



Instructions: The accurate interpretation and reporting of genetic results is contingent upon the reason for testing, clinical history, family history, and ancestry. To help provide the best possible service, supply the information requested below and **send paperwork with the specimen or return by fax to Mayo Clinic Laboratories, Attn: Molecular Technologies Laboratory Genetic Counselors at 507-284-1759. Phone: 800-533-1710 / International clients: +1-507-266-5700 or email MLIINT@mayo.edu**

Patient Information

| | | |
|---|--|-------------------------|
| Patient Name (Last, First Middle) | | Birth Date (mm-dd-yyyy) |
| Sex Assigned at Birth <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown <input type="checkbox"/> Choose not to disclose | Legal/Administrative Sex <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Nonbinary | |

Referring Provider Information

| | | |
|--|-------|------|
| Referring Provider Name (Last, First) | Phone | Fax* |
| Other Contact/Genetic Counselor Name (Last, First) | Phone | Fax* |

*Fax number given must be from a fax machine that complies with applicable HIPAA regulations.

Reason for Testing

Diagnosis Family history** Other; specify: _____

**Genetic testing should be performed on an affected family member first, when available. FMTT / Familial Variant, Targeted Testing should be ordered when there is a previous positive genetic test result in the family.

Clinical History Check all that apply.

Patient's clinical status: Asymptomatic Symptomatic Other: _____

Indicate whether the following are present:

| | | |
|--|---|---|
| <input type="checkbox"/> Warts | <input type="checkbox"/> Myelodysplastic syndrome (MDS) | <input type="checkbox"/> Viral infections |
| <input type="checkbox"/> Cytopenias (neutropenia, monocytopenia, B cell/NK cell/CD4+ T cell lymphopenia) | <input type="checkbox"/> Mycobacterial disease | <input type="checkbox"/> Pulmonary alveolar proteinosis (PAP) |
| <input type="checkbox"/> Other malignancies; specify: _____ | <input type="checkbox"/> Acute myeloid leukemia (AML) | <input type="checkbox"/> Emberger syndrome |
| <input type="checkbox"/> Lymphedema (eg, legs, genitals); specify: _____ | | |

Preliminary screening results:

WBC: _____; Absolute Neutrophil count: _____; Monocyte count: _____; Absolute Lymphocyte count: _____

B cell count: _____; CD4 T cell count: _____; CD8 T cell count: _____; NK cell count: _____

Dendritic cell count: _____; Dendritic cell phenotyping: _____

Patient treatment history:

No treatment Chemotherapy Allogeneic Hematopoietic cell transplant (blood, BM, cord); transplant date (mm-dd-yyyy): _____

Treatment for infections; specify: _____

Other relevant clinical history: _____

Diagnosis date, if applicable (mm-dd-yyyy): _____

Family History

Are there similarly affected relatives? Yes No
If "Yes," indicate relationship and symptoms: _____

Have any family members had genetic testing? Yes*** No Unknown

*****FMTT / Familial Variant, Targeted Testing should be ordered when there is a previous positive genetic test result in the family. Contact the lab for ordering assistance.**

Ancestry

| | | | | |
|---|-------------------------------------|---|--|---|
| <input type="checkbox"/> African/African American | <input type="checkbox"/> East Asian | <input type="checkbox"/> Latinx/Latine | <input type="checkbox"/> South Asian | <input type="checkbox"/> Choose not to disclose |
| <input type="checkbox"/> Ashkenazi Jewish | <input type="checkbox"/> European | <input type="checkbox"/> Middle Eastern | <input type="checkbox"/> None of the above | <input type="checkbox"/> Unknown |

New York State Patients: Informed Consent for Genetic Testing is required. See Informed Consent for Genetic Testing (T576), or Informed Consent for Genetic Testing – Spanish (T826).