



Patient Name: _____ DOB: ____/____/____

Date of Last Infusion: ____/____/____ Height _____ Weight _____

Infusion Location: (state and Site) _____

Feraheme® (ferumoxytol) Infusion Orders

Diagnoses- please provide ICD-10 and description in spaces provided

BOTH PRIMARY AND SECONDARY DIAGNOSES ARE REQUIRED

____ Iron Deficiency Anemia (ICD-10)	A N D	_____ (ICD-10) Description of underlying disease
____ Chronic Kidney Disease: Stage _____ (ICD-10)	A N D	c D63.1 Anemia in CKD
____ Description of underlying disease (ICD-10)	A N D	c D63.8 Anemia in Chronic Disease
Other: (BOTH primary and secondary dx including ICD-10 codes):		

Hold infusion and notify provider for:

- ☐ Hypotension (SBP less than 90 mmHg).
- ☐ History of allergy to other IV iron product

- Place patient in reclined or semi-reclined position and record vital signs before, after and every 30mins.
- Instruct patient to complete follow-up lab testing as ordered below
- If related reaction occurs, stop infusion and initiate Hypersensitivity Reaction Management Protocol

Pre-medications: (consider in presence of risk factors for hypersensitivity reaction i.e. multiple drug allergies).

▪ Solu-Medrol 125 mg IVP once 30 minutes prior to infusion ▪ Other: _____

Dosing:

- ☐ Administer **TWO (2) DOSES of Feraheme 510 mg** separated by 3-8 days.

Dilute in 100 ml 0.9% sodium chloride and infuse over 15-30 minutes.

- ☐ Administer **SINGLE DOSE of Feraheme 510 mg**.

Dilute in 100 ml 0.9% sodium chloride and infuse over 15-30 minutes.

Observation Period Mandatory for all patients every visit. If patient refuses to stay, ROR form may be signed to waive observation period after _____ infusions:

- Monitor patient for hypersensitivity reaction for a period of 30 minutes following each infusion.
- Record vital signs prior to discharge.

Follow-up Lab Orders: At least one month following last iron infusion, draw the following:

- ☐ CBC w/diff, ferritin, transferrin saturation, TIBC

(RN: Provide patient with order and fill in date: Draw on or after ____/____/____)

Provider name (print): _____ Date: _____

Provider signature: _____ Time: _____