

Client Information

Required Information

Account #: Account Name: Street Address: City, ST, ZIP: Phone: Fax: Additional Reporting Fax:

Requisition Completed by: Date: Ordering Physician: NPI #: Treating Oncologist/Physician: NPI #: Authorized Signature: Date:

Billing Information

Required: Please include face sheet and front/back of patient's insurance card.

Patient Status (Must Choose 1): Hospital Patient (in) Hospital Patient (out) Non-Hospital Patient Bill to: Client Bill Insurance Medicare Medicaid Patient/Self-Pay

Prior Authorization # See neogenomics.com/billing section for more info.

Clinical Information

Required: Please attach patient's pathology report (required), clinical history, and other applicable report(s).

ICD 10 (Diagnosis) Code/Narrative (Required): Reason for Referral: New Diagnosis Relapse/Refractory Monitoring MRD

Bone Marrow Transplant (required information for Oncology Cytogenetics):

None Autologous Allogeneic Sex Mismatch

Consultation

COMPASS Comprehensive evaluation including morphology

Blood and/or Bone Marrow Paraffin block for Morphology to follow

COMPASS Select (Without morphology)

Blood and/or Bone Marrow - Morphology performed by client (Morphology report required. Please fax to avoid testing delays.)

Lymphoma Consult

Lymph Node/Tissue for Lymphoma\* Split fresh specimens to RPMI and formalin Paraffin block for Morphology to follow

A NeoGenomics pathologist will select medically necessary tests (with any exceptions noted or marked by the client) to provide comprehensive analysis and professional interpretation for the materials submitted.

Please attach CBC for Blood and Bone Marrow (required).

Do not add NGS Profile without prior approval

Flow Cytometry Please attach CBC with all flow requests on blood or bone marrow (required).

Follow-Up/Add-On panels are available in conjunction with, or after, a Main Panel result has been reported by NeoGenomics or client.

Diagnostic/Prognostic Panels

Standard L/L Panel (24 Markers) Extended L/L Panel (31 Markers) N/A CD4/CD8 Ratio for BAL High Sensitivity PNH T&B Tissue Panel TRBC1/T-Cell Lymphoma Companion

Add-On Tubes

AML B-ALL CLL/Mantle Cell Companion Erythroid-Mega Hairy Cell Mast Cell Plasma Cell T-ALL TRBC1/LGL

Follow-Up Panels

AML B-ALL B-ALL Hairy Cell Plasma Cell T-ALL

MRD Panels

NA B-ALL MRD Panel (BM) NA B-ALL MRD Panel (PB) NA CLL MRD Panel NA Myeloma (MM) MRD Panel

G - Global T - Tech-Only

Specimen Hold Option: Refrigerate and Hold

Tech-Only Opt Out Option: To avoid delay in patient care and as medically necessary for an individual patient, additional markers will be added by the flow lab when abnormal populations are detected. Please refer to NeoGenomics Flow Cytometry Guidelines for additional information on tech-only add-on medical necessity criteria.

Tech-only clients may instruct NeoGenomics to not follow this stated criteria by checking this box.

FISH G - Global T - Tech-Only

HemeFISH Panels

Anaplastic Large Cell Lymphoma (ALCL) ALL - Adult ALL - Pediatric ALL, Ph-Like AML Standard AML Favorable-Risk ALL AML Non-Favorable Risk CLL Eosinophilia High-Grade/Large B-Cell Lymphoma Reflex to BCL6/MYC, IGK/MYC, IGL/MYC if MYC+ and IGH/MYC-

Plasma Cell Myeloma Panels

Plasma Cell Enrichment will be performed on all bone marrow samples having plasma cell FISH tests.

Plasma Cell Myeloma Do not reflex to IgH Complex (applies to global only; tech-only will not reflex) Plasma Cell Myeloma IgH Complex Plasma Cell Myeloma Prognostic Panel

Individual Probes

11q Aberration in NHL 1p36 Deletion ALK for Lymphoma BCL6/MYC t(3;8) BIRC3 (AP12)/MALT1 t(11;18) BCR/ABL1/ASS1 t(9;22) CDKN2A (p16) Deletion for ALL DUSP22-IRF4 Rearrangement IgH/MAFB t(14;20) IGK/MYC t(2;8) IGL/MYC t(8;22) JAK2 (9p24.1) MYC/IgH/CEN8 t(8;14) NUP98 TCL1 (14q32.1) TP63 Rearrangement PML/RARA t(15;17) Other

Specimen Hold Option: Direct Harvest and Hold Plasma Cell Enrichment and Hold

Patient Information

Last Name: First Name: M.I. Medical Record #: Date of Birth: Other Pt ID/Acct #: Client represents it has obtained informed consent from patient to perform the services described herein.

Specimen Information

Specimen ID: Block ID: Fixative/Preservative: Collection Date: Collection Time: Retrieved Date: Hospital Discharge Date: Body Site: Primary Metastasis - If Metastasis, list Primary:

Bone Marrow [must provide CBC and Path Report]: Green Top(s) Purple Top(s) Core Biopsy Clot Peripheral Blood: Green Top(s) Purple Top(s) Other Fresh Tissue (Media Type required): Fluid: CSF Pleural Other FNA cell block: Smears: Air Dried Fixed Stained (type of stain) Slides # Unstained Stained H&E Paraffin Block(s) #:

Choose best block (for global molecular/NGS testing only) 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.

