

## Ferriprox [deferiprone]

## **Prior Authorization Request**

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-866-814-5506**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name:Patient's ID:		Date:Patient's Date of Birth:
Sp	ecialty:	NPI#:
	ysician Office Telephone:	Physician Office Fax:
Re	equest Initiated For:	
1.	What is the prescribed drug? $\Box$ Ferriprox $\Box$ defe	eriprone
2.	What is the diagnosis?  ☐ Transfusional iron overload due to a thalassemia syndrome ☐ Transfusional iron overload due to sickle cell disease or other anemias ☐ Hereditary hemochromatosis ☐ Other	
3.	What is the ICD-10 code?	
Co	emplete the following sections based on the prescribe	ed product, if applicable.
Sec	ction A: Ferriprox (Brand) Requests	
	Is the product being requested for the treatment of	chronic iron overload?
	$\square$ Yes $\square$ No If No, skip to section B.	
5.	The preferred products for your patient's health plan are generic deferasirox, deferiprone, and deferoxamine. Can the patient's treatment be switched to a preferred product? <i>If deferasirox or deferoxamine, please call 1-866-814-5506 to have the updated form faxed to your office OR you may complete the PA electronically (ePA). You may sign up online via CoverMyMeds at: www.covermymeds.com/epa/caremark/ or call 1-866-452-5017.</i> Yes - generic deferasirox  Yes - deferoxamine  Yes - deferiprone, <i>skip to section B</i> No - Continue request for Ferriprox	
6.	Has the patient experienced a documented intoleral REQUIRED: If Yes, attach supporting chart note	ble adverse event to the preferred product deferiprone? <i>ACTION</i> (s). $\square$ Yes $\square$ No
7.		expected adverse event attributed to the active ingredient as <i>N REQUIRED: If No, attach supporting chart note(s)</i> .

8. Does the patient have a documented inadequate response or intolerable adverse event with deferasirox?

Send completed form to: Case Review Unit, CVS Caremark Prior Authorization Fax: 1-866-249-6155

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	ACTION REQUIRED: If Yes, attach supporting chart note(s) and skip to #10. □ Yes □ No
9.	Does the patient have any of the following documented clinical reasons to avoid deferasirox products?  **ACTION REQUIRED: If Yes, attach supporting chart note(s).  **Estimated glomerular filtration rate (GFR) less than 40 mL/min/1.73 m²  **Poor performance status**  **High-risk myelodysplastic syndrome**  **Advanced malignancy**  **Platelet count less than 50 x 10 <sup>9</sup> /L  **Known hypersensitivity to deferasirox or any components of drug formulations**  **Severe (Child-Pugh C) hepatic impairment**  **None of the above**
10.	Does the patient have a documented inadequate response or intolerable adverse event with deferoxamine? <i>ACTION REQUIRED: If Yes, attach supporting chart note(s) and skip to section B.</i> $\square$ Yes $\square$ No
11.	Does the patient have any of the following documented clinical reasons to avoid deferoxamine products? <i>ACTION REQUIRED: If Yes, attach supporting chart note(s)</i> .  ☐ Severe renal disease ☐ Anuria ☐ Known hypersensitivity to deferoxamine ☐ None of the above
	tion B: All Requests  Is this a request for continuation of therapy with the requested drug?   Yes   No If No, skip to #16
13.	If patient has diagnosis of Hereditary hemochromatosis, is the patient experiencing benefit from therapy as evidenced by decreased serum ferritin levels as compared to pretreatment baseline?  Yes No No further questions
14.	Is the patient experiencing benefit from therapy as evidenced by a decrease in serum ferritin levels as compared to pretreatment baseline? <i>ACTION REQUIRED: If Yes, attach supporting laboratory report or chart notes with current serum ferritin level.</i> $\square$ Yes $\square$ No
15.	Is the patient's serum ferritin level consistently below 500 mcg/L? ☐ Yes ☐ No No further questions.
16.	Does the member have transfusional iron overload due to myelodysplastic syndrome or Diamond Blackfan anemia $^{\circ}$ Yes $^{\square}$ No
17.	Is the patient's pretreatment serum ferritin level consistently greater than 1000 mcg/L? <i>Note: If the patient is currently on therapy for iron overload, provide the patient's serum ferritin level before patient initiated therapy.</i> ACTION REQUIRED: If Yes, attach supporting laboratory report or chart notes with pretreatment serum ferritin level.   Yes No
18.	Will the dose of the requested drug exceed 99 mg/kg per day? ☐ Yes ☐ No
19.	Has the patient had an unsatisfactory response to phlebotomy? If Yes, no further questions ☐ Yes ☐ No
20.	Is phlebotomy not an option for the patient (e.g., poor candidate due to underlying medical conditions)? ☐ Yes ☐ No
	ttest that this information is accurate and true, and that documentation supporting this formation is available for review if requested by CVS Caremark or the benefit plan sponsor.
Х	
_	escriber or Authorized Signature Date (mm/dd/yy)

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