

PATIENT INFORMATION (PLEASE PRINT IN BLACK INK)

Last Name First MI
Address Birth Date (MM/DD/YYYY) Sex M F
City SS # International Patient? Yes No
State Zip Home Phone
Hospital/Physician Office Patient ID # Accession #

MEDICAL NECESSITY NOTICE: When ordering tests for which Medicare reimbursement will be sought, physicians (or other individuals authorized by law to order tests) should only order tests that are medically necessary for the diagnosis or treatment of a patient, rather than for screening purposes.

INSURANCE BILLING INFORMATION (PLEASE ATTACH CARD OR PRINT IN BLACK INK)

BILL TO: Client/Institution Medicare Insurance (Complete insurance information below) Patient
PATIENT STATUS: Inpatient Outpatient Non-Hospital Patient Hospital discharge date: MM/DD/YYYY
WORKERS COMP: Yes No DOI:
PRIMARY: Medicare Medicaid (Ohio only) Other Ins. Self Spouse Child
Subscriber Last Name First MI
Beneficiary / Member # Group #
Claims Address City State Zip
SECONDARY: No Yes (if yes, please attach) ABN: Yes No

DIAGNOSIS CODE (REQUIRED) ICD-10 Codes 1. 2. 3.

CLINICAL INFORMATION

Blank lines for clinical information.

COMPREHENSIVE SERVICES The Cleveland Clinic Hematopathologist is authorized to add other testing as needed to assist in evaluation. REQUIRED: Copy of most recent WBC/CBC, peripheral blood smears, two 4 ml green top tubes and one 4 ml lavender top for bone marrow aspirate.

Comprehensive Primary Bone Marrow Diagnostic Analysis (Hematopathologist Directed) Includes clinical history review, morphology/microscopy, flow cytometry, cytogenetic analysis & summary report with correlation of all findings. May include FISH and/or molecular testing as medically necessary.

INDIVIDUAL DIAGNOSTIC TESTS

FLOW CYTOMETRY: Leukemia/Lymphoma Panel RLLLP, Lymphoma Panel for Tissue/Fluid RLLYMP, PNH, High Sensitivity, FLAER, Peripheral Blood Only PNHPNL
CHROMOSOME ANALYSIS: Chromosome Analysis, Bone Marrow, Bone Marrow CHRBMH, Reflex to AML FISH CHRAML, Reflex to MDS FISH CHRMD5, Reflex to Microarray BMCHF
FLUORESCENCE IN SITU HYBRIDIZATION (FISH): FISH for Acute Myeloid Leukemia, FISH for B Lymphoblastic Leukemia, FISH for Chronic Lymphocytic Leukemia, FISH for Myeloproliferative Neoplasms, FISH for Myeloma Panel FSHPCM, IgH/CCND1, PML/RARA

MOLECULAR TESTING

CC-SIGN® NEXT-GENERATION SEQUENCING PANELS: PLEASE SELECT ONLY ONE NGS PANEL*
Acute Leukemia NGS Panel, 63-gene (DNA) and 107-gene (RNA)
Chronic Myeloid NGS Panel, 63-gene (DNA)
Hematologic Neoplasm Fusion NGS Panel, 107-gene (RNA)**
*NGS panels overlap; please order only one. Refer to the NGS Panel Table at the top of Page 2 for intended use.
**For selected indications only; refer to Page 2 for intended use.

AML/MDS: FLT3 ITD/D835 Mutation F3ITDM
CLL: IG VH Sequencing IGVH
HEMOGLOBINOPATHY: Alpha Thalassemia Gene Deletions ATHALS
LYMPHOMA: B-Cell Clonality Using BIOMED-2 PCR Primers, Immunoglobulin Heavy Chain using Biomed-2 PCR Primers, Immunoglobulin Kappa Chain using Biomed-2 PCR Primers
MYELOPROLIFERATIVE NEOPLASMS: MPN NGS Panel, 3-Gene (DNA)* - includes JAK2, CALR, MPL
MYD88 L265P Mutation Analysis: Blood MYD88, Bone Marrow MYD88M
JAK2 V617F Mutation Detection: Blood JAK2, Bone Marrow JAK2M
KIT D816V Mutation Detection: Blood K816PB, Bone Marrow K816BM
BCR/ABL1: BCR/ABL1 p210 and p190 Diagnostic PCR, BCR/ABL1 p210 Quantitative PCR, BCR/ABL1 p190 Quantitative PCR, BCR/ABL1 Kinase Domain Mutation Analysis KINASE

