

Bile Acid Synthesis Disorders

Atypical Bile Acid Test

REPORT TO	PATIENT INFORMATION
Physician Name (print):	THE FOLLOWING INFORMATION IS REQUIRED FOR EACH SAMPLE
Clinic/Institution Name:	Patient Name:,,,
Address	Last First M
Address:	,
City: State: Zip:	Patient ID/Med Rec #:
Phone: ()	Address:
Email:	City: State: Zip:
NPI #	Preferred Phone: ()
NPI#	Other # Where Patient can be Reached: ()
FAX NUMBER FOR RESULTS:	_ Sex: ☐ Male ☐ Female ☐ Unknown
The Laboratory DOES NOT bill patients or insurance companies This is a program supported by Travere Therapeutics , Inc.	Parent Name (if patient is minor):
SAMPLE/SPECIMEN INFORMATION	Spouse:
Sample Type: Urine (1 – 25 mL)	Ethnicity of Patient (check all that apply):
Sample Collection Date (MM/DD/YYYY)://	□African American □ Asian □ Caucasian NW European □ E Indian
Internal Use only:	☐ Hispanic ☐ Ashkenazi Jewish ☐ Sephardic Jewish ☐ Mediterranean
Received date:	☐ Native American ☐ NativeHawaiian/Other Pacific Islander ☐ Other
FL#:	Because Ursodeoxycholic acid can mask detection of bile acid synthetic
FAB#:	disorders, the patient should be temporarily taken off URSO® or ACTIGALL®
SHIPPING INFORMATION	(ursodiol) for 5 DAYS before sample collection.
Shipment Requirements:	List Medications:
• US SHIPMENTS ONLY	Is the patient currently on URSO® or ACTIGALL® (ursodiol), or has been within
• SHIP FROZEN	the past month? If yes, please provide the DATES of medication:
- ON ICE PACKS OR - DRY ICE	
• OVERNIGHT EXPRESS	Clinical History/Preliminary Diagnosis:
- NO WEEKEND DELIVERY	
Ship to:	
Clinical Mass Spectrometry Facility, MLC 7019	
Department of Pathology and Laboratory Medicine	
Cincinnati Children's Hospital Medical Center	
240 Albert Sabin Way Cincinnati, OH 45229-3039	ICD-10:
Phone: (513) 636-4203 Fax: (513) 803-5014	

CRITERIA FOR FREE TESTING

Please check boxes and attest:
Patient must meet one of the following:
☐ Pathogenic Variant or VOUS from a genetic test on one of the following general PSD3B7, AKR1D1, AMACR, CYP7B1
☐ Negative result on genetic test but patient has GGT≤150 IU/L and direct

bilirubin >1 mg/dL

I hereby attest that the patient meets the attached criteria and is a candidate for the Atypical Bile Acid Test via FAB-MS. I understand the diagnostic testing services offered under this program are directional in nature and that they do not eliminate the need for additional medical management or replace any existing diagnostic methods. I further understand that neither Travere Therapeutics, Inc. nor Cincinnati Children's Hospital makes any claims as to the usefulness of this test.

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Signature: _

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