



**Instructions:** Accurate interpretation and reporting of the genetic results is contingent upon the reason for referral, clinical information, ethnic background, and family history information provided. Supply the information requested below and **send a completed copy of this form with the specimen.**

**Patient Information** (required)

Patient Name <i>(Last, First, Middle)</i>	Birth Date <i>(mm-dd-yyyy)</i>	Sex <input type="checkbox"/> Male <input type="checkbox"/> Female
Referring Provider Name <i>(Last, First)</i>	Phone	Fax*
Other Contact Name <i>(Last, First)</i>	Phone	Fax*

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

**Reason for Testing and Clinical History** (check all that apply)

<b>Treatment History</b>	Date Started <i>(mm-dd-yyyy)</i>
Previous diagnosis of X-linked Agammaglobulinemia (XLA)? <input type="checkbox"/> Yes <input type="checkbox"/> No	Date Last Received <i>(mm-dd-yyyy)</i>
Immunoglobulin treatment? <input type="checkbox"/> Yes <input type="checkbox"/> No	

**Pertinent Clinical and Laboratory History**

Hypogammaglobulinemia (low IgG, IgM, IgA) <input type="checkbox"/> Yes <input type="checkbox"/> No	Sinusitis <input type="checkbox"/> Yes <input type="checkbox"/> No
Common Variable Immunodeficiency (CVID) <input type="checkbox"/> Yes <input type="checkbox"/> No	Tonsils present <input type="checkbox"/> Yes <input type="checkbox"/> No
Recurrent infections <input type="checkbox"/> Yes <input type="checkbox"/> No	Lymph nodes present <input type="checkbox"/> Yes <input type="checkbox"/> No
Pneumonia <input type="checkbox"/> Yes <input type="checkbox"/> No	Splenomegaly <input type="checkbox"/> Yes <input type="checkbox"/> No
CD19+ B-cells present in blood (>1%) <input type="checkbox"/> Yes <input type="checkbox"/> No	
Btk protein by flow cytometry <input type="checkbox"/> Present <input type="checkbox"/> Absent <input type="checkbox"/> Equivocal <input type="checkbox"/> Carrier <input type="checkbox"/> Unknown	

Other Diagnosis

Other Information (such as allogeneic stem cell transplant; indicate type [myeloablative vs. non-myeloablative] and date)

**Ethnic Background** (Ethnic background may assist with interpretation of test results.)

- European/Caucasian, list countries of origin: \_\_\_\_\_  
 African American  Hispanic  Asian  Other, specify: \_\_\_\_\_

**Family History**

Normal <input type="checkbox"/> Father <input type="checkbox"/> Mother <input type="checkbox"/> Siblings
Hypogammaglobulinemia (low IgG and/or IgM, IgA) <input type="checkbox"/> Father <input type="checkbox"/> Mother <input type="checkbox"/> Siblings
CVID <input type="checkbox"/> Father <input type="checkbox"/> Mother <input type="checkbox"/> Siblings
Recurrent infections <input type="checkbox"/> Father <input type="checkbox"/> Mother <input type="checkbox"/> Siblings
Are other male relatives known to be affected? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, indicate their relationship to the patient:
Are other female relatives known to be a carrier? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, indicate their relationship to the patient:
Have other relatives had molecular genetics testing? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, indicate their relationship to the patient:
If the relative was tested at Mayo Clinic, include the name of the family member: