

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



ACTION	Intra	Intravenous Iron Therapy Outpatient Clinic Order Set							
	Iron Therapy								
	<ul> <li>Iron Sucrose (Venofer) (Patient to supply)</li> <li>Previous dose received in ED = mg on (date)</li> <li>(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</li> <li>OR</li> <li>Iron Sucrose OR </li> <li>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)</li> <li>Iron Isomaltoside (Monoferric) (Patient to supply)</li> </ul>								
									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 1,00 500 mg 7 days later)	00 mg then				
	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 1,00 doses 7 days apart)	00 mg x 2				
	<ul> <li>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</li> <li>Repeat x 1 dose in 7 days OR days OR days after initial dose (total dose = 2,000 mg)</li> <li>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</li> <li>Repeat x 1 dose in 7 days OR days OR days after initial dose (total dose = 1,000 mg)</li> <li>Give x 1 dose in 7 days OR days after initial 1,000 mg dose (total dose = 1,500 mg)</li> </ul>								
	Submitter Name: Co-Signer Signature:		e Order Verified by S e Scanner Signature		Date & Time				
			Transcriber Signate		Date & Time				



	Intravenous Ir	on Therapy Outpati	ient Clinic Order Set			
Adv	Adverse Reaction Management					
ph If s ar Ac Ce	in patients with Fishbane reaction significant hypotension occurs (Bl sysician signs of anaphylactic reaction (pe gioedema) – <b>STOP</b> infusion IMM otify physician STAT.	n P drop of 25 mmHg or more from p ersistent significant hypotension, sy IEDIATELY and administer <b>EPINEF</b> RN for pain or fever (start 4 hours a for urticarial/pruritus	present as it may contribute to hypotension ore-treatment value) – stop infusion and notify ncope, urticaria, pruritus, bronchospasm, PHrine (1 mg/mL) 0.5 mg = 0.5 mL IM STAT fter premedication dose if given)			
Oth	er Drug Therapy					
Ad   dij   Ce   Fa   Hy	as premedication only if patient has estaminophen 650 mg PO x 1 dos ohenhydr <b>AMINE</b> 25 mg IV x 1 dos estirizine 10 mg PO x 1 dose PRN umotidine 20 mg PO OR IV x 1 dose vdrocortisone 100 mg IV x 1 dose menhy <b>DRINATE</b> 25-50 mg PO/IV	se PRN ose PRN 9 PRN				
Disc	Discharge Instructions					
Instru	ct patient to follow up with 🗌 Pre	escribing Physician or 🗌 Family Pr	ovider			
Add	Additional Orders					
DO N	OT USE: <, >, SC, SQ, SUBQ, U	, IU, zero <u>after</u> decimal (write 1 mg	) ALWAYS USE: zero <u>before</u> decimal (0.5			
er Nam		e & Time Order Verified	by Signature: Date & Time			

Co-Signer Signature:

Date & Time

Scanner Signature:

Transcriber Signature:

Date & Time

Date & Time



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