

Oregon Health & Science University
Hospital and Clinics Provider's Orders



ADULT AMBULATORY INFUSION ORDER Epoetin alfa-epbx (RETACRIT) Injection Maintenance for Oncology Patients ACCOUNT NO.
MED. REC. NO.
NAME
BIRTHDATE

Patient Identification

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ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK (✓) TO BE ACTIVE.

Weight:		kg	Height: _		cm						
Allergies:											
	Code:										
Treatment	Start Date: _			Patient to	follow up v	with provid	der on d	ate:			
This ord	n will expiro ler set is fo N order set ions	r MAIN	ITENANC	E DOSING	G ONLY. F	Patients s	should l	nave re	ceived f	first dos	
□ Ch	ON: <i>(Must o</i> emotherapy mptomatic a	-induc	ed ánemia		elodysplast	ic syndroi	me (MD	S)			
GUIDELIN	IES FOR OF	RDERI	NG								
	nd FACE SI										
	ISU's formul					,	•		`	,	

- 2. OHSU's formulary erythropoiesis stimulating agent (ESA) is darbepoetin alfa (ARANESP). All orders for epoetin alfa-epbx (RETACRIT) will be converted to darbepoetin alfa using equivalent therapeutic interchange dosing listed in the table below. **Providers who prefer to use epoetin alfa-epbx must specify a reason for its use. REASON FOR EPOETIN USE:**
- 3. Patients receiving concurrent treatment with Iron Sucrose (VENOFER) and/or Vitamin B12 cannot receive ESA treatment on the same day. Patients may be on prophylactic oral iron supplementation concurrent with ESA treatment as long as supplementation for the prevention of iron deficiency is necessary due to ESA therapy alone.
- 4. Serum ferritin and transferrin saturation (TSAT) must be performed every month during initial (ESA) treatment and at least every 3 months during stable ESA treatment (serum ferritin greater than 100 ng/mL, and TSAT greater than or equal to 20%). Therapy with epoetin alfa-epbx may continue only if hemoglobin DOES NOT equal or exceed 11 g/dL.
- 5. **For patients with chemotherapy-induced anemia**: The medical record must document the provider's rationale for determining the anemia is "chemotherapy-induced." Anemia must be secondary to myelosuppressive anticancer chemotherapy in solid tumors, multiple myeloma, lymphoma, or lymphocytic leukemia. Treatment should be limited to the 8 weeks following myelosuppressive chemotherapy.
- 6. **For patients with symptomatic anemia from MDS**: The patient must be symptomatic and his/her life expectancy must be >3 months. The medical record must display documentation that a bone marrow biopsy has been reviewed by a provider and is consistent with the diagnosis of MDS. The marrow blast count must be <5%



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LAB2:	
	Hemoglobin & Hematocrit, Routine, ONCE, (weekly) or every (visit)(days)(weeks)(months) -
	Circle One
	CMP, Routine, ONCE, (weekly) or every (visit)(days)(weeks)(months) – Circle One
	Ferritin (serum), Routine, ONCE, every 12 weeks
	Iron and TIBC (serum), Routine, ONCE, every 12 weeks

NURSING ORDERS:

1. Patients receiving concurrent treatment with Iron Sucrose (VENOFER) and/or Vitamin B12 cannot receive ESA treatment on the same day.

☐ Labs already drawn. Date: _____ (Must have been drawn at least 14 days after the last dose)

- 2. TREATMENT PARAMETERS Hold treatment and call provider if hemoglobin is greater than or equal to 11, serum ferritin is less than or equal to 100 ng/mL, transferrin saturation is less than 20% or if blood pressure is greater than 180 systolic or 100 diastolic
- 3. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.

OTHER:

Conversion from epoetin alfa-epbx (RETACRIT) to darbepoetin alfa (ARANESP): Initial adult dosing

Epoetin alfa-epbx dose	Darbepoetin alfa dose (mcg/week)
(units/week)	, ,
<1500	6.25
1500-2499	6.25
2500-4999	12.5
5000-10,999	25
11,000-17,999	40
18,000-33,999	60
34,000-89,999	100
≥90,000	200

In patients receiving epoetin alfa-epbx 2-3 times weekly, darbepoetin should be given once weekly. If epoetin is administered once weekly, darbepoetin should be given once every 2 weeks. Darbepoetin dosing every 2 weeks should be determined by adding the 2 weekly epoetin alfa-epbx doses, then convert to appropriate corresponding darbepoetin dose. Doses should be titrated to hemoglobin response thereafter



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MEDICATIONS:

epoetin alfa-epbx (RETACRIT), subcutaneous, ONCE
 Pharmacist will round dose to nearest vial size if within 10% of original dose during verification

Weight based regimen:

weignt ba	asea regime	en:		
Dose				
		units/kg =	units	
Interv	al:	-		
	Weekly x _	weeks		
		times per week x		weeks
Fixed dos	se regimens	s:		
Dose:				
	2,000 units	•		
	3,000 units			
	4,000 units			
	10,000 unit			
	20,000 unit			
	40,000 unit			
Interv	•	.0		
	Weekly x	weeks		
		times per week x		weeks



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that corresponds with state where you provide state if not Oregon);	medicine in: □ Oregon □(eare to patient and where you are currently license	ed. Specify			
My physician license Number is # PRESCRIPTION); and I am acting within my so medication described above for the patient iden	(MUST BE COMPLETED TO BE A Vope of practice and authorized by law to order Infutified on this form.	' <u>ALID</u> usion of the			
Provider signature:	Date/Time:				
Printed Name:	Phone: Fax:				
OLC Central Intake Nurse: Phone: 971-262-9645 (providers only) Fax: 503 Please check the appropriate box for the pa					
□ Beaverton OHSU Knight Cancer Institute 15700 SW Greystone Court Beaverton, OR 97006 Phone number: 971-262-9000 Fax number: 503-346-8058	NW Portland Legacy Good Samaritan campus Medical Office Building 3, Suite 150 1130 NW 22nd Ave. Portland, OR 97210 Phone number: 971-262-9600 Fax number: 503-346-8058				
☐ Gresham Legacy Mount Hood campus Medical Office Building 3, Suite 140 24988 SE Stark Gresham, OR 97030 Phone number: 971-262-9500 Fax number: 503-346-8058	□ Tualatin Legacy Meridian Park campus Medical Office Building 2, Suite 140 19260 SW 65th Ave. Tualatin, OR 97062 Phone number: 971-262-9700 Fax number: 503-346-8058				

Infusion orders located at: www.ohsuknight.com/infusionorders