



MonoFerric
Patient Solutions®

MONOFERRIC PATIENT SOLUTIONS® (MPS) PROGRAM

Rapid Response, Customized Solutions

Services offered through the MPS Program include insurance coverage and reimbursement information, patient assistance program, copay assistance, and more

Sample MonoFerric® (ferric derisomaltose) Patient That May Qualify for Patient Services Diagnosed with Iron Deficiency Anemia (IDA):

Patient who has intolerance to oral iron or have had unsatisfactory response to oral iron

OR

Patient who has non-hemodialysis chronic kidney disease (NDD-CKD)

To learn more, call **1-800-992-9022** or visit [monoferric-patient-solutions.com](https://www.monoferric-patient-solutions.com)



INDICATION

MonoFerric is indicated for the treatment of iron deficiency anemia (IDA) in adult patients:

- who have intolerance to oral iron or have had unsatisfactory response to oral iron
- who have non-hemodialysis dependent chronic kidney disease (NDD-CKD)

IMPORTANT SAFETY INFORMATION

CONTRAINDICATIONS

MonoFerric is contraindicated in patients with a history of serious hypersensitivity to MonoFerric or any of its components. Reactions have included shock, clinically significant hypotension, loss of consciousness, and/or collapse.

Please see additional Important Safety Information throughout.
Please see Full Prescribing Information [here](#).

PHARMACOSMOS
THERAPEUTICS



MonoFerric®
(ferric derisomaltose)
injection



About Monoferric (ferric derisomaltose) injection

- Monoferric provides a single, rapid infusion and is administered by intravenous infusion over at least 20 minutes as a single dose

Dosing

- For patients weighing 50 kg or more: Administer 1,000 mg of Monoferric as an intravenous infusion
- For patients weighing less than 50 kg: Administer Monoferric as 20 mg/kg actual body weight as an intravenous infusion
- Repeat Monoferric treatment if iron deficiency anemia reoccurs
- Monitor patients for signs and symptoms of hypersensitivity during and after Monoferric administration for at least 30 minutes and until clinically stable following completion of the infusion

IMPORTANT SAFETY INFORMATION (continued)

WARNINGS AND PRECAUTIONS

Hypersensitivity Reactions

Serious hypersensitivity reactions, including anaphylactic-type reactions, some of which have been life-threatening and fatal, have been reported in patients receiving Monoferric. Patients may present with shock, clinically significant hypotension, loss of consciousness, and/or collapse. Monitor patients for signs and symptoms of hypersensitivity during and after Monoferric administration for at least 30 minutes and until clinically stable following completion of the infusion. Only administer Monoferric when personnel and therapies are immediately available for the treatment of serious hypersensitivity reactions.

**Please see additional Important Safety Information throughout.
Please see Full Prescribing Information [here](#).**

Monoferric Patient Solutions® Program

MPS is committed to providing a seamless access journey to patients and healthcare providers

Live representatives are available to support patients, caregivers, and healthcare providers through the various MPS offerings:

- ✔ Benefits verification
- ✔ Claims and appeals assistance
- ✔ Sample letters of appeal and/or medical necessity
- ✔ Prior authorization assistance
- ✔ Financial assistance
- ✔ In-person support for providers' offices by providing access to a local field reimbursement manager



To learn more about this support program, or to enroll your patient, please call **1-800-992-9022**

IMPORTANT SAFETY INFORMATION (continued)

Hypersensitivity Reactions (continued)

Monoferric is contraindicated in patients with prior serious hypersensitivity reactions to Monoferric or any of its components. In clinical trials in patients with IDA and CKD, serious or severe hypersensitivity were reported in 0.3% (6/2008) of the Monoferric treated subjects. These included 3 events of hypersensitivity in 3 patients; 2 events of infusion-related reactions in 2 patients and 1 event of asthma in one patient.

Please see additional Important Safety Information throughout.
Please see Full Prescribing Information [here](#).

How to enroll in any of the available support services (Reimbursement, Patient Assistance Program, Copay):

Enrollment form options:

- You may enroll your patient online via the portal monoferricpatientsolutionsportal.com
- You may also visit monoferric-patient-solutions.com to download the editable form, complete the required fields, and fax to **1-833-888-8837**

MPS may contact you and/or your patient for additional information in order to initiate next steps.



Monoferric Patient Solutions® Patient Assistance Program

Manageable Requirements to Facilitate Access

Eligibility Criteria:

- ✓ Fall within the income guidelines*
- ✓ Uninsured or underinsured (patients with claims covered, paid or reimbursed, in whole or in part, by Medicaid, Medicare, or other federal or state healthcare programs are not eligible for this program)
- ✓ Must be 18 years or older
- ✓ Prescribed Monoferric for an on-label diagnosis
- ✓ Patient must be a resident of the United States (residency includes anyone who lives in one of the U.S. states, the District of Columbia, Puerto Rico, and U.S. Virgin Islands). Citizenship or legal status is not a requirement



To learn more, call **1-800-992-9022**
or visit monoferric-patient-solutions.com



* Total household income is at or below 300% of the federal poverty level (FPL). Visit <https://aspe.hhs.gov/poverty-guidelines>, which lists the current FPL guidelines. Pharmacosmos Therapeutics Inc. and its authorized third-party agents will use the patient's date of birth or social security number and/or additional demographic information as needed to access credit information and information derived from public and other sources to estimate income in conjunction with the eligibility determination process. As a soft credit inquiry, this option will not impact credit scores. Pharmacosmos Therapeutics Inc. and its authorized third-party agents reserve the right to ask for additional documents and information at any time. Note: Patients may retroactively qualify for assistance under the Patient Assistance Program if the patient's healthcare provider submits an explanation of benefits (EOB) statement from the patient's commercial insurance provider within 120 days of the date of service.

