

ONCOLOGY GENETIC TESTING REQUISITION

All Information Must Be Completed Before Sample Can Be Processed

PATIENT INFORMATION

Patient Name: _____
Last First MI
Address: _____
Home Phone: _____ MR#: _____
Date of Birth: ____/____/____ Gender: Male Female

INDICATIONS/DIAGNOSIS/ICD-9 CODE

- | | | | |
|---|--|--|---|
| <input type="checkbox"/> Acute Myelogenous Leukemia | <input type="checkbox"/> Glioma | <input type="checkbox"/> Lymphoproliferative Disorder | <input type="checkbox"/> Pancytopenia |
| <input type="checkbox"/> Acute Promyelocytic Leukemia | <input type="checkbox"/> Hodgkin Lymphoma | <input type="checkbox"/> Malignant Melanoma | <input type="checkbox"/> Polycythemia Vera (PV) |
| <input type="checkbox"/> Adenopathy | <input type="checkbox"/> Langerhans Cell Histiocytosis (LCH) | <input type="checkbox"/> Medulloblastoma | <input type="checkbox"/> Sarcoma |
| <input type="checkbox"/> Anemia | <input type="checkbox"/> Leukemia | <input type="checkbox"/> Monoclonal Gammopathy | <input type="checkbox"/> Thrombocytopenia |
| <input type="checkbox"/> Burkitt Lymphoma | <input type="checkbox"/> Leukocytosis | <input type="checkbox"/> Multiple Myeloma | <input type="checkbox"/> Thrombocytosis |
| <input type="checkbox"/> Chronic Lymphocytic Leukemia | <input type="checkbox"/> Leukopenia | <input type="checkbox"/> Myelodysplastic Syndrome/Disease (MDS) | <input type="checkbox"/> Wilms Tumor |
| <input type="checkbox"/> Chronic Myelogenous Leukemia | <input type="checkbox"/> Lung Cancer | <input type="checkbox"/> Myeloproliferative Disease (MPS or MPD) | <input type="checkbox"/> Other _____ |
| <input type="checkbox"/> Colorectal Cancer | <input type="checkbox"/> Lymphocytosis | <input type="checkbox"/> Neutropenia | _____ |
| <input type="checkbox"/> Ewing Sarcoma | <input type="checkbox"/> Lymphoma | <input type="checkbox"/> Non-Hodgkin Lymphoma (NHL) | _____ |

ETHNIC/RACIAL BACKGROUND (Choose All)

- | | |
|--|--|
| <input type="checkbox"/> European American (White) | <input type="checkbox"/> African-American (Black) |
| <input type="checkbox"/> Native American or Alaskan | <input type="checkbox"/> Asian-American |
| <input type="checkbox"/> Pacific Islander | <input type="checkbox"/> Ashkenazi Jewish ancestry |
| <input type="checkbox"/> Latinx-Hispanic _____
(specify country/region of origin) | |
| <input type="checkbox"/> Other _____
(specify country/region of origin) | |

REFERRING PHYSICIAN

Physician Name (print): _____
Address: _____
Phone: (_____) _____ Fax: (_____) _____
Email: _____
Genetic Counselor/Lab Contact Name: _____
Phone: (_____) _____ Fax: (_____) _____
Email: _____
Date: ____/____/____
Referring Physician Signature (REQUIRED)

Contact Information for Results/Questions (if different than ordering provider) :

Name & Title: _____
Phone: (_____) _____ Fax: (_____) _____
Email: _____

BILLING INFORMATION (Choose ONE method of payment):

REFERRING INSTITUTION

Institution: _____
Address: _____
City/State/Zip: _____
Accounts Payable Contact Name: _____
Phone: _____
Fax: _____
Email: _____

COMMERCIAL INSURANCE

Insurance can only be billed if requested at the time of service.

Policy Holder Name: _____
Gender: _____ Date of Birth ____/____/____
Authorization Number: _____
Insurance ID Number: _____
Insurance Name: _____
Insurance Address: _____
City/State/Zip: _____
Insurance Phone Number: _____

See page 3 for more billing information

SAMPLE/SPECIMEN INFORMATION

Has patient received a bone marrow transplant? Yes No

If yes, date of bone marrow transplant _____

Percent engraftment _____

Specimen Date: _____ **Time:** _____

DRAWN BY: _____

***Phlebotomist must initial tube of specimen to confirm sample identity**

DISEASE STATUS:

