
		Page 2 for intended use. **For selected indications only; refer to Page 2 for intended use.		nality (<i>TCRB</i> and <i>TCRG</i> Gene ement) <i>TCBMD</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
Chronic LPD	LPMNGS (M) LPPNGS (PB) LPONGS (O)	Known or suspected Chronic Lymphoproliferative Disorders (LPDs)	7 (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
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 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