Comments:

Cytogenetics

Oncology Chromosome Analysis Reflex to FISH if cytogenetics is normal (reflex FISH panel must be marked) Reflex to FISH if cytogenetics is incomplete (<20 metaphases) G T MDS Standard FISH G T MDS Extended FISH Other:

Specimen Hold Option: Culture and Hold (liquid samples & lymph nodes; n/a for solid tissues)

Molecular Genetics

ABL1 Kinase Domain (Gleevec resistance)\* CXCR4 Mutation Analysis FLT3 Mutation Analysis MPN JAK2 V617F with Sequential Reflex to JAK2 B-Cell Gene Rearrangement IDH1/IDH2 by PCR NPM1 Mutation Analysis BCR-ABL1 Standard p210, p190\* IgH Clonality by NGS \* Baseline testing of original primary sample required NPM1 MRD Analysis BCR-ABL1 Standard p210, p190\* with reflex to ABL1 Kinase Domain if positive IgVH Mutation Analysis\* PML- RARA, t(15;17)\* BCR-ABL1 Standard p210, p190\* with reflex to BCR-ABL1 Non-Standard p230 if negative inv(16) CBFB-MYH11\* JAK2 V617F - Qualitative\* If negative, reflex to JAK2 Exon 12-13 Molecular only BCR-ABL1 Non-Standard p230\* If negative, reflex to CALR If negative, reflex to MPL RUNX1-RUNX1T1 (AML1-ETO), t(8;21)\* BRAF Mutation Analysis JAK2 V617F - Quantitative T-Cell Receptor Gamma BTK Inhibitor Acquired Resistance Panel JAK2 Exon 12-13\* T-Cell Receptor Beta Calreticulin (CALR)\* KIT (c-KIT) Mutation Analysis TP53 Mutation Analysis CEBPA Mutation Analysis MPL Mutation Analysis Other

Specimen Hold Options: Extract & Hold - DNA Extract & Hold - RNA

\* Test is RNA-based. † Test is both DNA- and RNA-based

NeoTYPE and Neo Comprehensive Cancer Profiles G - Global T - with Tech-Only FISH

AITL/Peripheral T-Cell Lymphoma ALL Profile Follicular Lymphoma Profile N/A Lymphoid Disorders Profile N/A AML Prognostic Profile N/A Lymphoma Profile N/A AML Prognostic Profile + FLT3 by PCR\* N/A MDS/CMML Profile N/A CLL Profile N/A MDS/CMML Profile + FLT3 by PCR\* Add IgVH Mutation Analysis N/A Neo Comprehensive - Heme Cancers N/A Neo Comprehensive - Heme Cancers + FLT3 by PCR\* N/A Neo Comprehensive - Myeloid Disorders N/A Neo Comprehensive - Myeloid Disorders + FLT3 by PCR\*

Specimen Hold Option: Extract & Hold - TNA (all tests are TNA-based)

\*Please see back page for detailed info on Intended Use and Billing for FLT3 by PCR

FlexREPORT

FlexREPORT: Please add summary report option to this case.

## 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.

## Specimen Hold Option Descriptions

To preserve the integrity of samples and avoid unnecessary testing, NeoGenomics Laboratories offers the option of processing samples to maintain specimen integrity for extended periods, without a test order. Any hold order will result in billed charges to the ordering client if testing is not ordered/performed. Specimen Hold Options include:

**FISH: Direct Harvest and Hold:** FISH specimens will be minimally processed and directly harvested while the cells are still viable. Analysis is not performed until the client test order is received. Processed samples will be retained for 28 days.

**Plasma Cell Enrichment and Hold:** Plasma cells will be isolated for bone marrow specimens. Sample should be received at NeoGenomics Laboratories within 72 hours of collection. Analysis is not performed until the client test order is received. Processed samples will be retained for 28 days.

**Flow Cytometry: Refrigerate and Hold:** Flow cytometry samples will be refrigerated and retained for 28 days, however, optimal stability is within 72 hours of collection.

**Molecular Testing: Extract Nucleic Acid and Hold:** Nucleic acid (DNA or RNA or TNA) will be isolated from viable cells and stored in a freezer. **Use this option when it is known which test(s) may be added.** Analysis is not performed until the client test order is received. Processed samples will be retained for 28 days.

## Test Descriptions

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

### FlexREPORT®

FlexREPORT can be ordered on any global or tech-only testing referred to NeoGenomics. This report template can be used to import data and images collected from testing performed outside of NeoGenomics, and incorporated into a one page summary report. Client logo and contact information will be in the header of the FlexREPORT.

### 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.

### FLT3 Testing with NeoTYPE and Neo Comprehensive profiles

**The FLT3 Mutation Analysis test is available as client-bill only when ordered with NeoTYPE and Neo Comprehensive.** The Molecular case reports separately from the NeoTYPE or Neo Comprehensive Profile (which also includes FLT3 gene by NGS) for the purpose of prompt therapy selection in patients with a *new* diagnosis of AML.