

Medical Drug Clinical Criteria

Subject: Iron Agents
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Overview

This document addresses the use of injectable agents for the treatment of iron deficiency anemia (IDA). Agents addressed in this document include:

- Feraheme (ferumoxytol)
- Ferrlecit (sodium ferric gluconate/sucrose complex)
- Infed (iron dextran)
- Injectafer (ferric carboxymaltose)
- Monoferric (ferric derisomaltose)
- Venofer (iron sucrose)

Iron is a mineral in the body that is an essential component for blood production, enabling them to carry oxygen throughout the body. The majority of body iron are found in circulating red blood cells called hemoglobin, while the remaining is stored as ferritin or bound to myoglobin in muscle cells. Individuals with iron deficiency anemia may have mild to severe symptoms, ranging from fatigue, shortness of breath, and chest pain to heart failure and developmental delays in children (NHLBI 2019).

The causes of iron deficiency anemia are numerous, including gastrointestinal bleeding or other blood loss, chronic kidney disease, celiac disease, multiple blood donations, and cancer or chemotherapy-related etiologies. Diagnosis of IDA is typically confirmed by evaluating levels of serum ferritin, transferrin saturation (TSAT), absence of stainable iron in the bone marrow, or resolution of anemia upon iron administration (Auerbach 2020).

While the 2012 Kidney Disease Improving Global Outcomes (KDIGO) guidelines for anemia in chronic kidney disease do not provide any guidance on preference of available IV iron agents over another, they do suggest that a trial oral iron for 1 to 3 months can be appropriate for individuals with IDA prior to initiating IV iron. The National Comprehensive Cancer Network (NCCN) guidelines for Hematopoietic Growth Factors provides a category 2A recommendation for use of Feraheme, Ferrlecit, Infed (IV only; IM not recommended), Injectafer, Venofer, and Monoferric for the management of cancer- and chemotherapy-induced anemia. NCCN also suggests that a trial of oral iron for at least 4 weeks can be appropriate prior to initiating IV iron.

Both Feraheme and Infed have black box warnings for fatal and serious hypersensitivity reactions including anaphylaxis, and as such, the administration of which should only occur when personnel and therapies are immediately available for the treatment of anaphylaxis and other hypersensitivity reactions.

Clinical Criteria

When a drug is being reviewed for coverage under a member's medical benefit plan or is otherwise subject to clinical review (including prior authorization), the following criteria will be used to determine whether the drug meets any applicable medical necessity requirements for the intended/prescribed purpose.

Feraheme (ferumoxytol), Ferrlecit (sodium ferric gluconate/sucrose complex), Venofer (iron sucrose)

Requests for Feraheme (ferumoxytol), Ferrlecit (sodium ferric gluconate/sucrose complex), Venofer (iron sucrose) may be approved if the following criteria are met:

