

## PATIENT INFORMATION

Last Name \_\_\_\_\_

First Name \_\_\_\_\_ M.I. \_\_\_\_\_

DOB \_\_\_\_/\_\_\_\_/\_\_\_\_ Gender:  Male  Female  Other \_\_\_\_\_

Address \_\_\_\_\_

City \_\_\_\_\_ State \_\_\_\_\_ Zip \_\_\_\_\_

Phone \_\_\_\_\_ Patient ID \_\_\_\_\_

## BILLING INFORMATION

**Required: Please attach a copy of the patient's insurance and demographic information.**

Bill to:  Insurance  Medicare  Referring Facility (Hospital/Client)  Patient

Patient Status:  Inpatient  Outpatient  Non-hospital patient

## PHYSICIAN INFORMATION

Authorized Signature \_\_\_\_\_ Date \_\_\_\_\_

Please Fax Duplicate Report to Physician \_\_\_\_\_ Fax \_\_\_\_\_

## CLINICAL INFORMATION (PLEASE PROVIDE DIAGNOSIS/INDICATION/PATIENT HISTORY/CBC/ICD 10 CODES)

## SPECIMEN INFORMATION

Date Collected \_\_\_\_/\_\_\_\_/\_\_\_\_ Time Collected \_\_\_\_:\_\_\_\_  AM  PM Specimen ID \_\_\_\_\_

**BONE MARROW (LEFT/RIGHT)** **PERIPHERAL BLOOD** **FLUID**

Aspirate  Clot  Core Smears # \_\_\_\_\_ Purple Tops # \_\_\_\_\_ Green Tops # \_\_\_\_\_ **PARAFFIN BLOCKS #** \_\_\_\_\_

Smears # \_\_\_\_\_ Touch Preps # \_\_\_\_\_ **LYMPH NODE** **SLIDES** \_\_\_\_\_

Green Tops # \_\_\_\_\_ Purple Tops # \_\_\_\_\_ **FINE NEEDLE ASPIRATE** **OTHER** \_\_\_\_\_

## TEST REQUESTED

**20/20 CORE EVALUATION REPORT**

Pathology Consultation with Morphologic Interpretation and/or Ancillary Studies (Flow Cytometry, Routine Chromosome Analysis, FISH and/or PCR); if clinically indicated.

**MORPHOLOGY**

Bone Marrow  Blood  Second Opinion  Surgical Pathology

**FLOW CYTOMETRY**

Please select service:  Global  TC  PC

Acute Leukemia Panel (ALL, AML, MDS, MPN)  CLL Panel

Lymphoma Panel (B-NHL, T-NHL, NK Cell Neoplasm, HCL)  Myeloma Panel

Immune Deficiency (HIV)  PNH Panel

Other: \_\_\_\_\_

**MOLECULAR**

Quantitative BCR-ABL p210/p190  T-Cell Clonality Assessment

ABL1 Kinase Domain Mutation  B-Cell Clonality Assessment

CALR Mutation Analysis  IgVH Hypermutation Analysis

MPL Mutation Analysis  FLT3/NPM1 Mutation Analysis

JAK2 Exon 12-13  FLT3 Mutation Analysis

JAK2 V617F Mutation Analysis  NPM1 Mutation Analysis

If negative reflex to:  cKIT (D816V) Mutation Analysis

CALR  PML/RARA Quantitative

MPL  CEBPA Mutation Analysis

JAK2 Exon 12  MLL-PTD Mutation Analysis

MPN Molecular Panel (JAK2 V617F). If negative, reflex to CALR; if negative, reflex to MPL; if negative, reflex to JAK2 Exon 12 Mutation Analysis.

Other: \_\_\_\_\_

**NEXT-GEN SEQUENCING**

Myeloid Molecular Panel  MDS Molecular Panel

MPN Molecular Panel  Lymphoid Molecular Panel

AML Molecular Panel  Other: \_\_\_\_\_

**CYTOGENETICS**

Routine Chromosome Analysis  Specimen Culture and Hold

**FISH PANELS: HEMATOLOGY**

ALL  CLL  Myeloma  MPN Eosinophilia

AML  CML  MDS  B-Cell NHL

Specimen Culture and Hold

**FISH PROBES: HEMATOLOGY**

ALK/Lymphoma (2p23)  IGH/CCND1 t(11;14)  RUNX1T1/RUNX1

ATM (11q22)  IGH/FGFR3 t(4;14)  (ETO/AML1) t(8;21)

BCL6 (3q27)  IGH/MAF t(14;16)  TP53 (17p13)

