US Family Health Plan Prior Authorization Request Form for ferric maltol (Accrufer)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

Initial and renewal prior authorization expires after 6 months. For renewal of therapy, an initial USFHP prior authorization approval is required.

author	ızatıon	approval is required.						
Step	Please complete patient and physician information (please print):							
.1	Patient Name: Phy		rsician Name:					
	Address:		Address:					
	Sponsor ID#		Phone #:					
	Date of Birth:		Secure Fax #:					
Step 2	Please complete the clinical assessment:							
	1.	Has the patient received this medication under the USFHP benefit in the last 6 months? Please choose "No" if the patient did not previously have a USFHP approved PA for Accrufer.	☐ Yes	□No				
			(subject to verification)	Proceed to question 4				
			Proceed to question 2	4.00000				
	2.	Is the patient still iron deficient?	□ Yes	□No				
			Proceed to question 3	STOP				
				Coverage not approved				
	3.	Has there been clinically significant improvement in the patient's iron deficiency? Please Note: Medical documentation specific to your response to this question must be attached to this case or your request could be denied.	☐ Yes	□ No				
			Sign and date below	STOP				
				Coverage not approved				
	4.	Does the patient have a documented diagnosis of iron deficiency?	□ Yes	□ No				
			Proceed to question 5	STOP				
				Coverage not approved				
	5.	Is the patient 18 years of age or older?	☐ Yes	□No				
			Proceed to question 6	STOP				
				Coverage not approved				
	6.	Has the patient tried and failed two oral iron products (must be different salts) of at least six weeks in duration for each?	☐ Yes	□ No				
			Proceed to question 8	Proceed to question 7				

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	7.	Has the patient had a contraindication to or experienced clinically significant adverse effects		☐ Yes Proceed to question 8	□ No STOP			
		to two oral iron products (n salts)?	must be different	Proceed to question a	Coverage not approved			
	8.	Please provide the date of or clinically significant adventors. Note: The dates for each me each medication must be p	n, or the contraindication n medication.					
	Oral iro	on product:	Date of trial and failure:					
	Contraindication to medication or clinically significant adverse effect:							
	Oral iron product:		Date of trial and failure:					
	Contraindication to medication or clinically significant adverse effect:							
	Sign and date below							
Step 3	I certify the above is true to the best of my knowledge. Please sign and date:							
		Prescriber Signa	ature	Date				
		_			.[30 July 2021]			