

## **MEDICARE FORM**

## Feraheme® (ferumoxytol) and Injectafer® (ferric carboxymaltose) Medication Precertification Request

Page 1 of 2

(All fields must be completed and legible for precertification review.)

For Ohio MMP:

**FAX:** 1-855-734-9389 **PHONE:** 1-855-364-0974

For other lines of business:

Please use other form.

Note: Feraheme, Injectafer, and Monoferric are non-preferred. The preferred products are Ferrlecit (sodium ferric gluconate), Infed,

and Venofer.

Please indicate:	☐ Start of treatment: Start d	ate / /	<del>-</del>					
	☐ Continuation of therapy, □	ate of last treatment	1 1					
Precertification R	Requested By:		Phone:		Fax:			
A. PATIENT INFO	ORMATION							
First Name:		Last Name:			DOB:			
Address:		l	City:		State:	ZIP:		
Home Phone:	Work Ph	one:	Cell Phone:		Email:			
Patient Current We	eight: lbs or	kas Patient Height:	inches or	ms Allergi	es:			
B. INSURANCE II		<u> </u>	<u> </u>	3				
	#:	Does nationt have	e other coverage?	Yes 🗌 No				
	π		_					
Insured:		Insured:	r Od	Carrier Name:				
	S ☐ No If yes, provide ID #:		Medicaid: ☐ Yes ☐ N	lo If yes prov	vide ID #:			
C. PRESCRIBER			Wieulcalu.   Tes   N	io ii yes, pro	vide ID #.			
First Name:	INFORMATION	Last Name:		(Check Or	A). 🗆 M D	☐ D.O. ☐ N.P. ☐ P.A		
		Last Name.	0.1	(Check On	_	1		
Address:			City:	1	State:	ZIP:		
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	1	UPIN:		
Provider Email:		Office Contact Na	ime:		Phone:			
Specialty (Check of	one): 🗌 Hematologist 🔃 Int	ernal Medicine 🔲 Ot	ther:					
D. DISPENSING	PROVIDER/ADMINISTRATION	INFORMATION						
Place of Administ	tration:		Dispensing Prov	rider/Pharmac	y: Patient S	Selected choice		
☐ Self-administere	ed Physician's Office		☐ Physician's O	ffice	☐ Retail Ph	armacv		
☐ Outpatient Infus					_ ☐ Other	,		
	me:			-				
☐ Home Infusion (	Center Phone:		Name:					
Agency Na			Address:					
☐ Administration c	code(s) (CPT):					ZIP:		
Address:					Fax:			
	State:				PIN:			
	Fax:							
NPI:	PIN:							
E. PRODUCT INF	ORMATION							
	Feraheme  Injectafer Do	)SO.	Frequency: _					
-	FORMATION - Please indicate							
Primary ICD Code	_	Secondary ICD			ICD Code:			
•		<del></del>	·					
	ORMATION - Required clinical i		neted in its <u>entirety</u> for all p	precertification	requests.			
•	(clinical documentation require		d de . de Fermile - 14 (-			ford and Manager		
	njectafer, and Monoferric are not s the patient had prior therapy with				liuconate), in	tea, and venoter.		
	s the patient had prior therapy with	` -	• •	•				
		, ,	• '	•				
Yes No Has the patient had prior therapy with Monoferric (ferric derisomaltose injection) within the last 365 days?  Yes No Has the patient had a trial and failure, intolerance, or contraindication to any of the following? (select all that apply)								
	Ferrlecit (sodium ferric gluconate)		-	g. (coloct all til	at apply)			
	ere are any other medical reason(			erred products	when indicate	d for the patient's		
diagnosis (select al	,	-, pason oannot u	protecting protecting					
│ <sup>¯</sup> ` □	Ferrlecit (sodium ferric gluconate)	) ☐ Infed (iron dextran)	☐ Venofer (iron sucrose)					
	<u> </u>	· , ,	·			<u>_</u>		
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Patient First Name	Patient Last Name	Patient Phone	Patient DOB					
G. CLINICAL INFORMATION (conti	<b>nued)</b> – Required clinical information	on must be completed in its <u>entire</u>	$\underline{\mathbf{v}}$ for all precertification requests.					
Please indicate the patient's serum f Please indicate the patient's transfer Yes No Was the serum ferri Yes No Is this a request for	rrin saturation (TSAT) level: tin and/or transferrin saturation le continuation of therapy?	vel drawn within the last 30 day						
Yes No Does the patient have a contraindication, intolerance or ineffective response to Ferrlecit, Infed, or Venofer?  For chronic kidney disease indications only:								
Yes No Does the patient have iron deficiency anemia associated with chronic kidney disease?  Yes No Is the patient non-dialysis dependent (NDD) or undergoing peritoneal dialysis?  Please explain: The patient is non-dialysis dependent (NDD) The patient is undergoing peritoneal dialysis								
For all other non- chronic kidney disease indications:								
☐ The patient is unable to tolerate oral iron compounds								
☐ The patient is losing iron (blood) at a rate that is too rapid for oral intake to compensate for the loss ☐ The patient has a gastrointestinal tract disorder, such as inflammatory bowel disease (ulcerative colitis, and Crohn's disease) that may be aggravated by oral iron therapy								
☐ The patient is unable to maintain iron balance on treatment with hemodialysis								
The patient is donating large amounts of blood for autologous programs								
The patient has failed to heed instructions for oral iron supplementation or are incapable of accepting or following them								
The patient has heart failure and iron deficiency with or without anemia								
☐ The patient has iron deficiency and chemotherapy-induced anemia ☐ The patient has iron deficiency anemia due to heavy uterine bleeding								
☐ The patient has iron deficiency anemia due to heavy diefine bleeding ☐ The patient has iron deficiency following gastric bypass surgery and/or subtotal gastric resection and who exhibited decreased absorption of oral iron								
H. ACKNOWLEDGEMENT								
Request Completed By (Signature	Required):		Date:/ /					
Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.								

The plan may request additional information or clarification, if needed, to evaluate requests.