

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 may be delayed.**

Drug Requested: (Select drug below)

<input type="checkbox"/> deferiprone (Ferriprox®) tablets	<input type="checkbox"/> Ferriprox® (deferiprone) solution	<input type="checkbox"/> Ferriprox® 2-day (deferiprone) tablets
---	--	---

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: Maximum 99 mg/kg/day (actual body weight) in two divided doses

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. **Check the diagnosis below that applies.**

Diagnosis: Transfusional hemosiderosis due to thalassemia syndrome

Initial Authorization: 6 months

- Member is 3 years of age or older
- Member has a diagnosis of transfusional hemosiderosis due to thalassemia syndrome (i.e., transfusion of ≥ 100 mL/kg of packed red blood cells, approximately 20 units for a 40 kg patient)

(Continued on next page)

- Member's serum ferritin levels are consistently >1,000 mcg/L (**submit serum ferritin labs done within the last 30 days**)
- Member's current weight: _____
- Member is currently established on **ONE** of the following chelator therapies for at least 90 consecutive days (**verified by pharmacy paid claims**):
 - deferoxamine (Desferal)
 - deferasirox (Exjade, Jadenu)
- Member continues to have **ONE** of the following after three (3) months of chelator therapy:
 - Serum ferritin in excess of 2,500 mcg/L
 - Liver iron concentration is >7 mg Fe/g dry weight (**submit liver biopsy, MRI or other FDA-approved test results**)
 - Cardiac magnetic resonance imaging (MRI T2*) is ≤ 10 ms (**submit MRI T2* lab results**)
- Baseline absolute neutrophil count (ANC) is $> 1.5 \times 10^9/L$ and ANC will continue to be monitored weekly while on therapy (**submit current labs**)
- If requesting brand Ferriprox, documentation of trial and intolerable life-endangering adverse event with generic deferiprone must be submitted
- Ferriprox solution may be approved for members aged 3-10 years only. If requesting Ferriprox solution for members ≥ 11 years of age, documentation that member is unable to ingest any solid dosage form must be submitted

Diagnosis: Transfusional hemosiderosis due to thalassemia syndrome

Reauthorization: 12 months.

- Member's ANC is $> 1.5 \times 10^9/L$ (**submit current lab results**)
- Liver iron concentration is ≤ 5 mg of Fe/g of dry weight (**submit current liver biopsy, MRI or other FDA-approved test results**)
- Treatment will be withheld if serum ferritin falls consistently below 500 mcg/L
- Serum ferritin has decreased by $\geq 20\%$ from baseline or has been maintained at a level that is $\geq 20\%$ below baseline level (**submit current serum ferritin labs**)

Diagnosis: Transfusional iron overload in members with sickle cell disease or other anemias

Initial Authorization: 12 months

- Member is 3 years of age or older
- Member has a diagnosis of transfusional iron overload associated with sickle cell disease or other anemia diagnosis
- Baseline liver iron concentration > 7 mg of Fe/g dry weight (**submit current MRI results**)

(Continued on next page)

- ❑ Member has received no less than 20 transfusions of RBCs
- ❑ Baseline absolute neutrophil count (ANC) is $> 1.5 \times 10^9/L$ and ANC will continue to be monitored weekly while on therapy (**submit current labs**)

❑ **Diagnosis: Transfusional iron overload in members with sickle cell disease or other anemias**

Reauthorization: 12 months.

- ❑ Liver iron concentration has decreased by at least 4 mg of Fe/g dry weight from baseline or has been maintained at a level that is at least 4 mg of Fe/g dry weight below baseline level since last approval (**submit current MRI results**)
- ❑ Member's ANC is $> 1.5 \times 10^9/L$ (**submit current lab results**)
- ❑ Treatment will be withheld if serum ferritin falls consistently below 500 mcg/L

Medication being provided by a 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.****