CLIENT INFORMATION

ORDERING PHYSICIAN CONTACT

Physician Name
Physician NPI#
Physician Phone
Physician Email
Call results to phone number: ()
Fax report to: ()

SPECIMEN INFORMATION

PLEASE INDICATE NUMBER OF TUBES, VIALS, SLIDES, TISSUE BLOCKS PROVIDED.
Collection Date: / / Time:
Body Site: Specimen ID (#):
Transport Temperature: Ambient Refrigerated
Bone Marrow Biopsy*: Core Clot
Bone Marrow Aspirate: Green top(s) Purple top(s) Other
Peripheral Blood: Green top(s) Purple top(s) Other
Smears: Air dried Fixed Stained (type of stain)
Fluids: CSF Pleural FNA Other
Fresh Tissue: Tumor or Lymph Node
Paraffin blocks: Tissue block(s) Cell block(s)
Slides: Unstained Stained

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 (O)	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 (O)	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</i> , <i>CALR</i> , <i>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 <i>PDGFRA</i> , <i>PDGFRB</i> or <i>FGFR1</i> ; <i>BCR-ABL1</i> -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 (O) = Other: FFPE, Clot sections

SNV = 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) <i>If an IgH translocation is present that is not IgH/CCND1, then will reflex to IgH/MMSET (4;14) and IgH/MAF (14;16).</i>		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	<i>BCL2</i>	18q21	BCL2FH	BCL2FM	BCL2FT
FISH for BCL6	<i>BCL6</i>	3q27	BCL6FH	BCL6FM	BCL6FT
FISH for BCR/ABL1	<i>BCR/ABL1</i>	t(9;22)	BCRFSH	BCRFSM	
FISH for BIRC3/MALT1	<i>BIRC3/MALT1 (API2/MALT1)</i>	t(11;18)			T1118
FISH for CBFB/MYH11	<i>CBFB</i>	inv(16)	INV16F	INV16M	
FISH for CCND1	<i>CCND1</i>	11q13	CCND1F	CCND1M	CCND1T
FISH for CRLF2 Rearrangement	<i>CRLF2</i>	Xp22.33/Yp11.32	CRLF2B	CRLF2M	
FISH for IGH::CRLF2 Rearrangement	<i>CRLF2, IGH</i>	Xp22.33/Yp11.32 and 14q32.33	CRIGHB	CRIGHM	
FISH for del (5q)	<i>EGR1</i>	5q31	5QFSH	5QFSBM	
FISH for del (7q)	<i>D7S486</i>	7q31	FISH7Q	FSH7QM	
FISH for DUSP22/IRF4	<i>DUSP22/IRF4</i>	6p25.3			DUIRFH
FISH for ETV6/RUNX1	<i>ETV6/RUNX1 (TEL/AML1)</i>	t(12;21)	1221FH	1221FM	
FISH for FGFR1	<i>FGFR1</i>	8p12	FGFR1F	FGFR1M	
FISH for IGH/BCL2	<i>IGH/BCL2</i>	t(14;18)	FSHFCL	FSFCLM	T1418
FISH for IGH/CCND1	<i>IGH/CCND1</i>	t(11;14)	FSHMCL	FSMCLM	T1114
FISH for IGH/MYC/CEP8	<i>IGH/MYC</i>	t(8;14)	814FSH	814FSM	T814
FISH for MALT1	<i>MALT1</i>	18q21			MALT1
FISH for MLL	<i>MLL</i>	11q23	MLLFSH	MLLFBM	
FISH for MYC(8q24)	<i>MYC</i>	8q24	MYCFSH	MYCFSM	MYC
FISH for PDGFRA	<i>PDGFRA</i>	4q12	PDGFRA	PGFRAM	
FISH for PDGRRB	<i>PDGRRB</i>	5q33	PDGFRB	PDGFBM	
FISH for PML/RARA	<i>PML/RARA</i>	t(15;17)	APLFSH	APLFBM	
FISH for RARA	<i>RARA</i>	17q21	RARFSH	RARFSM	
FISH for 8;21 Translocation for AML	<i>RUNX1/RUNX1T1 (AML1/ETO)</i>	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	<i>TP63</i>	3q28			TP63FH