Fabry Disease: Globotriaosylceramide (Gb3) Quantification, Urine

Test Code: BGBCU
Turnaround time: 7 days - 10 days
CPT Codes: 82542 x1, 82570 x1

Notice: This test will soon be discontinued.
As of 02/01/2020, EGL can no longer accept samples for this test. For questions, please call: 470-378-2200.

Fabry disease, the only X-linked sphingolipidosis, is associated with severe multiorgan dysfunction. Its incidence has been estimated from 1 in 40,000 to 60,000 live births for males. Heterozygous females can be symptomatic. Although clinical onset often occurs in childhood, disease presentation may be subtle, leading to delayed diagnosis or misdiagnosis. The primary defect is a deficiency of the lysosomal enzyme, alpha-galactosidase A which releases galactose from ceramide trihexoside (globotriaosylceramide, Gb3) and related glycosphingolipids (especially galabiosylceramide, Gb2), due to mutations in the GLA gene. This results in progressive accumulation of Gb3 in vascular endothelial cells, epithelial and smooth muscle cells, leading to ischemia and infarction especially in the kidney, heart and brain. Large amounts of Gb3 are excreted by untreated male hemizygotes (except patients with a renal graft and those with a cardiac variant), and smaller but still significant amounts by heterozygote females, symptomatic or not. At present, urinary Gb3 quantification also provides additional diagnostic information for manifesting female carriers of Fabry disease, as well as patients with multiple glycolipid storage conditions. Measurement of Gb3 has also been used by some groups to monitor enzyme replacement therapy and disease progression.1,2


References:

Genes
GLA

Indications
This test is indicated for:
- Males who have clinical symptoms of Fabry disease
- Females, both symptomatic and asymptomatic, who may be carriers of Fabry disease
- Males and females on enzyme replacement therapy (ERT) for Fabry disease

Methodology
LC-MS/MS

Detection
- Fabry patients not on treatment may have elevated Gb3 levels.
- Fabry patients on treatment have significantly lower to normal Gb3 levels.
- Mildly elevated Gb3 levels may also be caused by other renal diseases or injury to the urinary tract.
- This assay may be affected by urine that is stored at room temperature for a prolonged period of time.

Reference Range
1-4 years old: 0.0-17 ug/mmol Cr
>4 years old: 0.0-8.0 ug/mmol Cr

Specimen Requirements
Submit only 1 of the following specimen types

Type: Urine

Specimen Requirements:
Clean container without additives
1-5 ml
Freeze sample. Fasting or first void sample is preferable.

Disclaimer: This information is confidential and subject to change without notice. It may not be reproduced in whole or part unless authorized in writing by an authorized EGL representative.
Ship frozen sample on dry ice with overnight delivery.

**Special Instructions**

Clinical information is required for complete interpretation.

**Related Tests**

**Biochemical:**

- Fabry Disease: Alpha-Galactosidase A Enzyme Activity (LB)

**Molecular:**

- Fabry Disease: GLA Full Gene Sequencing (DG)
- Fabry Disease: GLA Gene Deletion/Duplication (KX)