



# **Oncology Office Hematology Requisition**

Client Information	Patient Information
Required Information	Last Name:
Account #: Account Name:	First Name: M.I Medical Record #:
Street Address:	Date of Birth: mm / dd / yyyy Other Pt ID/Acct #:
City, ST, ZIP:	Client represents it has obtained informed consent from patient to perform the services described herein.
Phone: Fax:	Specimen Information
Additional Reporting Fax:	☐ Mobile Phlebotomy Request  NeoGenomics will reach out to patient to schedule appointment - Patient Phone:
Requisition Completed by: Date:	Specimen ID: Block ID:
Ordering Physician: NPI #:	Fixative/Preservative:  Collection Date: mm
Treating Oncologist/Physician: NPI #:	Netrieved Date. IIIII/ dd/ yyyy
(please print: Last, First):  The undersigned certifies that he/she is licensed to order the test(s) listed below and that such test(s) are medi-	Hospital Discharge Date: mm/dd/yyyyBody Site:
cally necessary for the care/treatment of this patient.  Authorized Signature:	□ Primary □ Metastasis – If Metastasis, list Primary:
	☐ Bone Marrow [must provide CBC Report]:
Billing Information	Green Top(s)         Purple Top(s)         Core Biopsy         Clot           □ Peripheral Blood: Green Top(s)         Purple Top(s)         Other
Required: Please include face sheet and front/back of patient's insurance card.	☐ Smears: Air Dried Fixed Stained (type of stain)
Patient Status (Must Choose 1):	☐ Slides # Unstained Stained ☐ H&E
T OP Molecular to MCR, all other testing to Client	☐ Paraffin Block(s) #: ☐ Choose best block (for global molecular/NGS testing only)
☐ Hospital Patient (out)  See back for definitions.  ☐ Hospital Patient (out) ☐ Bill charges to other Hospital/Facility: ☐ Bill charges to other Hospital/Facility:	Submit ≤4 blocks. Blocks will be combined for molecular testing when necessary.  For all other testing, specify which block to use for each if sending multiple blocks. See back for details.
Prior Authorization # See neogenomics.com/billing for more info.	Specimen Retrieval
Clinical Information	Client Services will request specimen from Pathology site.
Required: Please attach patient's pathology report (required), clinical history, and other applicable report(s).	Pathology Site:
ICD 10 (Diagnosis) Code/Narrative (Required):	Address:
Reason for Referral:	Required Items
□ New Diagnosis □ Relapse/Refractory □ Monitoring □ MRD	☐ Patient Demographics ☐ Pathology Report
Bone Marrow Transplant ☐ None ☐ Autologous ☐ Allogeneic ☐ Sex Mismatch	☐ Copy of Insurance Card ☐ Clinical History ☐ CBC Within Last 30 Days ☐ Relevant Treatment History
□ Blood and/or Bone Marrow □ Paraffin block for Morphology to follow Lymphoma Consult □ Lymph Node/Tissue for Lymphoma* *Split fresh specimens to RPMI and formalin □ Paraffin block for Morphology to follow  or marked by the client) to provide on the materials submitted.  Please attach CBC for Blood and E □ Do not add NGS Profile without	
Morphology	Molecular Genetics
☐ Blood and/or Bone Marrow	☐ ABL1 Kinase Domain (Gleevec® resistance) ☐ JAK2 V617F - Qualitative
NeoTYPE® and Neo Comprehensive™ Cancer Profiles	☐ BCR-ABL1 Standard p210, p190 ☐ If negative, reflex to CALR
☐ AITL/Peripheral T-Cell Lymphoma Profile ☐ Neo Comprehensive – Heme Cancers	☐ BCR-ABL1 Standard p210, p190 with reflex to ABL1 Kinase Domain if positive ☐ JAK2 V617F - Quantitative
☐ ALL Profile ☐ Neo Comprehensive – Myeloid Disorders	☐ BCR-ABL1 Standard p210, p190 with reflex to ☐ KIT (c-KIT) ☐ MPL Mutation Analysis
☐ AML Prognostic Profile ☐ Follicular Lymphoma Profile (FFPE only)	☐ BCR-ABL1 Non-Standard p230 ☐ MPN JAK2 V617F with Sequential Reflex
☐ CLL Profile ☐ Lymphoid Disorders Profile ☐ Lymphoma Profile ☐ Lymphoma Profile	□ BRAF to JAK2 Exon 12-13, CALR, & MPL □ BTK Inhibitor Acquired Resistance Panel □ MYD88 Mutation Analysis
☐ MDS/CMML Profile	☐ Calreticulin (CALR) Mutation Analysis ☐ NPM1 MRD Analysis ☐ NPM1 Mutation Analysis ☐ NPM1 Mutation Analysis
Flow Cytometry	☐ CXCR4 Mutation Analysis ☐ PML- RARA, t(15;17)
Please attach CBC with all flow requests on blood (required).	□ FLT3 Mutation Analysis       □ Rapid AML Therapeutic Panel         □ IDH1/IDH2 by PCR       □ RUNX1-RUNX1T1 (AML1-ETO), t(8;21)
Diagnostic/Prognostic Panels MRD Panels	☐ IgH Clonality by NGS  • Baseline testing of original primary sample required  ☐ T-Cell Receptor Gamma  ☐ T-Cell Receptor Beta
☐ Standard L/L Panel (24 Markers) ☐ AML MRD Panel	☐ IgVH Mutation Analysis ☐ TP53 Mutation Analysis
☐ Extended L/L Panel (31 Markers) ☐ B-ALL MRD (Bone Marrow) ☐ B-ALL MRD (Peripheral Blood)	□ inv(16) CBFB-MYH11 □ Other
High Sensitivity PNH	HemeFISH®  ☐ Apaplastic Large Cell Lymphoma (ALCL) ☐ High-Grade B-Cell Lymphoma Reflex
☐ Myeloma (MM) MRD	□ ALL - Adult □ Low-Grade/Small B-Cell Lymphoma
Cytogenetics	☐ ALL - Pediatric ☐ MDS Extended ☐ MDS Standard
☐ Oncology Chromosome Analysis	☐ AML Standard ☐ MPN
☐ Reflex to FISH if cytogenetics is normal (reflex FISH panel must be selected)	☐ AML Non-Favorable Risk ☐ NHL
☐ Reflex to FISH if cytogenetics is incomplete (<20 metaphases) ☐ MDS Standard FISH	☐ CLL ☐ Do not reflex to IgH Complex
☐ MDS Extended FISH	☐ Eosinophilia ☐ Plasma Cell Myeloma IgH Complex
☐ Follow-up Constitutional Chromosome Analysis	☐ High-Grade/Large B-cell Lymphoma ☐ Plasma Cell Myeloma Prognostic Panel ☐ Reflex to BCL6/MYC, IGK/MYC, IGL/MYC if ☐ Other ☐
(only if recommended by Oncology Chromosome Analysis)	MYC+ and IGH/MYC-

