

# SENTARA HEALTH PLANS

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

**Directions:** The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-800-750-9692.** No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If the information provided is not complete, correct, or legible, the authorization process can be delayed.**

**Drug Requested:** Sucraid® (sacrosidase)

**MEMBER & PRESCRIBER INFORMATION:** Authorization may be delayed if incomplete.

Member Name: \_\_\_\_\_

Member Sentara #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

**DRUG INFORMATION:** Authorization may be delayed if incomplete.

Drug Form/Strength: \_\_\_\_\_

Dosing Schedule: \_\_\_\_\_ Length of Therapy: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code, if applicable: \_\_\_\_\_

Weight: \_\_\_\_\_ Date: \_\_\_\_\_

**Quantity Limit:** 236mL/30 days

**CLINICAL CRITERIA:** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

**Initial Approval: 60 days**

- Patient is 5 months of age or older and has a diagnosis of congenital sucrase-isomaltase deficiency (CSID) confirmed by a gastroenterologist, endocrinologist, or genetics specialist

**AND**

- Patient has documented chronic symptoms of CSID including watery diarrhea, abdominal pain, gas/bloating after sucrose/starch ingestion (must submit chart notes documenting symptoms following sucrose/starch ingestion)

- Number of severe GI events within the last 2 months: \_\_\_\_\_ (must be documented in submitted chart notes)

**AND**

(Continued on next page)

- ❑ A low sucrose and low starch diet has been attempted with improvement in patient symptoms, and patient will continue to follow a low sucrose, low starch diet while on therapy

**AND**

- ❑ Patient does not have lactose intolerance or a secondary sucrose deficiency associated with any of the following: celiac disease, Crohn's disease, autoimmune gastroenteropathy, eosinophilic gastroenteropathy, short bowel syndrome, Giardiasis, small intestinal bacterial overgrowth (SIBO), acute gastroenteritis, or enteropathy associated with acquired immune deficiency syndrome

**AND (ALL 4 below MUST be met):**

<input type="checkbox"/> Stool pH < 6.0	<input type="checkbox"/> Increase in breath hydrogen of > 10 ppm when challenged with sucrose after fasting
<input type="checkbox"/> Genetic test results confirm diagnosis of CSID	<input type="checkbox"/> Negative lactose breath test

**OR (BOTH below MUST be met)**

- ❑ Small bowel biopsy documents intestinal sucrose activity of <25 U/g protein (must be greater than 2 standard deviations below the mean) with normal or decreased maltase and isomaltase levels, normal levels of other disaccharides, and normal villous architecture of the small intestine on biopsy
- ❑ Genetic testing results document sucrose-isomaltase deficiency (CSID)

**Reauthorization Approval: 12 months.** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

- ❑ Patient has had a 50% reduction in all symptoms of CSID, including watery diarrhea, abdominal pain, gas/bloating; etc. (**improvement from baseline must be noted in submitted chart notes**)
  - Number of severe GI events within the last 2 months: \_\_\_\_\_ (**must be documented in submitted chart notes**)

**AND**

- ❑ Patient will continue to follow a low sucrose, low starch diet while on therapy.

**Medication being provided by Specialty Pharmacy - PropriumRx**

*Not all drugs may be covered under every Plan*

*If a drug is non-formulary on a Plan, documentation of medical necessity will be required.*

*\*\*Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.\*\**

*\*Previous therapies will be verified through pharmacy paid claims or submitted chart notes.\**