



Instructions: Accurate interpretation and reporting of genetic results is contingent upon the reason for testing, clinical information, 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 800-533-1710. Phone: 507-266-5700 / International clients: +1-507-266-5700 or email MLIINT@mayo.edu**

Patient Information

Patient Name <i>(Last, First, Middle)</i>		Birth Date <i>(mm-dd-yyyy)</i>
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 <i>(Last, First)</i>	Phone	Fax*
Genetic Counselor Name <i>(Last, First)</i>	Phone	Fax*

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

Reason for Testing Specify below or attach relevant clinic note.

Confirm clinical diagnosis; specify diagnosis: _____ Age of onset: _____

Family history**; describe: _____

Other; specify: _____

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

Clinical Presentation

Epstein Barr Virus (EBV) susceptibility Familial hemophagocytic lymphohistiocytosis (F-HLH)

Other viral susceptibility; specify: _____ Other; specify: _____

Lymphoproliferative disorder

Clinical Features Check all that apply.

<input type="checkbox"/> Abnormal bleeding	<input type="checkbox"/> Fulminant viral hepatitis	<input type="checkbox"/> Pityriasis-like lesions
<input type="checkbox"/> Abnormal pigmentation	<input type="checkbox"/> Hemophagocytosis	<input type="checkbox"/> Severe influenza pneumonia
<input type="checkbox"/> Brainstem encephalitis	<input type="checkbox"/> Herpes simplex encephalitis	<input type="checkbox"/> Severe mononucleosis
<input type="checkbox"/> Critical COVID-19 pneumonia	<input type="checkbox"/> Hypogammaglobulinemia	<input type="checkbox"/> Splenomegaly
<input type="checkbox"/> Disseminated intravascular coagulation	<input type="checkbox"/> Live-attenuated viral vaccine strain disease	<input type="checkbox"/> Varicella zoster virus encephalitis and cerebellitis
<input type="checkbox"/> Epidermodyplasia verruciformis	<input type="checkbox"/> Lymphoproliferation	<input type="checkbox"/> Warts
<input type="checkbox"/> Fever	<input type="checkbox"/> Neurological symptoms	<input type="checkbox"/> Other; specify: _____

Oncologic History

<input type="checkbox"/> Myelodysplasia/AML	<input type="checkbox"/> Leukemia; specify: _____
<input type="checkbox"/> Lymphoma; specify: _____	<input type="checkbox"/> Skin cancer; specify: _____
<input type="checkbox"/> Solid tumor; specify: _____	<input type="checkbox"/> Other; specify: _____

Viral Susceptibility, Lymphoproliferation, and Hemophagocytic Lymphohistiocytosis Patient Information (continued)

Patient Treatment History

Has the patient received an allogenic stem cell transplant***? <input type="checkbox"/> No <input type="checkbox"/> Yes; transplant date (mm-dd-yyyy): _____
Is the patient transfusion-dependent***? <input type="checkbox"/> No <input type="checkbox"/> Yes; last transfusion date (mm-dd-yyyy): _____ Was this transfusion leukoreduced***? <input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Unknown
Chemotherapy: <input type="checkbox"/> No <input type="checkbox"/> Yes; date (mm-dd-yyyy): _____
***Results may be inaccurate due to the presence of donor DNA if the patient has received an allogeneic hematopoietic stem cell transplant or non-leukocyte reduced blood products. Call Mayo Clinic Laboratories for instructions for testing patients who have received a bone marrow transplant.

General History

<input type="checkbox"/> Anemia (Hemoglobin < 9 g/dL; neonates < 10 g/dL)	<input type="checkbox"/> Hyperferritinemia (≥ 500 mg/nL; ≥ 500 μ g/L)
<input type="checkbox"/> Thrombocytopenia (Platelets < 100×10^9 /L)	<input type="checkbox"/> Reduced or absent NK-cell cytotoxicity
<input type="checkbox"/> Neutropenia (Neutrophils < 1×10^9 /L)	<input type="checkbox"/> Elevated soluble CD25 (soluble IL-2 receptor)
<input type="checkbox"/> Hypertriglyceridemia (≥ 265 mg/dL; ≥ 3 mmol/L)	<input type="checkbox"/> Viral infection; specify: _____
<input type="checkbox"/> Hypofibrinogenemia (≤ 150 mg/dL; ≤ 1.5 g/L)	<input type="checkbox"/> Other infections; specify: _____

Family History

Are there similarly affected relatives? <input type="checkbox"/> Yes <input type="checkbox"/> No If "Yes," indicate relationship, and diagnosis or symptoms: _____
Have any family members had genetic testing? <input type="checkbox"/> Yes**** <input type="checkbox"/> No <input type="checkbox"/> Unknown ****FMTT / Familial Mutation, Targeted Testing should be ordered when there is a previous positive genetic test result in the family. Contact the lab for ordering assistance.
History of consanguinity: <input type="checkbox"/> No <input type="checkbox"/> Yes; relationship details: _____

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"/> Unknown
<input type="checkbox"/> Ashkenazi Jewish	<input type="checkbox"/> European	<input type="checkbox"/> Middle Eastern	<input type="checkbox"/> None of the above	<input type="checkbox"/> Choose not to disclose

New York State patients: Informed Consent for Genetic Testing is required. See Informed Consent for Genetic Testing (T576), Informed Consent for Genetic Testing – Spanish (T826), or Informed Consent for Genetic Testing for Deceased Individuals (T782).