

## **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 New Jersey HMO D-SNP: FAX: 1-833-322-0034

**PHONE**: 1-844-362-0934

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 treatmen							
	Continuation of t		of last treatment			_		
	lequested By:			Phone:		Fax:		
A. PATIENT INFO	RMATION							
First Name:			Last Name:	<del></del>		DOB:		
Address:				City:		State:	ZIP:	
Home Phone:		Work Phone:		Cell Phone:	Т	Email:		
Patient Current We	ight: lbs or	kgs	Patient Height:	_inches orcm	ns Allergie	es:		
B. INSURANCE IN	NFORMATION							
Aetna Member ID	#:		Does patient have other coverage? ☐ Yes ☐ No					
	Group #:		If yes, provide ID#: Carrier Name:					
Insured:			Insured:					
	□ No If yes, provid	de ID #:	M	<b>ledicaid:</b> ☐ Yes ☐ No	If yes, prov	ide ID #:		
C. PRESCRIBER	INFORMATION							
First Name:			Last Name:		(Check On	1	D.O.   N.P.   P.A.	
Address:				City:		State:	ZIP:	
Phone:	Fax:		St Lic #:	NPI #:	DEA #:		UPIN:	
Provider Email:			Office Contact Name	э:		Phone:		
Specialty (Check of	one):	st 🗌 Internal	I Medicine ☐ Othe	er:				
D. DISPENSING P	PROVIDER/ADMINIS	TRATION INFO						
Place of Administ	ration:			Dispensing Provid	der/Pharmac	y: Patient Sele	ected choice	
☐ Self-administere	ed 🔲 Physicia	an's Office		☐ Physician's Offi		- □ Retail Pharn		
☐ Outpatient Infusi		one:		_ ☐ Specialty Pharr		Other	-	
	me:			Name:	-			
Home Infusion C		one:		Address:				
Agency Na				_   -			ZIP:	
Address:							Σπ	
		State:	ZIP:					
TIN:	i	PIN:		— NPI:				
NPI:				_				
E. PRODUCT INFO	ORMATION							
-	Feraheme 🗌 Injec			Frequency:				
F. DIAGNOSIS IN	FORMATION - Please	e indicate prima	ry ICD code and speci	fy any other where applic	cable.			
Primary ICD Code	»: 🔲		Secondary ICD Co	ode :	Other	ICD Code:		
G. CLINICAL INFO	ORMATION - Require	ed clinical inform	nation must be complet	ed in its <u>entirety</u> for all pr	ecertification	requests.		
	(clinical documentat	•			_	_	_	
Note: Feraheme, Injectafer, and Monoferric are non-preferred. The preferred products are Ferrlecit (sodium ferric gluconate), Infed, and Venofer.								
Yes No. Has the patient had prior therapy with Feraheme (ferumoxytol injection) within the last 365 days?								
Yes No. Has the patient had prior therapy with Injectafer (ferric carboxymaltose injection) within the last 365 days?								
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)								
Please explain if there are any other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's								
diagnosis (select all that apply).								
	☐ Ferrlecit (sodium ferric gluconate) ☐ Infed (iron dextran) ☐ Venofer (iron sucrose)							



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB						
G CLINICAL INFORMATION (confin	wood) Required clinical information mus	et he completed in its entire	ty for all procertification requests						
G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.									
Please indicate the patient's serum ferritin level:									
Please indicate the patient's transferrin saturation (TSAT) level:									
Yes No Was the serum ferritin and/or transferrin saturation level drawn within the last 30 days?									
Yes No Is this a request for continuation of therapy?									
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 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.