

PATIENT INFORMATION (PLEASE PRINT IN BLACK INK)
Last Name First MI
Address Birth Date Sex M F
City SS #
State Zip Home Phone
Hospital/Physician Office Patient ID # Accession #

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

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.

ORDERING PHYSICIAN CONTACT

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: / /
ABN: Yes No WORKERS COMP: Yes No DOI:
PRIMARY: Medicare Medicaid 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)

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

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

SPECIMEN INFORMATION

CLINICAL INFORMATION

Please indicate number of tubes, vials, slides, tissue blocks provided.
Collection Date: / / Time:
Body Site: Specimen ID (#):
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

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

INDIVIDUAL DIAGNOSTIC TESTS

- FLOW CYTOMETRY
Leukemia/Lymphoma Panel RLLIIP
Lymphoma Panel for Tissue/Fluid RLLYMP
PNH, High Sensitivity, FLAER, Peripheral Blood Only PNHPNL
CHROMOSOME ANALYSIS
Chromosome Analysis, Bone Marrow, Reflex to AML FISH CHRAML
Chromosome Analysis, Bone Marrow, Reflex to MDS FISH CHRMD5
Chromosome Analysis, Bone Marrow, Reflex to Microarray BMCHF
Cytogenetic Analysis, Bone Marrow CHRBMH
Cytogenetic Analysis, Leukemic Blood CHRBL
Cytogenetic Analysis Lymph Node CHRSL

- MOLECULAR TESTING
CC-SIGN™ NEXT-GENERATION SEQUENCING PANELS
Hematologic Neoplasm NGS Panel 62-gene, Blood HNPNGS
Hematologic Neoplasm NGS Panel 62-gene, Bone Marrow HNMNGS
ALL NGS Panel 25-gene, Bone Marrow ALLMRW
ALL NGS Panel 25-gene, Blood ALLPBL
Myeloid NGS Panel 50-gene, Bone Marrow MYMNGS
Myeloid NGS Panel 50-gene, Blood MYPNGS
Chronic LPD NGS Panel 7-gene, Bone Marrow LPMNGS
Chronic LPD NGS Panel 7-gene, Blood LPPNGS
MPN NGS Panel 3-gene, Blood MPNP
MPN NGS Panel 3-gene, Bone Marrow MPNM
FLUORESCENCE IN SITU HYBRIDIZATION (FISH)
BCR/ABL1, t(9;22) BCRFSH
FISH for Aggressive B-cell Lymphoma FABCEL
FISH for AML Panel FAMLPN
FISH for B-ALL Panel FSHLL
FISH for CLL Panel (peripheral blood only) CLLFSH
FISH for MDS Panel FSHMDS
FISH for MPN Panel MPNFSH
FISH for Myeloma Panel FSHPCM
IGH/CCND1, t(11;14) FSHMCL
PML/RARA, t(15;17) APLFSH
Other
Other

- ALL
B-ALL Fusion Detection Full Panel, Multiplex RT-PCR BALLFP
BCR/ABL1 p190 RT-PCR, Quantitative I90PCR
AML/MDS
CEBPA Mutation CEBPA
FLT3 ITD/D835 Mutation FLT3
NPM1 Mutation NPM1
CLL
IGHV Sequencing IGVH
HEMOGLOBINOPATHY
Alpha Thalassemia Gene Deletions ATHALS
LYMPHOMA
B-Cell Clonality (IGH and IGH Gene Rearrangement) BCBMD
IGH Gene Rearrangement IGHPCR
IGK Gene Rearrangement IGKPCR
MYD88 L265P Mutation Detection MYD88
T-Cell Clonality (TCRB and TCRG Gene Rearrangement) TCBMD
TCRB Gene Rearrangement TCRB
TCRG Gene Rearrangement TGAMMA
MEYLOPROLIFERATIVE NEOPLASMS
BCR/ABL Kinase Domain Mutation Analysis KINASE
BCR/ABL1 Qualitative Multiplex RT-PCR BCRQL
BCR/ABL1 p210 RT-PCR, Quantitative BCRPCR
CALR Mutation Analysis CALR
JAK2 Exon 12-15 Sequencing JAKNON
JAK2 V617F Mutation Detection JAK2
KIT D816V PCR KITMST
MPL Mutation Sequencing MPL