- I. Individual has a diagnosis of chronic kidney disease (CKD); **AND**
 - A. Individual is dialysis dependent; **AND**
 - B. Individual has iron deficiency anemia (IDA);
- OR**
- II. Individual has a diagnosis of iron deficiency anemia (IDA) or iron deficiency; **AND**
- III. Individual is non-dialysis dependent; **AND**
- IV. Diagnosis is confirmed by one of the following:
 - A. For IDA or iron deficiency associated with CKD or inflammatory conditions (for example, inflammatory bowel disease [IBD], heart failure), individual meets *one* of the following within the last ~~four~~eight (48) weeks (De Franceschi 2017):
 - 1. Serum ferritin levels less than 100 ng/mL; **OR**
 - 2. TSAT levels less than 20%; **OR**
 - 3. Serum ferritin is less than or equal to 500 ng/mL **and** TSAT is less than or equal to 30% (KDIGO 2012); **OR**
 - 4. Bone marrow demonstrates inadequate iron stores; **OR**
 - 5. Hemoglobin (HGB) less than 13 g/dl (less than 130 g/l) in males or less than 12 g/dl (less than 120 g/l) in females and TSAT 30% or less and ferritin 500ng/ml or less (500 µg/l or less) (Ko 2020); **OR**
 - B. For IDA or iron deficiency associated with cancer/chemotherapy or non-inflammatory conditions (for example, blood loss, malabsorption, malnutrition), individual meets *one* of the following within the last ~~four~~eight (48) weeks (NCCN 2024, De Franceschi 2017):
 - 1. Serum ferritin levels less than 30 ng/mL; **OR**
 - 2. TSAT levels less than 20%; **OR**
 - 3. Bone marrow demonstrates inadequate iron stores; **AND**
- V. Individual has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (NCCN 2024, KDIGO 2012); **OR**
- VI. Individual is unable to use oral iron supplementation for one of the following reasons:
 - A. Malabsorption conditions; **OR**
 - B. Gastric Surgery (DeFlipp 2013);
- OR**
- VII. Individual has iron deficiency anemia or iron deficiency in pregnancy; **AND**
- VIII. Diagnosis is confirmed by one of the following:
 - A. Serum ferritin levels less than 30 ng/mL; **OR**
 - B. TSAT levels less than 20%; **OR**
 - C. Bone marrow demonstrates inadequate iron stores; **AND**
- IX. Individual is past 14 weeks of pregnancy and has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (Muñoz 2017); **OR**
- X. Individual is past 14 weeks of pregnancy and individual is unable to use oral iron supplementation due to malabsorptive conditions or gastric surgery (Muñoz 2017, DeFlipp 2013); **OR**
- XI. Individual is past 14 weeks of pregnancy and diagnosed with severe iron deficiency anemia, defined as Hemoglobin (HGB) less than 8 g/dL; **OR**
- XII. Individual is past 34 weeks of pregnancy.

Feraheme (ferumoxytol), Ferrlecit (sodium ferric gluconate/sucrose complex), or Venofer (iron sucrose) may not be approved when the above criteria are not met and for all other indications.

Approval Duration (dialysis-dependent use excluded)
6 months

Infed (iron dextran)

Requests for Infed (iron dextran) may be approved if the following criteria are met:

- I. Individual has a diagnosis of chronic kidney disease (CKD); **AND**
 - A. Individual is dialysis dependent; **AND**
 - B. Individual has iron deficiency anemia (IDA);

OR

- II. Individual has a diagnosis of iron deficiency anemia (IDA) or iron deficiency; **AND**
- III. Individual is non-dialysis dependent; **AND**
- IV. Diagnosis is confirmed by one of the following:
 - A. For IDA or iron deficiency associated with CKD or inflammatory conditions (for example, inflammatory bowel disease [IBD], heart failure), individual meets *one* of the following within the last ~~four~~**eight** (48) weeks (De Franceschi 2017):
 - 1. Serum ferritin levels less than 100 ng/mL; **OR**
 - 2. TSAT levels less than 20%; **OR**
 - 3. Serum ferritin is less than or equal to 500 ng/mL **and** TSAT is less than or equal to 30% (KDIGO 2012); **OR**
 - 4. Bone marrow demonstrates inadequate iron stores; **OR**
 - 5. Hemoglobin (HGB) less than 13 g/dl (less than 130 g/l) in males or less than 12 g/dl (less than 120 g/l) in females and TSAT 30% or less and ferritin 500ng/ml or less (500 µg/l or less) (Ko 2020); **OR**
 - B. For IDA or iron deficiency associated with cancer/chemotherapy or non-inflammatory conditions (for example, malabsorption, malnutrition), individual meets *one* of the following within the last ~~four~~**eight** (48) weeks (NCCN 2024, De Franceschi 2017):
 - 1. Serum ferritin levels less than 30 ng/mL; **OR**
 - 2. TSAT levels less than 20%; **OR**
 - 3. Bone marrow demonstrates inadequate iron stores; **AND**
- V. Individual has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (NCCN 2024, KDIGO 2012); **OR**
- VI. Individual is unable to use oral iron supplementation for one of the following reasons:
 - A. Malabsorption conditions; **OR**
 - B. Gastric Surgery (DeFlipp 2013);

