



MEDICARE FORM

Feraheme® (ferumoxylol) and
Injectafer® (ferric carboxymaltose)
Medication Precertification Request

Page 1 of 2

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

Virginia (HMO D-SNP)
FAX: 1-833-280-5224
PHONE: 1-855-463-0933

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: Email:
Patient Current Weight: lbs or kgs Patient Height: inches or cms Allergies:

B. INSURANCE INFORMATION
Aetna Member ID #: Does patient have other coverage?
Group #: If yes, provide ID#: Carrier Name:
Insured: Insured:
Medicare: Yes No If yes, provide ID #: Medicaid: Yes No If yes, provide ID #:

C. PRESCRIBER INFORMATION
First Name: Last Name: (Check One): M.D. D.O. N.P. P.A.
Address: City: State: ZIP:
Phone: Fax: St Lic #: NPI #: DEA #: UPIN:
Provider Email: Office Contact Name: Phone:
Specialty (Check one): Hematologist Internal Medicine Other:

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION
Place of Administration: Self-administered Physician's Office
Outpatient Infusion Center Phone:
Center Name:
Home Infusion Center Phone:
Agency Name:
Administration code(s) (CPT):
Address: City: State: ZIP:
Phone: Fax: TIN: PIN: NPI:
Dispensing Provider/Pharmacy: Patient Selected choice
Physician's Office Retail Pharmacy
Specialty Pharmacy Other
Name:
Address:
City: State: ZIP:
Phone: Fax: TIN: PIN: NPI:

E. PRODUCT INFORMATION
Request is for: Feraheme Injectafer Dose: Frequency:

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.
Has the patient had prior therapy with Feraheme (ferumoxylol injection) within the last 365 days?
Has the patient had prior therapy with Injectafer (ferric carboxymaltose injection) within the last 365 days?
Has the patient had prior therapy with Monoferric (ferric derisomaltose injection) within the last 365 days?
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)

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
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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.