

**Reporting Title:** Oligosaccharide Screen, U  
**Performing Location:** Rochester

**Ordering Guidance:**  
This is the recommended test when clinical features are suggestive of, or when molecular testing results suggest, an oligosaccharidosis disorder that can be identified by this test.

The recommended screening test for the initial workup of a suspected lysosomal storage disorder, particularly when clinical features are nonspecific, is LSDS / Lysosomal Storage Disorders Screen, Random, Urine.

**Necessary Information:**  
1. Patient's age is required.  
2. [Biochemical Genetics Patient Information](#) (T602) is recommended. This information aids in providing a more thorough interpretation of results. Send information with specimen.

**Specimen Requirements:**  
**Supplies:** Urine Tubes, 10 mL (T068)  
**Container/Tube:** Plastic, 10-mL urine tube  
**Specimen Volume:** 8 mL  
**Pediatric Volume:** 2 mL  
**Collection Instructions:**

- 1. Collect a random urine specimen.
- 2. No preservative
- 3. Immediately freeze specimen.

**Forms:**  
1. [Biochemical Genetics Patient Information](#) (T602)  
2. If not ordering electronically, complete, print, and send a [Biochemical Genetics Test Request](#) (T798) with the specimen.

Specimen Type	Temperature	Time	Special Container
Urine	Frozen (preferred)	365 days	
	Refrigerated	15 days	
	Ambient	7 days	

**Result Codes:**

Result ID	Reporting Name	Type	Unit	LOINC®
64889	Oligosaccharide Screen, U	Alphanumeric		49284-3

LOINC® and CPT codes are provided by the performing laboratory.

**Supplemental Report:**  
No

**CPT Code Information:**

84377

**Reference Values:**

An interpretive report will be provided.