OR

- VII. Individual has iron deficiency anemia or iron deficiency in pregnancy; **AND**
- VIII. Diagnosis is confirmed by one of the following:
 - A. Serum ferritin levels less than 30 ng/mL; **OR**
 - B. TSAT levels less than 20%; **OR**
 - C. Bone marrow demonstrates inadequate iron stores; **AND**
- IX. Individual is past 14 weeks of pregnancy and has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (Muñoz 2017); **OR**
- X. Individual is past 14 weeks of pregnancy and individual is unable to use oral iron supplementation due to malabsorptive conditions or gastric surgery (Muñoz 2017, DeFlipp 2013); **OR**
- XI. Individual is past 14 weeks of pregnancy and diagnosed with severe iron deficiency anemia, defined as Hemoglobin (HGB) less than 8 g/dL; **OR**
- XII. Individual is past 34 weeks of pregnancy;

OR

- XIII. Individual is diagnosed with iron deficiency due to blood loss; **AND**
- XIV. Diagnosis is confirmed by one of the following:
 - A. Serum ferritin levels less than 100 ng/mL; **OR**
 - B. TSAT levels less than 20%; **OR**
 - C. Serum ferritin is less than or equal to 500 ng/mL **and** TSAT is less than or equal to 30% (KDIGO 2012); **OR**
 - D. Bone marrow demonstrates inadequate iron stores.

Infed (iron dextran) may not be approved when the above criteria are not met and for all other indications.

Approval Duration (~~dialysis-dependent use excluded~~)
6 months

Injectafer (ferric carboxymaltose)

Requests for Injectafer (ferric carboxymaltose) may be approved if the following criteria are met:

- I. Individual has a diagnosis of chronic kidney disease (CKD); **AND**
 - A. Individual is dialysis dependent; **AND**
 - B. Individual has iron deficiency anemia (IDA);

OR

- II. Individual has a diagnosis of iron deficiency anemia (IDA) or iron deficiency; **AND**
- III. Individual is non-dialysis dependent; **AND**
- IV. Diagnosis is confirmed by one of the following:
 - A. For IDA or iron deficiency associated with CKD or inflammatory conditions (for example, inflammatory bowel disease [IBD], heart failure), individual meets *one* of the following within the last **four**eight (48) weeks (De Franceschi 2017):
 - 1. Serum ferritin levels less than 100 ng/mL; **OR**
 - 2. TSAT levels less than 20%; **OR**
 - 3. Serum ferritin is less than or equal to 500 ng/mL **and** TSAT is less than or equal to 30% (KDIGO 2012); **OR**
 - 4. Bone marrow demonstrates inadequate iron stores; **OR**
 - 5. Hemoglobin (HGB) less than 13 g/dl (less than 130 g/l) in males or less than 12 g/dl (less than 120 g/l) in females and TSAT 30% or less and ferritin 500ng/ml or less (500 µg/l or less) (Ko 2020); **OR**
 - B. For IDA or iron deficiency associated with cancer/chemotherapy or non-inflammatory conditions (for example, blood loss, malabsorption, malnutrition), individual meets *one* of the following within the last **four**eight (48) weeks (NCCN 2024, De Franceschi 2017):
 - 1. Serum ferritin levels less than 30 ng/mL; **OR**
 - 2. TSAT levels less than 20%; **OR**
 - 3. Bone marrow demonstrates inadequate iron stores; **AND**
- V. Individual has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (NCCN 2024, KDIGO 2012); **OR**
- VI. Individual is unable to use oral iron supplementation for one of the following reasons:
 - A. Malabsorption conditions; **OR**
 - B. Gastric Surgery (DeFlipp 2013);

OR

- VII. Individual has iron deficiency anemia or iron deficiency in pregnancy; **AND**
- VIII. Diagnosis is confirmed by one of the following:
 - A. Serum ferritin levels less than 30 ng/mL; **OR**
 - B. TSAT levels less than 20%; **OR**
 - C. Bone marrow demonstrates inadequate iron stores; **AND**
- IX. Individual is past 14 weeks of pregnancy and has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (Muñoz 2017); **OR**
- X. Individual is past 14 weeks of pregnancy and individual is unable to use oral iron supplementation due to malabsorptive conditions or gastric surgery (Muñoz 2017, DeFlipp 2013); **OR**
- XI. Individual is past 14 weeks of pregnancy and diagnosed with severe iron deficiency anemia, defined as Hemoglobin (HGB) less than 8 g/dL; **OR**
- XII. Individual is past 34 weeks of pregnancy;