New diagnosis Remission Relapse E(COG) study COG patient

SPECIMEN TYPE — SEE PAGE 3 FOR SPECIMEN REQUIREMENTS

Bone marrow Oncology blood Lymph node

Formalin fixed paraffin embedded tissue Touch prep Smear

Estimated percent of tumor in sample: _____

Solid tumor (specify): _____ If in media, type: _____

Estimated percent of tumor in sample: _____

Other: _____ WBC _____ % Blasts _____

TEST(S) REQUESTED

SEE PAGE 3 FOR SPECIMEN AND TEST DETAILS

Cytogenetic Chromosome and Microarray Analysis

Oncology Chromosome Analysis

Constitutional (blood) Chromosome Analysis

Oncology Microarray

[Additional 3 mL blood or bone marrow (NaHep) if ordered without chromosomes] — % **Tumor:** _____

FISH

(Additional FISH probes available. See page 3 for details.)

t(9;22) [*BCR/ABL1*]

11q23 [*KMT2A*]

X/Y [Opposite sex BMT]

t(15;17) [*PML/RARα*]

Other (please call lab) _____

Hematologic FISH Panels

(All probes available individually. Please see page 3 for panel descriptions)

ALL Hyperdiploid

Fanconi anemia

ALL Risk Stratification

Multiple myeloma

B-Cell ALL

MDS

Ph-like ALL

Myeloid Malignancy

AML

MPD

APL

SDS

Burkitt Lymphoma

Large cell NHL

CLL

Small cell NHL

Double Hit Lymphoma

Combined NHL

Eosinophilia

T-Cell Lymphoma/Leukemia

Non-Hematologic FISH

Fresh tumor or FFPE slides (include 1 marked H&E slide with FFPE)

See page 3 for specimen requirements

BRAF (7q34) FISH

Ependymoma FISH Panel

High-Grade Glioma FISH Panel

Low-Grade Glioma FISH Panel

Lung Cancer FISH Panel

Medulloblastoma FISH Panel

Melanocytic Tumor FISH Panel

Pilocytic Astrocytoma FISH Panel

Molecular Genetic Analysis (RNA assays)

Samples must be received within 24 hours of collection.

BCR/ABL - QUANTITATIVE (p210)

BCR/ABL - QUANTITATIVE (p190)

BCR/ABL - RT-PCR (QUALITATIVE)

PML/RARα - RT-PCR

Molecular Genetic Analysis (DNA assays)

Samples must be received within 48 hours of collection.

JAK2 QUANTITATIVE (V617F)

PTEN sequencing

Bone marrow engraftment (BME) by STR (Same sex donor & recipient)

Pre-transplant host sample

Post transplant sample

Donor sample

WBC sub-populations engraftment study*

STR (same sex)

X/Y FISH (opposite sex)

Cell Separation (for non-engraftment testing)*

***You MUST call the GENETICS LAB at 513-636-4474 to schedule this test prior to sample submission.**

Non-Hematologic Genetic Analysis

MAP2K1 full gene sequence analysis (Langerhans cell histiocytosis, colon, lung, melanoma) — % **Tumor:** _____

Please see the **Pediatric/Adult Requisition** for
Chromosome Breakage Study for Fanconi Anemia or contact the lab at:
www.cincinnatichildrens.org/cytogenetics or 513-636-4474

SPECIMEN REQUIREMENTS

Cytogenetic Analysis (Chromosome, FISH, and Microarray analysis):

3 mL blood or bone marrow (NaHep)
Chromosome analysis
Cell culture only
FISH probes and FISH panels

3 mL blood or bone marrow (EDTA)
Oncology microarray

Fresh Tumor or Lymph Nodes (1cm³ in sterile saline or sterile transport media)
Chromosome analysis
Cell culture
FISH probes and FISH panels

Molecular Genetic Analysis (RNA Assays): 5-10 mL blood or 3–5 mL bone marrow (EDTA) — Samples must be received within 24 hours of collection.

BCR/ABL — Quantitative (p210), *BCR/ABL* — Quantitative (p190), *BCR/ABL* — Qualitative, *PML/RAR α* — Quantitative

Molecular Genetics Analysis (DNA Assays): 3 mL bone marrow or blood (EDTA) — Samples must be received within 48 hours of collection.

JAK2 Quantitative (V617F), *PTEN* Seq, Bone marrow engraftment by STR, WBC sorted sub-populations engraftment study (by STR or FISH)

Non-Hematologic Genetic Analysis:

MAP2K1 full gene sequencing:

3 mL blood or bone marrow (EDTA), 1 cm³ fresh tumor or 10 formalin fixed paraffin embedded (FFPE) tissue scrolls. Please send additional scrolls (if possible) for extremely small tissue samples.

