


ACTION	Intravenous Iron Therapy Outpatient Clinic Order Set
Allergies	
<input type="checkbox"/> NKA <input type="checkbox"/> Allergies confirmed within Meditech <input type="checkbox"/> New Allergies to be entered into Meditech: _____	
Criteria for IV Iron	
<input type="checkbox"/> Failed oral iron therapy or unable to take oral therapy <input type="checkbox"/> Symptomatic <input type="checkbox"/> Evidence of iron deficiency anemia (IDA) (i.e. ferritin less than 10 ug/L or iron less than 10 umol/L; TIBC above upper limit of normal; iron saturation value below lower limit of normal)	
Clinic Appointment Booking	
<input checked="" type="checkbox"/> Book clinic appointment for initial IV Iron administration <input type="checkbox"/> Urgent _____ <input type="checkbox"/> Routine _____ <input type="checkbox"/> Book subsequent clinic appointment <input type="checkbox"/> q _____ weeks from initial visit x _____ <input type="checkbox"/> EAP completed (venofer) <input type="checkbox"/> Prescription written [Note: monoferric LU Code 610] (please send copy with this order set to the clinic) <input type="checkbox"/> Prescription faxed to retail pharmacy	
Lab Investigations	
<input type="checkbox"/> CBC <input type="checkbox"/> Iron <input type="checkbox"/> Ferritin To be drawn: _____ <input type="checkbox"/> Other: _____	
Monitoring	
<input checked="" type="checkbox"/> Observe patient during infusion and for 30 min post infusion for signs of hypotension or hypersensitivity reaction <input checked="" type="checkbox"/> Temp, HR, RR, BP pre-treatment, post infusion, and after 30 min observation period	
IV Therapy	
<input checked="" type="checkbox"/> 2/3 – 1/3 or 0.9% Sodium Chloride TKVO. Adjust rate as needed.	

Submitter Name: _____	Date & Time _____	Order Verified by Signature: _____	Date & Time _____
Co-Signer Signature: _____	Date & Time _____	Scanner Signature: _____	Date & Time _____
		Transcriber Signature: _____	Date & Time _____

ACTION

Intravenous Iron Therapy Outpatient Clinic Order Set

Iron Therapy

Iron Sucrose (Venofer) (Patient to supply)

Previous dose received in ED = _____ mg on _____ (date)

(total cumulative dose = 1,000 mg = 2 doses x 300 mg, then 1 dose x 400 mg; account for dose given in Emergency Dept, if applicable). Space doses 2 weeks apart.

Iron Sucrose 300 mg in 250 mL 0.9% Sodium Chloride IV over 2 hours x _____ dose(s) **then**

Iron Sucrose 400 mg in 500 mL 0.9% Sodium Chloride over 3 hours x 1 dose

OR

Iron Sucrose _____

Repeat q 2 weeks from initial visit x _____ **OR** q _____ weeks from initial visit x _____

OR

Iron Sucrose 200 mg in 100 mL 0.9% Sodium Chloride IV over 1 hour (consider for patients less than 50 kg)

Repeat every 2 weeks x _____ dose(s)

Iron Isomaltoside (Monoferic) (Patient to supply)

Contraindications include but are not limited to: pregnancy, prior allergic reaction to any IV iron product and multiple drug allergies

Hemoglobin (g/L)	Weight less than 50 kg	Weight 50-69 kg	Weight 70 kg or more
Hgb 100 or greater	500 mg	1,000 mg	1,500 mg (give as 1,000 mg then 500 mg 7 days later)
Hgb less than 100	1,000 mg	1,500 mg (give as 1,000 mg then 500 mg 7 days later)	2,000 mg (give as 1,000 mg x 2 doses 7 days apart)

Iron Isomaltoside 1,000 mg in 100 mL 0.9% Sodium Chloride IV x 1 dose. Starting rate 25 mL/h x 5 min then, if tolerated, give remainder over 60 min

Repeat x 1 dose in 7 days **OR** _____ days after initial dose (total dose = 2,000 mg)

Iron Isomaltoside 500 mg in 50 mL 0.9% Sodium Chloride IV x 1 dose. Starting rate 25 mL/h x 5 min then, if tolerated, give remainder over 30 min

Repeat x 1 dose in 7 days **OR** _____ days after initial dose (total dose = 1,000 mg)

Give x 1 dose in 7 days **OR** _____ days after initial 1,000 mg dose (total dose = 1,500 mg)

Submitter Name: _____

Date & Time _____

Order Verified by Signature: _____

Date & Time _____

Co-Signer Signature: _____

Date & Time _____

Scanner Signature: _____

Date & Time _____

Transcriber Signature: _____

Date & Time _____



