



The accurate interpretation and reporting of genetic results is contingent upon the reason for testing, clinical information, and family history. To help provide the best possible service, supply the information requested below and **send this paperwork with the specimen or return by fax to 507-284-1759.**

†Contact the Special Coagulation DNA Laboratory at 800-533-1710 with questions (International Clients +1-507-266-5700 or mclglobal@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*
Other Contact 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 Check one.

Patient has a diagnosis or suspected diagnosis of hemophilia A and you would like to identify the underlying mutation.
 Patient has a family history of hemophilia A.
 Patient is a known or suspected carrier for hemophilia A, and the mutation in the family has not been previously identified. If familial mutation has been identified, indicate it in the F8 Known Mutation box.

F8 Known Mutation (if applicable)

If a known variant is ordered, the following information **MUST** be provided or testing cannot be completed. Known familial variant:
 Intron 1 Inversion Intron 22 Inversion
 Other: _____
 Proband's relationship to this patient: _____

Clinical Information

Factor 8 Coagulant Activity Undetermined or unavailable 1%–5% of normal (moderately affected†)
 Less than 1% of normal (severely affected) More than 5% of normal (mildly affected†)
 Indicate any other relevant clinical information: _____

Pregnancy Information

Is patient or partner currently pregnant? Yes No If Yes, weeks gestation: _____
 Prenatal specimen? Yes No If Yes, specify specimen type: Chorionic villus sampling Amniotic fluid
 Cord blood specimen? Yes No

Family History

Are there relatives known to be affected or to be a carrier of hemophilia A? Yes No Unknown
 If Yes, indicate relationship (including degree) to patient or attach pedigree: _____
 Have other relatives had molecular genetic testing for hemophilia A? Yes No Unknown
 If Yes, provide results and attach a copy of the genetic test lab report, if available: _____
 If the relative was tested at Mayo Clinic, include the following information about the family member:
 Name *(Last, First, Middle)* _____ Birth Date *(mm-dd-yyyy)* _____

Affiliation

Hemophilia Center Affiliation Yes No If Yes, which center: _____