



MOLECULAR HEMATOLOGY REQUISITION

Scanning Label or Accession # (lab only)

| | | | | | | |
|--------------|---|-----------------|------------------|-------------|--|--|
| Patient | PHN | | Expiry: | | Date of Birth (dd-Mon-yyyy) | |
| | Legal Last Name | | Legal First Name | | Middle Name | |
| | Alternate Identifier | | Preferred Name | | <input type="checkbox"/> Male <input type="checkbox"/> Non-binary | <input type="checkbox"/> Female <input type="checkbox"/> Prefer not to disclose |
| | Address | | City/Town | | Prov | Postal Code |
| Provider (s) | Authorizing Provider Name (last, first, middle) | | | | Copy to Name (last, first, middle) | |
| | Address | | Phone | | Address | |
| | CC Provider ID | CC Submitter ID | Legacy ID | Phone | Phone | |
| | Clinic Name | | | Clinic Name | Clinic Name | |
| Collection | Date (dd-Mon-yyyy) | | Time (24 hr) | | Location | |
| Collector ID | | | | | | |

DO NOT use this requisition for bone marrow test requests; see form REQ9061BM *Bone Marrow Pathology Requisition*.

DO NOT use this requisition for **Chimerism Studies**

Screening/monitoring hematopoietic cell chimerism

For peripheral blood chimerism testing, see form FC3200 *Flow Cytometry Requisition*.

For bone marrow chimerism test requests, see form REQ9061BM *Bone Marrow Pathology Requisition*.

Philadelphia Chromosome Transcript Analysis (Nested PCR)

Screening for CML/ALL, BCR-ABL1 gene transcripts

☐ PHLR PB (Peripheral Blood)

Quantitative PCR Analysis of BCR-ABL1 Fusion Gene Transcripts (Q-PCR)

Monitoring residual disease in CML

☐ QPCRPH1 (Peripheral Blood)

APL Transcript Analysis (Nested PCR)

Screening/monitoring PML-RARA gene transcripts

☐ APL PB (Peripheral Blood)

JAK2-V617F Mutation Analysis

Screening for myeloproliferative neoplasms

☐ JAK2 PB (Peripheral Blood)

FLT3 and NPM1 Mutation Analysis*

Screening for FLT3 and NPM1 gene mutations

☐ FLT3 NPM1 PB (Peripheral Blood)

*FLT3/NPM1 mutation analysis may include CEBPA testing if clinically indicated

Molecular Hemostasis Studies (Factor VIII inversion, 2N vonWillebrand Factor, Multimers, Carrier Study: Hemophilia A, B or vWD)

☐ MH HEMST (Peripheral blood)

State test requested: _____

Family history: _____

Miscellaneous Test: Miscellaneous hemostasis (other than above) and oncology tests

☐ MH MISC (Peripheral blood)

State test requested: _____