



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.
-	Specimen Information
Phone: Fax:	
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 #:	
(please print: Last, First): Treating Oncologist/Physician: NPI #:	Collection Date: mm /dd /vvvv Collection Time: AM PM
(please print: Last, First):	Retrieved Date: mm / dd / yyyy / Hospital Discharge Date: mm / dd / yyyy / yyy
The undersigned certifies that he/she is licensed to order the test(s) listed below and that such test(s) are medically necessary for the care/treatment of this patient.	Body Site:
Authorized Signature: Date:	☐ Primary ☐ Metastasis – If Metastasis, list Primary:
	☐ Bone Marrow [must provide CBC Report]:
Billing Information	Green Top(s)
Required: Please include face sheet and front/back of patient's insurance card.	Peripheral Blood: Green Top(s) Other
Patient Status (Must Choose 1): Bill to:	☐ Smears: Air Dried Stained (type of stain)
□ Non-Hospital Patient □ Insurance □ Patient/Self-Pay □ Medicare □ Medicaid □ Client Bill	☐ Slides # Unstained Stained ☐ H&E ☐ Paraffin Block(s) #:
OP Molecular to MCR, all other testing to Client	Choose best block (for global molecular/NGS testing only)
☐ Bill charges to other Hospital/Facility:	Submit ≤4 blocks. Blocks will be combined for molecular testing when necessary.
See back for definitions.	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. Pathology Site:
Required: Please attach patient's pathology report (required), clinical history, and other applicable report(s).	Address:
ICD 10 (Diagnosis) Code/Narrative (Required):	Phone:Fax:
Reason for Referral:	Download House
☐ New Diagnosis ☐ Relapse/Refractory ☐ Monitoring ☐ MRD	Required Items ☐ Patient Demographics ☐ Pathology Report
Bone Marrow Transplant	☐ Copy of Insurance Card ☐ Clinical History
☐ None ☐ Autologous ☐ Allogeneic ☐ Sex Mismatch	☐ CBC Within Last 30 Days ☐ Relevant Treatment History
□ 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 the materials submitted. Please attach CBC for Blood and B Do not add NGS Profile without	
□ Paraffin block for Morphology to follow Morphology	Molecular Genetics
☐ Blood and/or Bone Marrow	☐ JAK2 Exon 12-13 ☐ ABL1 Kinase Domain (Gleevec® resistance) ☐ JAK2 V617F - Qualitative
a blood diligion boile marrow	B-Cell Gene Rearrangement If negative, reflex to JAK2 Exon 12-13
NeoTYPE® and Neo Comprehensive® Cancer Profiles	□ BCR-ABL1 Standard p210, p190 □ If negative, reflex to CALR □ If negative, reflex to MPL
☐ 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
☐ AML Prognostic Profile ☐ Follicular Lymphoma Profile (FFPE only)	BCR-ABL1 Non-Standard p230 if negative MPL Mutation Analysis BCR-ABL1 Non-Standard p230 MPN JAK2 V617F with Sequential Reflex
☐ CLL Profile ☐ Lymphoid Disorders Profile	BRAF to JAK2 Exon 12-13, CALR, & MPL
☐ Add IgVH Mutation Analysis ☐ Lymphoma Profile	☐ BTK Inhibitor Acquired Resistance Panel ☐ MYD88 Mutation Analysis
□ MDS/CMML Profile	☐ Calreticulin (CALR) Mutation Analysis ☐ NPM1 MRD Analysis ☐ NPM1 MRD Analysis ☐ NPM1 Mutation Analysis
	□ CEBPA Mutation Analysis □ NPM1 Mutation Analysis □ CXCR4 Mutation Analysis □ PML- RARA, t(15;17)
Flow Cytometry	☐ FLT3 Mutation Analysis ☐ Rapid AML Therapeutic Panel
Please attach CBC with all flow requests on blood (required).	☐ IDH1/IDH2 by PCR ☐ RUNX1-RUNX1T1 (AML1-ETO), t(8;21)
Diagnostic/Prognostic Panels MRD Panels	☐ IgH Clonality by NGS ☐ T-Cell Receptor Gamma Baseline testing of original primary sample required ☐ 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)	□ inv(16) CBFB-MYH11 □ Other
☐ High Sensitivity PNH ☐ B-ALL MRD (Peripheral Blood)	HemeFISH®
□ CLL MRD	☐ Anaplastic 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
	☐ AML Standard ☐ Reflex to ETV6 (12p13), MLL (11q23)
☐ Oncology Chromosome Analysis ☐ Pelloy to EISH if outconnotice is normal (reflex EISH panel must be selected)	Reflex to 5q-/-5, 7q/-7, DEK/NUP214 t(6;9), p53 and +19 if negative
☐ Reflex to FISH if cytogenetics is normal (reflex FISH panel must be selected) ☐ Reflex to FISH if cytogenetics is incomplete (<20 metaphases)	(17p13.1)NF1 (17q11) and NUP98 (11p15) if negative MPN
☐ MDS Standard FISH	BCR/ABL1/ASS1 t(9:22)
☐ MDS Extended FISH	CLL Do not reflex to left Complex
☐ Follow-up Constitutional Chromosome Analysis	☐ Eosinophilia ☐ Plasma Cell Myeloma IgH Complex
(only if recommended by Oncology Chromosome Analysis)	Reflex to BCL6/MYC, IGK/MYC, IGL/MYC if MYC+
Uther	and IGH/MYC- Plasma Cell Enrichment will be performed on all bone marrow samples having plasma cell FISH tests

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.