OR

- XIII. Individual is diagnosed with iron deficiency in adult patients with heart failure with New York Heart Association class II/III; **AND**
- XIV. Individual is using to improve exercise capacity; **AND**
- XV. Diagnosis is confirmed by one of the following (Heidenreich 2022):
 - A. Serum ferritin levels less than 100 µg/L; **OR**
 - B. TSAT levels less than 20% and ferritin level 100 to 300 µg/L.

Injectafer (ferric carboxymaltose) may not be approved when the above criteria are not met and for all other indications.

Approval Duration (dialysis-dependent use excluded)
6 months

Monoferic (ferric derisomaltose)

Requests for Monoferic (ferric derisomaltose) may be approved if the following criteria are met:

- I. Individual has a diagnosis of iron deficiency anemia (IDA) or iron deficiency; **AND**
- II. Individual is non-dialysis dependent; **AND**
- III. Diagnosis is confirmed by one of the following:

- A. For IDA or iron deficiency associated with CKD or inflammatory conditions (for example, inflammatory bowel disease [IBD], heart failure), individual meets *one* of the following within the last ~~four~~**eight** (48) weeks (De Franceschi 2017):
 1. Serum ferritin levels less than 100 ng/mL; **OR**
 2. TSAT levels less than 20%; **OR**
 3. Serum ferritin is less than or equal to 500 ng/mL **and** TSAT is less than or equal to 30% (KDIGO 2012); **OR**
 4. Bone marrow demonstrates inadequate iron stores; **OR**
 5. Hemoglobin (HGB) less than 13 g/dl (less than 130 g/l) in males or less than 12 g/dl (less than 120 g/l) in females and TSAT 30% or less and ferritin 500ng/ml or less (500 µg/l or less) (Ko 2020); **OR**
- B. For IDA or iron deficiency associated with cancer/chemotherapy or non-inflammatory conditions (for example, blood loss, malabsorption, malnutrition), individual meets *one* of the following within the last ~~four~~**eight** (48) weeks (NCCN 2024, De Franceschi 2017):
 1. Serum ferritin levels less than 30 ng/mL; **OR**
 2. TSAT levels less than 20%; **OR**
 3. Bone marrow demonstrates inadequate iron stores; **AND**
- IV. Individual has had a four (4) week trial of and inadequate response, or intolerance to oral iron supplementation (NCCN 2024, KDIGO 2012); **OR**
- V. Individual is unable to use oral iron supplementation for one of the following reasons:
 - A. Malabsorption conditions; **OR**
 - B. Gastric Surgery (DeFlipp 2013).

Monoferic (ferric derisomaltose) may not be approved for the following:

- I. Individual has hemodialysis dependent chronic kidney disease (CKD); **OR**
- II. When the above criteria are not met and for all other indications.

Approval Duration 6 months

Triferic/Triferic AVNU (ferric pyrophosphate citrate)

Requests for Triferic/Triferic AVNU (ferric pyrophosphate citrate) may be approved if the following criteria are met:

- I. ~~Individual has a diagnosis of chronic kidney disease (CKD);~~ **AND**
 - A. ~~Individual is hemodialysis dependent;~~ **AND**
 - B. ~~Individual has iron deficiency anemia (IDA).~~

Triferic/Triferic AVNU (ferric pyrophosphate citrate) may not be approved for the following:

- I. ~~Peritoneal dialysis;~~ **OR**
- II. ~~When the above criteria are not met and for all other indications.~~

Step Therapy

Summary of FDA-approved and NCCN 2A recommended indications for agents for Iron Deficiency Anemia (IDA):

