# PRIOR AUTHORIZATION CRITERIA

# **BRAND NAME** (generic)

**SUCRAID** (sacrosidase)

Status: CVS Caremark® Criteria

Type: Initial Prior Authorization with Quantity Limit

## **POLICY**

# **FDA-APPROVED INDICATIONS**

Sucraid (sacrosidase) Oral Solution is indicated as oral replacement therapy of the genetically determined sucrase deficiency, which is part of congenital sucrase-isomaltase deficiency (CSID).

#### **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
  - 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

Quantity Limits apply.

#### **QUANTITY LIMIT**

PLEASE NOTE: Since manufacturer package sizes may vary, it is the discretion of the dispensing pharmacy to fill quantities per package size up to these quantity limits. In such cases the filling limit and day supply may be less than what is indicated.

1 Month Limit\* 3 Month Limit\* Drug

Sucraid Multiple-Dose Bottle

(Each bottle contains 4 oz [118 mL total]) 354 mL / 25 days 1062 mL / 75 days

Sucraid Single-Use Container

(Each carton contains 150 single-use containers of 2 900 mL/ 63 days

300 mL/ 21 days mL each [300 mL total])

\*The duration of 25 days is used for a 30-day fill period and 75 days is used for a 90-day fill period to allow time for refill processing OR the duration of 21 days is used for a 25-day fill period and 63 days is used for a 75-day fill period to allow time for refill processing.

Duration of Approval (DOA):

3369-C: DOA: 12 months

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### **REFERENCES**

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- 2. Lexicomp Online, AHFS DI (Adult and Pediatric) Online. Waltham, MA: UpToDate, Inc.; 2023. https://online.lexi.com. Accessed August 23, 2023.
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- 4. National Organization for Rare Disorders (NORD). Congenital Sucrase-Isomaltase Deficiency. 2005. Available at https://rarediseases.org/rare-diseases/disaccharide-intolerance-i/. Accessed August 23, 2023.

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