MAYO CLINIC
Mayo Medical Laboratories

Laboratory Service Report 1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT,UGT2O	SA00055149	46	F	SA00055149
Ordering Phys UNKNOWN,PROVIDER				**DOB** 06/10/1966
Client Order # SA00055149	**Account Information**			**Report Notes**
Collected 03/15/2013 01:21	C7028846-DLMP Rochester 3050 Superior Drive Rochester, MN 55901			
Printed 05/28/2013 14:10				

Test	Flag	Results	Unit	Reference Value	Perform Site*
UGT1A1 Sequence, Hyperbili, Saliva			REPORTED 05/28/2013 13:41		
UGT1A1 Sequence, Hyperbilirubinemia					
UGT1A1 Hyperbilirubinemia Result					MCR
A mutation was NOT detected in the UGT1A1 gene.					
Reviewed by		Jamie Bruflat			MCR
UGT1A1 Hyperbilirubinemia Interp					MCR
Both copies of the UGT1A1 gene have the normal TA6 (*1) promoter repeat and no additional UGT1A1 mutations were identified, which is consistent with normal activity of the UGT1A1 enzyme and would not be expected to cause marked unconjugated hyperbilirubinemia. Bidirectional DNA sequence analysis was used to test for the presence of mutations in the promoter, exons, exon-intron boundaries, and 3'-untranslated region of the UGT1A1 gene that are associated with the diagnosis of unconjugated hyperbilirubinemia. A small percentage of individuals who have a diagnosis of unconjugated hyperbilirubinemia may have a mutation that is not identified by the methods described above. The presence of UGT1A1 mutations does not necessarily confirm a diagnosis of unconjugated hyperbilirubinemia. Breast-fed neonates may experience a physiologic unconjugated hyperbilirubinemia and jaundice from deconjugation of maternal bilirubin-glucuronides present in breast milk. Clinical correlation is recommended.					
A genetic consultation may be of benefit.					
A list of common polymorphisms identified for this patient is available from the laboratory upon request.					
CAUTIONS: Rare polymorphisms exist that could lead to false negative or positive results. Test results should be interpreted in the context of clinical findings, family history, and other laboratory data. Large deletions or rearrangements are not detected by this assay, and these may affect UGT1A1 protein expression, and the ability to conjugate bilirubin. Laboratory developed test.					
UGT, Full Gene Sequencing		Performed			MCR

* Performing Site:

MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester, MN 55905	Lab Director: Franklin R. Cockerill, III, M.D.

* Report times for Mayo performed tests are CST/CDT