



**Instructions:** The accurate interpretation and reporting of the genetic results is contingent upon the reason for referral, clinical information, ethnic background, and family history. 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: Personalized Genomics Laboratory Genetic Counselors at 507-284-1759. Phone: 507-266-5700 / International clients: +1-507-266-5700 or email [mclglobal@mayo.edu](mailto:mclglobal@mayo.edu)**

**Patient Information**

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 given must be from a fax machine that complies with applicable HIPAA regulation.

**Reason for Testing** Select one or more of the following indications for testing.

<input type="checkbox"/> Diagnosis/suspected diagnosis <input type="checkbox"/> Gilbert syndrome <input type="checkbox"/> Crigler-Najjar syndrome <input type="checkbox"/> Type I <input type="checkbox"/> Type II <input type="checkbox"/> Other, specify: _____	<input type="checkbox"/> Carrier testing	<input type="checkbox"/> Pharmacogenomic screening <input type="checkbox"/> Irinotecan <input type="checkbox"/> Atazanavir <input type="checkbox"/> Belinostat <input type="checkbox"/> Nilotinib <input type="checkbox"/> Pazopanib <input type="checkbox"/> Other, specify: _____
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**Clinical History**

<b>Laboratory Data</b> Date last bilirubin was performed <i>(mm-dd-yyyy)</i> : _____ Total bilirubin: _____ mg/dL Conjugated (direct) bilirubin: _____ mg/dL Older significant bilirubin levels: _____ Symptoms <input type="checkbox"/> Hyperbilirubinemia <input type="checkbox"/> Other, specify: _____	History of suspected UGT1A1-associated drug toxicity or adverse drug reaction? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide: <input type="checkbox"/> Irinotecan <input type="checkbox"/> Belinostat <input type="checkbox"/> Atazanavir <input type="checkbox"/> Nilotinib <input type="checkbox"/> Pazopanib <input type="checkbox"/> Other, specify: _____ Reaction/Symptoms <input type="checkbox"/> Neutropenia <input type="checkbox"/> Diarrhea <input type="checkbox"/> Other, specify: _____
Additional Comments	

**Family History and Ethnic Background**

<b>Family History</b> Other relatives known to be affected? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, indicate their relationship to the patient: _____ Have other relatives had UGT1A1 molecular genetic testing? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, indicate: Performing laboratory for relative's testing: _____ UGT1A1 variants detected in relative: _____ If relative was tested at Mayo Clinic, name and relationship of relative: _____ <p style="text-align: center;"><b>Attach a copy of the genetic test lab report, if available.</b></p> <b>Ethnic Background</b> <input type="checkbox"/> European/Caucasian <input type="checkbox"/> African American <input type="checkbox"/> Hispanic/Latino <input type="checkbox"/> Asian <input type="checkbox"/> Middle Eastern <input type="checkbox"/> Other, specify: _____ Indicate countries of origin: _____
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