Agent	Route	Oral iron intolerant or unresponsive IDA	CKD	Dialysis-dependent CKD only	Iron Replacement for Blood Loss	NCCN
Feraheme (ferumoxytol)	IV	x	x			x
Ferlecit (sodium ferric gluconate/sucrose complex)	IV			x*		x
Infed (iron dextran)	IV, IM	x*				x (IV only)
Injectafer (ferric carboxymaltose)	IV	x	x			x
Monoferic (ferric derisomaltose)	IV	x	x			x

Triferic, Triferic AVNU (ferric pyrophosphate citrate)	IV			x		
Venofer (iron sucrose)	IV		x*			x

*Includes FDA-approved pediatric indication

Note: When an IDA agent is deemed approvable based on the clinical criteria above, the benefit plan may have additional criteria requiring the use of a preferred¹ agent or agents.

Non-Preferred Iron Deficiency Anemia (IDA) Step Therapy

A list of the preferred iron deficiency anemia agents is available [here](#).

Requests for a non-preferred agent for IDA may be approved when the following criteria are met:

- I. Individual has had a trial and inadequate response or intolerance to two (2) preferred agents;

OR

- II. If Injectafer (ferric carboxymaltose) is designated as non-preferred, then it may be approved for the following:
 - A. Individual is being treated for iron deficiency in heart failure;

OR

- III. The preferred agent(s) are not acceptable due to concomitant clinical conditions, including but not limited to known hypersensitivity to any active or inactive component which is not also associated with the requested non-preferred agent;

OR

- IV. Individual is dialysis-dependent and using iron in conjunction with dialysis.

¹Preferred, as used herein, refers to agents that were deemed to be clinically comparable to other agents in the same class or disease category but are preferred based upon clinical evidence and cost effectiveness.

Quantity Limits

Iron Deficiency Anemia Agents Quantity Limits

Drug	Limit
Feraheme (ferumoxytol) 510 mg/17 mL vial*	1020 mg per 6 days [‡]
Ferlecit (sodium ferric gluconate/sucrose complex) 62.5 mg/5 mL vial*	1000mg per 8 weeks ^Δ
Injectafer (ferric carboxymaltose) 100mg/2ml vial*, 750 mg/15 mL vial*, 1000 mg/20 mL vial*	1500 mg per 7 days
Monoferic (ferric derisomaltose) 100 mg/mL vial, 500 mg/5 mL vial, 1000 mg/10 mL vial	1000 mg per day [‡]
Venofer (iron sucrose) 50 mg/2.5 mL vial*, 100 mg/5 mL vial*, 200 mg/10 mL vial*	1000 mg per 14 days [‡]
Override Criteria	
*Use in dialysis-dependent individuals excluded from quantity limits.	

[‡]Limit represents FDA-approved maximum dose recommendations per course of therapy (excluding dialysis-dependent diagnosis).

^ΔLimit according to NCCN guidelines for hematopoietic growth factors (v2.2023).

Coding

The following codes for treatments and procedures applicable to this document are included below for informational purposes. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member coverage or provider reimbursement policy. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

Feraheme (ferumoxytol), Ferrlecit (sodium ferric gluconate), Infed (iron dextran), Injectafer (ferric carboxymaltose), Venofer (iron sucrose)
HCPCS

<u>J1439</u>	<u>Injection, ferric carboxymaltose, 1 mg [Injectafer]</u>
<u>J1750</u>	<u>Injection, iron dextran, 50 mg [Infed]</u>
<u>J1756</u>	<u>Injection, iron sucrose, 1 mg [Venofer]</u>
<u>J2916</u>	<u>Injection, sodium ferric gluconate complex in sucrose injection, 12.5 mg [Ferrlecit]</u>
<u>Q0138</u>	<u>Injection, ferumoxytol, for treatment of iron deficiency anemia, 1 mg (non-ESRD use) [Feraheme]</u>
<u>Q0139</u>	<u>Injection, ferumoxytol, for treatment of iron deficiency anemia, 1 mg (for ESRD on dialysis) [Feraheme]</u>

