

# 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:** Vykat™ XR (diazoxide choline)

**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:** \_\_\_\_\_

**NPI #:** \_\_\_\_\_

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

**Drug Name/Form/Strength:** \_\_\_\_\_

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

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

**Weight (if applicable):** \_\_\_\_\_ **Date weight obtained:** \_\_\_\_\_

### **Recommended Dosing:**

Weight	Starting Dosage	Titration Dosage		Target Maintenance Dosage
	Weeks 1 and 2	Weeks 3 and 4	Weeks 5 and 6	
20 to < 30 kg	25 mg	50 mg	75 mg	100 mg
30 to < 40 kg	75 mg	150 mg	150 mg	150 mg
40 to < 65 kg	75 mg	150 mg	225 mg	225 mg
65 to < 100 kg	150 mg	225 mg	300 mg	375 mg
100 to < 135 kg	150 mg	300 mg	375 mg	450 mg
≥ 135 kg	150 mg	300 mg	450 mg	525 mg

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**Quantity Limits:**

- 25 mg – 4 tablets per day
- 75 mg – 7 tablets per day
- 150 mg – 3 tablets per day

**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 Authorization: 12 months**

- ☐ Member is > 4 years of age
- ☐ Member weighs > 20 kg
- ☐ Prescribed by or in consultation with a geneticist, endocrinologist, metabolic disorder subspecialist, neurologist, or a physician who specializes in the treatment of Prader-Willi syndrome
- ☐ Member has a confirmed diagnosis of Prader-Willi syndrome as established by a genetic test showing **ONE** of the following (**submit documentation**):
  - ☐ Deletion 15q11-q13 or Maternal disomy
  - ☐ Imprinting genes on chromosome 15
- ☐ Member has moderate to severe hyperphagia
- ☐ Provider must submit **ONE** of the following baseline assessments (**submit documentation**):
  - ☐ Hyperphagia questionnaire
  - ☐ Clinical Global Impression of Improvement (CGI-I)
  - ☐ Caregiver Global Impression of Change (GI-C)
  - ☐ Body fat measured

**Reauthorization: 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.

- ☐ Member has experienced a positive clinical response to Vykath<sup>TM</sup> XR therapy (e.g., improved behavior, metabolism improvement, reduced hyperphagia)

**Medication being provided by Specialty Pharmacy – Proprium Rx**

***\*\*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.\****