IMPORTANT SAFETY INFORMATION (continued)

Iron Overload

Excessive therapy with parenteral iron can lead to excess iron storage and possibly iatrogenic hemosiderosis or hemochromatosis. Monitor the hematologic response (hemoglobin and hematocrit) and iron parameters (serum ferritin and transferrin saturation) during parenteral iron therapy. Do not administer Monoferric to patients with iron overload.

Please see additional Important Safety Information throughout.
Please see Full Prescribing Information [here](#).

MonoFerric Patient Solutions® Copay Assistance Program

Patients pay as little as \$0 per dose†

Eligibility Information:



To be eligible to participate in the MPS Copay Assistance Program, the patient must:

- ✓ Have commercial health insurance (i.e., health insurance offered through an employer; NOT Medicare, Medicare Advantage, Medicaid, TRICARE, or Veteran Affairs healthcare)
- ✓ Reside in the United States or Puerto Rico
- ✓ Be treated by a healthcare professional in the United States or Puerto Rico
- ✓ Be 18 years of age or older
- ✓ Be prescribed MonoFerric for an on-label diagnosis



If a patient is eligible to participate:

- ✓ They will receive savings on out-of-pocket (OOP) expenses (i.e., deductible, copay, or coinsurance obligations) for MonoFerric of up to \$2,000 per dose†
- ✓ Copay assistance may be applied retroactively to prescription costs that occurred within 120 days prior to the date of enrollment and the patient met all of the eligibility criteria at the time of the infusion



To find out if a patient is eligible to enroll in the MPS Program, call **1-800-992-9022** or visit the online portal at **monoferricpatientsolutionsportal.com**

†If IDA returns within the coverage period, you would receive an annual maximum savings on OOP expenses of up to \$4,000. Additional restrictions apply. Please see full Terms and Conditions at monoferric-patient-solutions.com.

IMPORTANT SAFETY INFORMATION (continued)

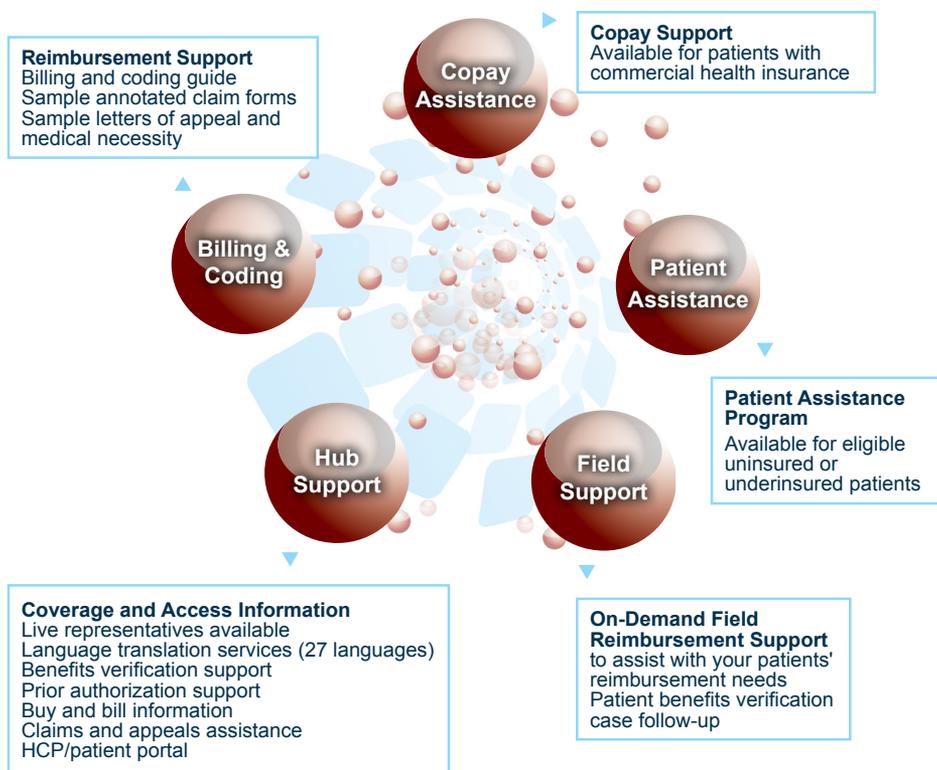
ADVERSE REACTIONS

Adverse reactions were reported in 8.6% (172/2008) of patients treated with MonoFerric. Adverse reactions related to treatment and reported by $\geq 1\%$ of the treated patients were nausea (1.2%) and rash (1%). Adjudicated serious or severe hypersensitivity reactions were reported in 6/2008 (0.3%) patients in the MonoFerric group. Hypophosphatemia (serum phosphate < 2.0 mg/dL) was reported in 3.5% of MonoFerric-treated patients in Trials 1 & 2.

Please see additional Important Safety Information throughout.
Please see Full Prescribing Information [here](#).

Monoferric Patient Solutions® Program Delivers Diverse Support to Address Monoferric Treatment Needs

Rapid Response, Customized Solutions



To report adverse events, please contact Pharmacosmos at 1-888-828-0655.
You may also contact the FDA at www.fda.gov/medwatch or call 1-800-FDA-1088.

Please see Full Prescribing Information [here](#).



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PHARMACOSMOS
THERAPEUTICS

Monoferric®
(ferric derisomaltose)
injection