ICD-10 Diagnosis

<u>D50.0-D50.9</u>	<u>Iron deficiency anemia</u>
<u>D63.0-C63.8</u>	<u>Anemia in chronic diseases classified elsewhere</u>
<u>D64.81</u>	<u>Anemia due to antineoplastic chemotherapy</u>
<u>E61.1</u>	<u>Iron deficiency</u>
<u>E83.10</u>	<u>Disorder of iron metabolism, unspecified</u>
<u>I50.1-I50.9</u>	<u>Heart failure</u>
<u>K50.00-K50.919</u>	<u>Crohn's disease [regional enteritis]</u>
<u>K51.00-K51.919</u>	<u>Ulcerative colitis</u>
<u>K90.0-K90.9</u>	<u>Intestinal malabsorption</u>
<u>K91.1</u>	<u>Postgastric surgery syndromes</u>
<u>K91.2</u>	<u>Postsurgical malabsorption, not elsewhere classified</u>
<u>N18.1-N18.9</u>	<u>Chronic kidney disease (CKD)</u>
<u>O99.011-O99.019</u>	<u>Anemia complicating pregnancy</u>
<u>T45.X5A-T45.X5D</u>	<u>Adverse effect of iron and its compounds [inadequate response or intolerance to oral iron supplementation]</u>
<u>Z98.84</u>	<u>Bariatric surgery status</u>

Monoferric (ferric derisomaltose)

HCPCS

<u>J1443</u>	<u>Injection, ferric pyrophosphate citrate solution, 0.1 mg of iron [Triferic]</u>
<u>J1445</u>	<u>Injection, ferric pyrophosphate citrate solution (Triferic AVNU), 0.1 mg of iron</u>
<u>J1437</u>	<u>Injection, ferric derisomaltose, 10 mg [Monoferric]</u>
<u>Q0138</u>	<u>Injection, ferumoxytol, for treatment of iron deficiency anemia, 1 mg (for non-ESRD on dialysis) [Feraheme]</u>
<u>J2916</u>	<u>Injection, sodium ferric gluconate complex in sucrose injection, 12.5 mg [Ferrlecit]</u>
<u>J1750</u>	<u>Injection, iron dextran, 50 mg [Infed]</u>
<u>J1756</u>	<u>Injection, iron sucrose, 1 mg [Venofer]</u>
<u>J1439</u>	<u>Injection, ferric carboxymaltose, 1 mg [Injectafer]</u>

ICD-10 Diagnosis

<u>D50.0-D50.9</u>	<u>Iron deficiency anemia</u>
<u>D63.0-D63.8</u>	<u>Anemia in chronic diseases classified elsewhere</u>
<u>D64.81</u>	<u>Anemia due to antineoplastic chemotherapy</u>

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E61.1	Iron deficiency
E83.10	Disorder of iron metabolism, unspecified
I50.1-I50.9	Heart failure
K50.00-K50.919	Crohn's disease [regional enteritis]
K51.00-K51.919	Ulcerative colitis
K90.0-K90.9	Intestinal malabsorption Celiac disease
K91.1	Postgastric surgery syndromes
K91.2	Postsurgical malabsorption, not elsewhere classified
N18.1-N18.59	Chronic kidney disease, stages I-V (CKD)
T45.X5A-T45.X5D	Adverse effect of iron and its compounds [inadequate response or intolerance to oral iron supplementation]
Z98.84	Bariatric surgery status
O99.011	Anemia complicating pregnancy, first trimester
O99.012	Anemia complicating pregnancy, second trimester
O99.013	Anemia complicating pregnancy, third trimester
O99.019	Anemia complicating pregnancy, unspecified trimester

Document History

Revised: 08/15/2025

Document History:

