



# MEDICARE FORM

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

Page 1 of 2

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

For Medicare Advantage Part B:  
FAX: 1-844-268-7263  
PHONE: 1-866-503-0857 (TTY: 711)

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 date \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
 Continuation of therapy, Date of last treatment \_\_\_\_ / \_\_\_\_ / \_\_\_\_

Precertification Requested By: \_\_\_\_\_ Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

### A. PATIENT INFORMATION

First Name:		Last Name:		DOB:	
Address:			City:	State:	ZIP:
Home Phone:		Work Phone:		Cell Phone:	
Patient Current Weight: _____ lbs or _____ kgs		Patient Height: _____ inches or _____ cms		Allergies:	

### B. INSURANCE INFORMATION

Aetna Member ID #: _____		Does patient have other coverage? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Group #: _____		If yes, provide ID#: _____ Carrier Name: _____	
Insured: _____		Insured: _____	
Medicare: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____		Medicaid: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____	

### C. PRESCRIBER INFORMATION

First Name:		Last Name:		(Check One): <input type="checkbox"/> M.D. <input type="checkbox"/> D.O. <input type="checkbox"/> N.P. <input type="checkbox"/> P.A.	
Address:			City:	State:	ZIP:
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	UPIN:
Provider Email:		Office Contact Name:		Phone:	
Specialty (Check one): <input type="checkbox"/> Hematologist <input type="checkbox"/> Internal Medicine <input type="checkbox"/> Other: _____					

### D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

<b>Place of Administration:</b> <input type="checkbox"/> Self-administered <input type="checkbox"/> Physician's Office <input type="checkbox"/> Outpatient Infusion Center Phone: _____ Center Name: _____ <input type="checkbox"/> Home Infusion Center Phone: _____ Agency Name: _____ <input type="checkbox"/> Administration code(s) (CPT): _____ Address: _____ City: _____ State: _____ ZIP: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____ NPI: _____		<b>Dispensing Provider/Pharmacy: Patient Selected choice</b> <input type="checkbox"/> Physician's Office <input type="checkbox"/> Retail Pharmacy <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Other Name: _____ Address: _____ City: _____ State: _____ ZIP: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____ NPI: _____	
--	--	--	--

### E. PRODUCT INFORMATION

Request is for:  Feraheme  Injectafer  Monoferric Dose: \_\_\_\_\_ Frequency: \_\_\_\_\_  
HCPCS Code: \_\_\_\_\_

### F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other where applicable.

Primary ICD Code:  \_\_\_\_\_ Secondary ICD Code : \_\_\_\_\_ Other ICD Code: \_\_\_\_\_

### G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests.

**For All Requests (clinical documentation required for all requests):**  
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)  Infed (iron dextran)  Venofer (iron sucrose)  
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)  
\_\_\_\_\_  
\_\_\_\_\_

Continued on next page



# MEDICARE FORM

## Feraheme® (ferumoxyl) and Injectafer® (ferric carboxymaltose) Monoferric® (ferric derisomaltose) Medication Precertification Request

Page 2 of 2

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

For Medicare Advantage Part B:  
FAX: 1-844-268-7263  
PHONE: 1-866-503-0857 (TTY: 711)

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.

Patient First Name	Patient Last Name	Patient Phone	Patient DOB
--------------------	-------------------	---------------	-------------

### 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.