### **Specimen Requirements**

Refrigerate specimen if not shipping immediately and use cool pack during transport. Please call the Client Services team with any questions regarding specimen requirements or shipping instructions at 866.776.5907 option 3. Please refer to the website for specific details on each specimen.

## **Additional Billing Information**

Any organization referring specimens for testing services pursuant to this Requisition Form ("Client") expressly agrees to the following terms and conditions.

- 1. Binding Service Order. This Requisition Form is a contractually binding order for the services ordered hereunder ("Services") and Client agrees that it is financially responsible for all tests billable to Client hereunder.
- 2. Third Party Billing by NeoGenomics and Right to Bill Client. Client agrees to accurately indicate on the front of the Requisition Form that either Client should be billed (e.g., Client receives reimbursement pursuant to a non-fee-for-service basis, including, but not limited to, a capitated, diagnostic related group ("DRG"), per diem, all-inclusive, or other such bundled or consolidated billing arrangement) or NeoGenomics should bill the applicable federal, state or commercial health insurer or other third party payer (collectively, "Payers") for all Services ordered pursuant to this Requisition Form. For all such Services billable to Payers, Client agrees to provide all billing information necessary for NeoGenomics to bill such payer. In the event NeoGenomics: (i) does not receive the billing information required for it to bill any Payers within ten days of the date that any Services are reported by NeoGenomics; (ii) the Services were performed for patients who have no Payer coverage arrangements; or (iii) the Payer identified by Client denies financial responsibility for the Services and indicates that Client is financially responsible, NeoGenomics shall have the right to bill such Services to Client.

# **Additional Specimen Information**

If submitting multiple blocks, clients must indicate either "Choose best block (global molecular/NGS testing only)", or assign the selection of blocks to individual tests. If multiple blocks are sent without a selection, they will be held until clarification is provided. Please call the Client Services team with any questions regarding specimen information.

# **Definitions of Patient Status for Specimen Origin**

Non-Hospital Patient: Patient is not registered at a hospital (neither an in-patient nor out-patient)

Hospital Patient (in): Patient is registered and admitted to a hospital overnight

Hospital Patient (out): Patient is registered and admitted to a hospital, then discharged before the end of the day

#### **Test Descriptions**

Please see complete test descriptions and all available tests at our website, www.neogenomics.com/test-menu.

### **Test Notations**

# Specimen Usage

NeoGenomics makes every effort to preserve and not exhaust tissue, but in small and thin specimens, there is a possibility of exhausting the specimen in order to ensure adequate material and reliable results.

#### FISH

Plasma cell myeloma FISH panels: Plasma cell enrichment will be performed on bone marrow samples having plasma cell FISH. Sample should be received at NeoGenomics Laboratories within 72 hours of collection.