Referral Status: 🗆 New Patient 📮 Updated Patient Information



CMC Infusion Center- Referral Face Sheet

Patient Name:	Patient D.O.B			
Current Height:	Current Weight			
Patient Phone # ()	Gender			
Patient Address				
CityState	_Zip Code			
Related Diagnosis Code (ICD-10 code):				
Referring Provider (Please Print)				
Office Phone # ()	Office Contact:			

If referring provider and/or patient is external to Conway Medical Center. Please include:

- Most recent history and physical including:
 - \circ Comprehensive medication list
 - o Past medical history
 - \circ Related past/ failed therapies (with dates)
 - Known allergies
- Most recent labs / test result (Ex. CBC, PPD, Hepatitis screening, etc.)
- Insurance Information

Please fax requested documentation, face sheet, and treatment plan to **843-234-5460** attention CMC Medication Management Specialist. Please feel free to call 843-234-8575 with any questions.

We look forward to the opportunity to be a part of your patient's treatment journey at Conway Medical Center.

Referral Status 🛛 New 🗖 Order Change 🗖 Order Renewal

IV Iron Treatment Plan



Patient Name:		Patient DOB
General		
Diagnosis		ICD-10 Code
Pt. Weight	Pt Height	Known Allergies
Requested Start Date		
Has patient previously re	ceived this medicatio	on \square No \square Yes, if so, date of last infusion//
		lobin and hematocrit results (minimum) in addition to any additional nce authorization process. H&H must have been obtained within the
Indication for treatment	<u>::</u>	
Oral iron ineffective	Oral iron not tolerat	ted \square Chronic kidney disease \square Oral iron drug interaction
•	th 1 to 12)	Temperature on admission to the unit and prior to infusion HR, BP every 30
Vital Signs (Month as needed		ces adverse reaction until one hour after symptom resolution
Pre- Treatment Medic	ations	
Benadryl (Month 1 to 12)	g, IV Push, Injection, I	ster 30 minutes before iron infusion
SOLU-Medrol (Month 1 to	g, IV Push, Injection, I Comments: Adminis	Day of Tx ster 30 minutes before iron infusion
0 125 m	Comments: Adminis g, IV Push, Powder-Ii	ster 30 minutes before iron infusion
Prescriber Signature	(No Stamped Sig	gnatures or Electronic Signatures)
Provider Signature		Provider Name (please print)
Date / /	Patient Nar	me Patient DOB

IV Iron Treatment Plan

Iron Sucrose (Venofer®)

200 mg, IV Piggyback, Injection, Day of Tx followed by _____ additional dose(s) {Max=4} Comments: In 100 mL NS over 30 minutes. Maximum dose is 1000mg in 5 divided doses over a 14 day period. Observe patients for at least 30 minutes post infusion

Ferric Carboxymaltose (Injectafer®)

750 mg, IV Piggyback, Soln-IV, Day of Tx [Greater Than or Equal To 50 kg] followed by __1____ additional dose(s) given 7 days apart Comments: administer as an IV infusion, dilute up to 750 mg in a maximum of 250 mL of 0.9%

sodium chloride injection to a concentration of 2-4 mg/mL; concentration should be =2 mg/mL

☐ 15 mg/kg, IV Piggyback, Soln-IV, Once **[Less Than 50 kg]** followed by **__1_** additional dose(s) given 7 days apart

Comments: administer as an IV infusion, dilute in a maximum of 250 mL of 0.9% sodium chloride injection to a concentration of 2-4 mg/mL; concentration should be =2 mg/mL

Ferumoxytol (Feraheme®)

□ 510 mg, IV Piggyback, Injection, Day of Tx with a second dose to be given 3-8 days later Comments: Mix in 100 mL NS, infuse over 30 minutes. Infuse at no greater than 1 mL/sec.

Communication Order (Month 1 to 12)

Patient must stay in infusion area for 30 min post infusion. If vital signs normal may be discharged to home. If abnormal, call physician.

Notify Provider (Month 1 to 12)

for fever/chills, chest pain, hypotension, hypertension, dyspnea, Pruritis, urticaria, persistent flushing Temperature > 100 or HR <50 or > 130

INF Infusion Room Orders Subphase (Month 1 to 12)

heparin flush 100 units/mL intravenous solution (Month 1 to 12)

5 mL, Intracatheter, Injection, Once, PRN other (see comment) Comments: To maintain central line patency for implanted ports, flush once monthly or prior to deaccess

Labs:

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Ordering provider acknowledges responsibility for laboratory monitoring including but not limited to CBC, Ferritin, and Iron studies as clinically appropriate. Note: Iron studies are not recommended to be completed sooner than 30 days following the completion of the iron infusions.

Provider Initials:

Prescriber Signature (No Stamped Signatures or Electronic Signatures)

Provider Signature	 _ Provider Name (please print)	

Date___/___/ Patient Name_____ Patient DOB_____