BCR-ABL/ASS1 t(9;22)  IGH/MYC t(8;14)  5q-/-5/+5

CBFβ/MYH11 inv(16)  MALT1 (18q21)  6q21/6q23

CKS1B/CDKN2C (1p/1q)  MLL (11q23)  7q-/-7

ETV6/RUNX1 t(12;21)  c-MYC (8q24)  +8

FGFR1 (8p11)  PDGFRA (4q12)  +12

IGH (14q32)  PDGFRB (5q32)  13q-/-13

IGH/BCL2 t(14;18)  PML/RARA t(15;17)  20q-

Other: \_\_\_\_\_

**FISH: SOLID TUMOR**

ALK Rearrangement  MET Amplification

HER2 Amplification  ROS1 Rearrangement

EWSR1 Rearrangement  Other: \_\_\_\_\_

**POC PLOIDY**

FISH on Formalin Fixed Products of Conception, Common Trisomies and Triploidies, Chromosomes 13/16/18/21/X/Y

**LABORATORY USE ONLY**

Bone Marrow Clinic Patient

## FLOW CYTOMETRY

**Acute Leukemia Panel (ALL, AML, MDS, MPN):** CD2, CD3, CD4, CD5, CD7, CD8, CD10, CD11C, CD13, CD14, CD15, CD16, CD19, CD20, CD22, CD23, CD25, CD33, CD34, CD45, CD56, CD57, CD117, CD200, HLA-DR, KAPPA, LAMBDA

**Acute Leukemia-Intracytoplasmic Panel:** CD3, CD22, CD79a, MPO, TdT

**Lymphoma Panel (B-NHL, T-NHL, NK Cell Neoplasms):** CD2, CD3, CD4, CD5, CD7, CD8, CD10, CD11C, CD16, CD19, CD20, CD22, CD23, CD25, CD34, CD45, CD56, CD57, CD200, HLA-DR, KAPPA, LAMBDA

**CLL-Prognostic Panel (Add to Lymphoma Panel):** CD3, CD19, CD38, CD45, CD49d

**Myeloma Panel (Add to Lymphoma Panel):** CD38, CD138, KAPPA(Cy), LAMBDA(Cy)

**PNH Panel:** CD14, CD15, CD24, CD45, CD59a, CD64, CD235a, FLAER

**Immune Deficiency (HIV):** CD3, CD4, CD8, CD45

## FISH PROBES (CALL 1.877.617.4445 FOR ANY PROBES NOT LISTED HERE)

**ALL Panel:** BCR/ABL (t(9;22)), D10Z1/D17Z1/CHIC2 (Hyperdiploidy), E2A Break-Apart (E2A Rearrangement), IGH Break-Apart (IGH Rearrangement), MLL Break-Apart (MLL Rearrangement), MYC Break-Apart (MYC Rearrangement), P16/D9Z3 (P16 Deletion), TEL/AML1 (t(12;21))

**AML Panel:** AML1/ETO (t(8;21)), CBFβ/MYH11 (Inv(16)), EGR1/5p15.31 (5q Deletion), MLL Break-Apart (MLL Rearrangement), P53/D17Z1 (P53 Deletion), PML/RARA (t(15;17)), PTPRT/MYBL2 (20q Deletion), RELN/TES (7q Deletion)

**CLL Panel:** 13q14.3 (RB1 Deletion), ATM/D11Z1 (ATM Deletion), D12Z1 (Trisomy 12), IGH/CCND1 (t(11;14)), MYB/D6Z1 (6q Deletion), p53/D17Z1 (P53 Deletion)

**CML Panel:** BCR/ABL1-ASS

**Myeloma Panel:** 13q14.3 (RB1 Deletion), 5q/-5/+5, CKS1B/CDKN2C, IGH Break-Apart (IGH Rearrangement), IGH/CCND1 (t(11;14)), IGH/FGFR3 (t(4;14)), IGH/MAF (t(14;16)), P53/D17Z1 (P53 Deletion)

**MDS Panel:** 13q14.3 (13q Deletion), CEP8 (Trisomy 8), EGR1/5p15.31 (5q Deletion), MLL Break-Apart (MLL Rearrangement), P53/D17Z1 (P53 Deletion), PTPRT/MYBL2 (20q Deletion), RELN/TES (7q Deletion)

**MPN Eosinophilia Panel:** FIP1L1/CHC2/PDGFRα Deletion/Fusion, FGFR1 Rearrangement, PDGFRB Rearrangement

**B-Cell NHL Panel:** BCL6 Rearrangement, IGH/BCL2 (t(14;18)), IGH/CCND1 (t(11;14)), MALT1 Rearrangement, MYC Rearrangement

