

# 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; **(Pharmacy) 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:** (Select drug below)

<input type="checkbox"/> <b>deferasirox (Exjade®) tablets for oral suspension</b>	<input type="checkbox"/> <b>deferasirox (Jadenu®) tablets</b>
<input type="checkbox"/> <b>deferasirox (Jadenu® sprinkle) packet</b>	

**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 Limits:

**deferasirox (Exjade):** Maximum of 40 mg/kg/day.

**deferasirox (Jadenu):** Maximum of 28 mg/kg/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.

**For diagnosis of transfusional iron overload (transfusional hemosiderosis), ALL of the following criteria must be met for initial 6 month approval:**

Member is  $\geq 2$  years of age and has a diagnosis of transfusional hemosiderosis (i.e., transfusion of  $\geq 100$  mL/kg of packed red blood cells, approximately 20 units for a 40kg patient)

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- Member's serum ferritin levels are consistently >1,000mcg/L (submit serum ferritin labs done within the last 30 days)
- Member's liver iron concentration (LIC) is >5mg of Fe/g of dry weight (submit liver biopsy, MRI or other FDA-approved test to document LIC [Fe/g of dry weight])
- Member's current weight must be noted: \_\_\_\_\_
- Member has an eGFR  $\geq$ 40mL/min/1.73m<sup>2</sup> (submit renal function labs)
- Member's baseline liver function labs must be submitted (submit ALT, AST, bilirubin)
- If requesting brand Jadenu or Exjade, documentation of trial and intolerable life-endangering adverse event with generic deferaxirox must be submitted

For diagnosis of non-transfusion-dependent thalassemia syndrome, **ALL** of the following criteria must be met **for initial 6-month approval**:

- Member is  $\geq$ 10 years of age and has a diagnosis of non-transfusion-dependent thalassemia syndrome
- Liver iron concentration (LIC) is  $\geq$ 5mg of Fe/g of dry weight (submit current liver biopsy, MRI or other FDA-approved test to document LIC)
- Serum ferritin is >300mcg/L (submit 2 serum ferritin labs, taken at least 1 month apart, from within the last 3 months)
- Member's current weight must be noted: \_\_\_\_\_
- Member has an eGFR  $\geq$ 40mL/min/1.73m<sup>2</sup> (submit renal function labs)
- Member's platelets levels are  $\geq$ 50 x 10<sup>9</sup>/L
- Member's baseline liver function labs must be submitted (submit ALT, AST, total bilirubin)
- If requesting brand Jadenu or Exjade, documentation of trial and intolerable life-endangering adverse event with generic deferaxirox must be submitted

**For 12 month reauthorization, ALL** of the following criteria must be met:

- Serum ferritin has decreased from baseline or last approval (submit current ferritin labs)
- Liver iron concentration (LIC) has decreased to < 5mg of Fe/g of dry weight or has been maintained at a level that is 30% lower than baseline level (submit liver biopsy, MRI or other FDA-approved test to document current LIC)
- If serum ferritin is <500 mcg/L or LIC is less than 3 mg Fe/g dw, deferaxirox therapy will be temporarily discontinued; if <300 mcg/L, deferaxirox therapy will be interrupted and LIC obtained
- Member's liver function continues to be monitored (submit current ALT, AST, bilirubin labs)
- Member's platelets levels are  $\geq$ 50 x 10<sup>9</sup>/L
- If requesting brand Jadenu or Exjade, documentation of trial and intolerable life-endangering adverse event with generic deferaxirox must be submitted

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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.\****