FISH (Fluorescence In Situ Hybridization)

NOTE: All FISH probes are available for individual testing

Hematologic FISH Panels — 3 mL blood or bone marrow (NaHep)

- ALL Hyperdiploid: trisomy 4, 10, 17
- ALL Risk Stratification: 4, 10, 17, t(1;19), t(12;21), t(9;22), *KMT2A*
- B-Cell ALL: *CRLF2*, t(1;19), *ABL2*, 4/10/17, *CSF1R*, *PDGFRB*, *JAK2*, *ABL1*, t(9;22), *KMT2A*, t(12;21), *IGH*, *EPOR*
- Ph-like ALL: *CRLF2*, *ABL2*, *PDGFRB*, *CSF1R*, *JAK2*, *ABL1*, *EPOR*
- AML: t(6;9), t(8;21), *NUP98*, *KMT2A*, inv(16)
- APL: t(15;17), *RAR α*
- Burkitt Lymphoma: t(8;14), *MYC*
- CLL: 13q14.3, 13q34, 12 centromere, *ATM*, *TP53*, t(11;14)
- Double Hit Lymphoma: *BCL6*, *MYC*, t(8;14), t(14;18)
- Eosinophilia: 4q12, *PDGFRB*, *FGFR1*, *CBFB*
- Fanconi Anemia: 1q25, 3q27, mono 7 / del(7q)
- Multiple Myeloma (CD138+): 1p32.3/1q21, t(4;14), t(11;14), monosomy 13/ del 13q, t(14;16), t(14;20), *TP53*
- MDS: mono 5/del 5q, mono 7/del 7q, tri 8, *TP53*, del (20q)
- Myeloid Malignancy: mono 5/del 5q, t(6;9), mono 7/del 7q, tri 8, t(8;21), *NUP98*, *KMT2A*, *ETV6*, t(15;17), inv(16), *TP53*, del (20q)

- MPD: 4q12, *PDGFRB*, *FGFR1*, *BCR/ABL1*
- SDS: mono 7/del 7q, tri 8, del (20q)
- Large cell NHL: t(11;14), t(14;18), *TP53*, *BCL6*, *ALK*
- Small B-cell NHL: t(11;14), t(14;18), 18q21 (*MALT1*), CLL Panel
- Combined NHL: (large and small cell NHL panels)
- T-Cell Lymphoma: *TRA/TRD*, *TRB*; *TRG*, *BCR/ABL1*, *KMT2A*

Non-Hematologic FISH Panels — 4-8 FFPE slides cut to 4 micron thickness and 1 marked H & E slide[†] — Fresh Tumor (1cm³)

- Ependymoma: *ABL2*, *CDKN2A*, *C11orf95*, *RELA*
- High-Grade Glioma: *PDGFRA*, *CDKN2A*, *NTRK2*, *MYCN*
- Low-Grade Glioma: *TP73/ABL2*, *FGFR1*, *MYB*, *BRAF*, *MYBL1*
- Lung Cancer: *ALK*, *ROS1*, *MET*, *RET*
- Medulloblastoma: *MYB*, *LIS1/RAR α* , *MYC*, *MYCN*
- Melanocytic Tumor: *RREB1*, *MYC*, *CDKN2A*, *CCND1*
- Pilocytic Astrocytoma: *BRAF*, *CDKN2A**

[†]For each probe ordered, send 2 unstained slides with one section cut to 4 micron thickness and mounted on a charged slide. Blocks are also accepted for processing.

*Pilocytic Astrocytoma FISH Panel only needs 2-4 FFPE slides and 1 marked H&E slide

SHIPPING INFORMATION

Local courier is available; please call 513-636-4474 for information.

Shipping for samples that arrive Monday–Saturday:

Cincinnati Children's
Genetics and Genomics Diagnostic Laboratory
3333 Burnet Ave. TCHRf 1042
DOCK 5
Cincinnati, OH 45229-3039

BILLING INFORMATION

* PLEASE NOTE:

- We will not bill Medicaid, Medicaid HMO, or Medicare except for the following: CCHMC Patients, CCHMC Providers, or Designated Regional Counties.
- If you have questions, please call 1-866-450-4198 for complete details.

Patient Signed Completed ABN:

Medical Necessity Regulations: At the government's request, the Molecular Genetics Laboratories would like to remind all physicians that when ordering tests that will be paid under federal health care programs, including Medicare and Medicaid programs, that these programs will pay only for those tests the relevant program deems to be (1) included as covered services, (2) reasonable, (3) medically necessary for the treatment and diagnosis of the patient, and (4) not for screening purposes.