## MOLECULAR STUDIES

Quantitative BCR-ABL p210/190  
 ABL1 Kinase Domain Mutation  
 CALR Mutation Analysis  
 MPL Mutation Analysis  
 JAK2 Exon 12-13

JAK2 V617F Mutation Analysis  
 MPN Molecular Panel  
 T-Cell Clonality Assessment  
 B-Cell Clonality Assessment  
 IgVH Hypermutation Analysis

FLT3/NPM1 Mutation Analysis  
 FLT3 Mutation Analysis  
 NPM1 Mutation Analysis  
 cKIT (D816V) Mutation Analysis  
 PML/RARA Quantitative

CEBPA Mutation Analysis  
 MLL-PTD Mutation Analysis

## NEXT-GEN SEQUENCING

**Myeloid Molecular Panel:** ASXL1, BCOR, BRAF, CALR, CEBPA, CBL, CSF3R, DNMT3A, ETV6, EZH2, GATA2, GNAS, IDH1, IDH2, JAK2, KIT, KRAS, MPL, NF1, NPM1, NRAS, PDGFRα, PDGFRβ, PHF6, PTPN11, RAD21, RUNX1, SETBP1, SF3B1, SMC1A, SMC3, SRSF2, STAT3, STAT5b, TET2, TP53, U2AF1, WT1, ZRSR2

**MPN Molecular Panel:** CALR, CSF3R, JAK2, MPL, SETBP1

**AML Molecular Panel:** ASXL1, BCOR, CEBPA, DNMT3A, EZH2, IDH1, IDH2, KIT, KRAS, NPM1, NRAS, PHF6, RUNX1, SF3B1, SRSF2, STAG2, TET2, TP53, U2AF1, WT1, ZRSR2

**MDS Molecular Panel:** ASXL1, ETV6, EZH2, RUNX1, TP53

**Lymphoid Molecular Panel:** ABL1, ALK, ATM, BCL2, BCL6, BCOR, BIRC3, BRAF, BTK, CARD11, CCND1, CD79A, CD79B, CDKN2A, CREBBP, CTCF, CXCR4, DDX3X, EP300, ETV6, EZH2, FAT1, FBXW7, FOXO1, GNAI3, GNAI2, HIST1H1E, ID3, IKZF1, ITPKB, JAK1, JAK2, JAK3, KMT2A/MLL, KRAS, MAPK1, MED12, MEF2B, MYC, MYD88, NF1, NOTCH1, NOTCH2, NRAS, P2RY8, PDGFRβ, PIK3CD, PIM1, PLCG2, POT1, PRDM1, PTEN, PTPN11, RB1, RHOA, RIPK1, S1PR2, SAMHD1, SF3B1, SGK1, SPEN, SRC, STAT3, STAT5B, SYK, TBL1XR1, TCF3, TNFAIP3, TNFRSF14, TP53, TRAF2, TRAF3, UBR5, XPO1, ZMYM3

## PREFERRED SPECIMEN REQUIREMENTS

Specimen Type	Flow Cytometry	Cytogenetics	FISH <sup>1</sup>	Molecular <sup>1</sup>	Immunostains
Peripheral Blood <sup>2</sup>	Lavender <sup>3</sup> or Green Top <sup>4</sup>			Lavender Top <sup>3</sup>	Not Applicable
Bone Marrow	Tissue Media			Tissue Media RNA Fixative or 10% Neutral Buffer	10% Formalin or Paraffin Block
Lymph Node or Tissue (Fresh)	Not Applicable			Paraffin Block or Unstained Slides	Paraffin Block or Unstained Slides
Formalin Fixed Paraffin Embedded	Tissue Media			Paraffin Block <sup>5</sup>	Cell Block
Fine Needle Aspirate	Sterile Container				
Body Fluid <sup>6</sup>					

<sup>1</sup> Decalcified samples not suitable | <sup>2</sup> Please provide copy of CBC if available | <sup>3</sup> Lavender Top: EDTA | <sup>4</sup> Green Top: Sodium Heparin | <sup>5</sup> Only for some tests; contact us for details | <sup>6</sup> Minimum of 5 mL is preferred

## SPECIMEN HANDLING AND TRANSPORTATION

**Storage:** All specimens should be stored at room temperature.

**Transportation:** Please use cold packs for transportation without placing packs in direct contact with specimens. Please fill out the requisition form completely and place all specimens in a CorePath shipping kit using the pre-printed air bills supplied by CorePath.

**Schedule a Pick-Up:** Call CorePath Laboratories at 1.877.617.4445 to schedule a pick-up. In the San Antonio area, call 210.617.4445 to schedule a courier pick-up.