FISH PROBES AND SQ CODES

Test Name	Gene(s)	Location	SQ Acronym	Blood/ Marrow	Paraffin
FISH for BCL2	<i>BCL2</i>	18q21	BCL2FH	Y	Y
FISH for BCL6	<i>BCL6</i>	3q27	BCL6FH	Y	Y
FISH for BCR/ABL1	<i>BCR/ABL1</i>	t(9;22)	BCRFISH	Y	N
FISH for BIRC3/MALT1	<i>BIRC3/MALT1 (API2/MALT1)</i>	t(11;18)		N	Y
FISH for CBFβ/MYH11	<i>CBFB</i>	inv(16)	INV16F	Y	N
FISH for CCND1	<i>CCND1</i>	11q13	CCND1F	Y	Y
FISH for del (5q)	<i>EGR1</i>	5q31	5QFSH	Y	N
FISH for ETV6/RUNX1	<i>ETV6/RUNX1 (TEL/AML1)</i>	t(12;21)	1221FH	Y	N
FISH for FGFR1	<i>FGFR1</i>	8p12	FGFR1F	Y	N
FISH for IGH	<i>IGH</i>	14q32		N	Y
FISH for IGH/BCL2	<i>IGH/BCL2</i>	t(14;18)	FSHFCL	Y	Y
FISH for IGH/CCND1	<i>IGH/CCND1</i>	t(11;14)	FSHMCL	Y	Y
FISH for IGH/MYC	<i>IGH/MYC</i>	t(8;14)	814FSH	Y	Y
FISH for MALT1	<i>MALT1</i>	18q21		N	Y
FISH FOR MLL	<i>KMT2A (MLL)</i>	11q23	MLLFSH	Y	N
FISH for MYC(8q24)	<i>MYC</i>	8q24	MYCFISH	Y	Y
FISH for PDGFRA	<i>PDGFRA</i>	4q12	PDGFRA	Y	N
FISH for PDGFRB	<i>PDGFRB</i>	5q33	PDGFRB	Y	N
FISH for PML/RARA	<i>PML/RARA</i>	t(15;17)	APLFSH	Y	N
FISH for RARA	<i>RARA</i>	17q21	RARFSH	Y	N
FISH for 8;21 Translocation for AML	<i>RUNX1/RUNX1T1 (AML1/ETO)</i>	t(8;21)	AMLFSH	Y	N
FISH for Trisomy 4, 10	Trisomy 4/10	4 cen, 10 cen	FHT410	Y	N

Panel Names:	Probes		SQ Acronym	Blood/ Marrow?	Paraffin?
FISH for Acute Myeloid Leukemia	t(15;17), t(8;21), inv(16), MLL		FAMLPN	Y	N
FISH for B Lymphoblastic Leukemia (B-ALL)	t(9;22), MLL, t(12;21), 4/10 cen		FSHBLL	Y	N
FISH for Chronic Lymphocytic Leukemia	17p (TP53), 11q (ATM), 12 cen, 13q (D13S319,LAMP1)		CLLFSH	Y – peripheral blood only	N
FISH for Myelodysplastic Syndrome	5q (EGR1), 7q (D7S486), 8 cen, 20q (D20S108)		FSHMDS	Y	N
FISH for Myeloma (on plasma cells)	17p (TP53), 13q (RB1), 14q32 (IGH). If IGH pos, add t(11;14), t(4;14), t(14;16)		FSHPCM	Y	N
FISH for Myeloproliferative Neoplasms	t(9;22), 4q12 (PDGFRA), 5q33 (PDGFRB), 8p12 (FGFR1)		MPNFSH	Y	N