

# PRIOR AUTHORIZATION CRITERIA

**BRAND NAME**  
(generic)

**SUCRAID**  
(sacrosidase)

**Status: CVS Caremark Criteria**

**Type: Initial Prior Authorization with Quantity Limit**

## POLICY

### FDA-APPROVED INDICATIONS

Sucraid is indicated as oral replacement therapy of the genetically determined sucrase deficiency, which is part of congenital sucrase-isomaltase deficiency.

### COVERAGE CRITERIA

The requested drug will be covered with prior authorization when the following criteria are met:

- The patient has a diagnosis of congenital sucrase-isomaltase deficiency
- AND**
- The diagnosis of congenital sucrase-isomaltase deficiency was confirmed by small bowel biopsy
- OR**
- The diagnosis of congenital sucrase-isomaltase deficiency was confirmed by genetic testing
- OR**
- The diagnosis of congenital sucrase-isomaltase deficiency was confirmed by sucrose hydrogen breath test

Quantity Limits apply.

3 bottles (354 mL) per month

### REFERENCES

1. Sucraid [package insert]. Vero Beach, FL: QOL Medical, LLC; September 2018.
2. Lexicomp Online, AHFS DI (Adult and Pediatric) Online. Hudson, OH: Wolters Kluwer Clinical Drug Information, Inc. <http://online.lexi.com/>. Accessed August 2020.
3. Micromedex (electronic version). Truven Health Analytics, Greenwood Village, Colorado, USA. <http://www.micromedexsolutions.com/>. Accessed August 2020.
4. National Organization for Rare Disorders (NORD). Congenital Sucrase-Isomaltase Deficiency. 2005. Available at <https://rarediseases.org>. Accessed August 2020.
5. Genetic and Rare Diseases Information Center. Congenital sucrose-isomaltase deficiency. 2020. Available at <https://rarediseases.info.nih.gov>. Accessed August 2020.