- 08/15/2025 – Annual Review: Remove approval duration time frames. Remove/Archive Triferic/Triferic AVNU PA, since it is no longer available. Update time frame to obtain lab values from 4 weeks to 8 weeks. Coding Reviewed: Removed HCPCS J1443 and J1445. Added HCPCS Q0139. Updated description for HCPCS Q0138. Consolidated codes O99.011-O99.019 into one range. Separated Monoferic from other iron agents and removed O99.011-O99.019 from Monoferic. Added ICD-10-CM E83.10, I50.1-I50.9, K51.00-K51.919, K91.1, K91.2, N18.6, N18.9, T45.X5A-T45.X5D, Z98.84 to all iron agents. Updated description for ICD-10-CM K90.0-K90.8.
- 08/16/2024 – Annual Review: Add iron deficiency clarification, add oral iron exception requirements to anemia in pregnancy, clarify blood loss iron deficiency for Infed. Coding Reviewed: Add ICD-10-CM E61.1.
- 12/01/2023 – Step therapy table updates.
- 09/11/2023 – Select Review: Add override criteria for Infed in step for iron deficiency due to blood loss, wording and formatting. Coding Reviewed: No changes.
- 08/18/2023 – Annual Review: Add hemoglobin in diagnosis, edit oral iron requirement, update Infed criteria to include iron deficiency from blood loss, Add Injectafer criteria related to heart failure, add override criteria for Injectafer in step edit, quantity limits, increase approval length. Step therapy table updates. Coding Reviewed: No changes.
- 05/15/2023 – Step therapy table updates.
- 05/01/2023 – Step therapy table updates.
- 03/27/2023 – Step therapy table updates.
- 01/25/2023 – Step therapy table updates.
- 01/11/2023 – Step therapy and step therapy table updates.
- 12/01/2022 – Step therapy table updates.
- 09/15/2022 – Select Review: add quantity limit for Injectafer 100mg/2ml vial. Coding Reviewed: No changes.
- 08/19/2022 – Annual Review: Add criteria for iron deficiency anemia in pregnancy, wording and formatting changes. Coding reviewed: Added ICD-10-CM O99.011, O99.012, O99.013, O99.019.
- 04/25/2022 – Step therapy table updates.
- 03/28/2022 – Step therapy table updates.
- 08/20/2021 – Annual Review: Update criteria to update approval durations to three months for all requests, and remove continuation approval duration. Add timeframe parameters for lab values for Feraheme, Infed, Venofer, Ferlecit, Injectafer, and Monoferic. Update Monoferic non-approvable criteria to restrict use in hemodialysis patients per label. Remove QL override criteria for Monoferic. Add generically available version of Feraheme to step therapy criteria. Update Injectafer QL to include new strength. Clarify use in

hemodialysis dependent CKD patients. Update references, and wording and formatting changes. Coding reviewed: Added HCPCS J1445.

- 08/23/2021 – Step therapy table updates.
- 07/26/2021 – Step therapy table updates.
- 04/26/2021 - Step therapy table updates.
- 08/21/2020 – Annual Review: Add new clinical criteria document, including PA, step therapy, and quantity limits, for Injectafer, Infed, Venofer, Triferic/Triferic AVNU, Feraheme, Monoferic, and Ferlecit. Coding Reviewed: Added HCPCS codes- J1443, J1437, Q0138, J2916, J1750, J1439, J1756. Added ICD-10-CM codes-D50.0-D50.9, D63.0-D63.8, D64.81, K50.00-K50.919, K90.0, K90.4, K90.9, N18.1-N18.5. Effective 2/1/2021 extended ICD-10-CM K50.00-K50.919, Extended K90.0-K90.9, Removed K90.4, and K90.9.

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CC-0182 Agents for Iron Deficiency Anemia

Commercial Medical Benefit

Effective Date	Preferred Agents	Non-Preferred Agents
03/01/2023	Ferrlecit Feraheme Venofer	Infed Injectafer Monoferric
10/01/2023	Ferrlecit Feraheme Infed Venofer	Injectafer Monoferric

Medicaid Medical Benefit

Effective Date	Preferred Agents	Non-Preferred Agents
09/18/2023: AR, CA, DC, FL Healthy Kids, GA, IA, KY, LA, MD, NJ, NV, NY, OH, SC, TN, VA, WI, WNY	Ferrlecit Feraheme Infed Venofer	Injectafer Monoferric

Medicare Medical Benefit

Effective Date	Preferred Agents	Non-Preferred Agents
03/01/2023	Ferrlecit Feraheme Venofer	Infed Injectafer Monoferric
12/01/2023	Ferrlecit Feraheme Infed Venofer	Injectafer Monoferric