

<div style="display: flex; align-items: center;"> <div> <b>Oregon Health &amp; Science University</b>  <b>Hospital and Clinics Provider's Orders</b> </div> </div> <div style="display: flex; align-items: center; margin-top: 10px;"> <div style="writing-mode: vertical-rl; transform: rotate(180deg); font-size: small; margin-right: 5px;">PO7071</div> </div> <div style="text-align: center; margin-top: 10px;"> <b>ADULT AMBULATORY INFUSION ORDER</b>  <b>Epoetin Alfa-epbx (RETACRIT)</b>  <b>Injection</b>        Page 1 of 4     </div>	<div style="display: flex; flex-direction: column; justify-content: space-between;"> <div>ACCOUNT NO.</div> <div>MED. REC. NO.</div> <div>NAME</div> <div>BIRTHDATE</div> </div> <div style="text-align: right; font-size: x-small; margin-top: 20px;">Patient Identification</div>
<b>ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK ( ✓ ) TO BE ACTIVE.</b>	

Weight: \_\_\_\_\_ kg      Height: \_\_\_\_\_ cm  
 Allergies: \_\_\_\_\_  
 Diagnosis Code: \_\_\_\_\_  
 Treatment Start Date: \_\_\_\_\_ Patient to follow up with provider on date: \_\_\_\_\_

**\*\*This plan will expire after 365 days at which time a new order will need to be placed\*\***

**INDICATION: (Must check one)**

- ☐ Chemotherapy-induced anemia  
**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.
- ☐ Symptomatic anemia associated with myelodysplastic syndrome (MDS)  
**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%.
- ☐ Anemia of Chronic Kidney Disease (CKD)  
**For patients with anemia of CKD:** The medical record must display documentation that anemia is clearly attributed to a CKD diagnosis. The specific CKD stage must be moderate (stage III) to end stage.

**GUIDELINES FOR ORDERING:**

1. Send **FACE SHEET and H&P or most recent chart note detailing treatment indication and plan.**
2. Hemoglobin and hematocrit must be obtained within 1 week of therapy initiation. Hemoglobin must be < 10 g/dL or hematocrit must be < 30% prior to initiation.
3. Serum ferritin and transferrin saturation (TSAT) must be performed every 3 months during erythropoiesis stimulating agent (ESA) treatment (serum ferritin ≥ 100 ng/mL, and TSAT ≥ 20%). Therapy with ESA may continue only if hemoglobin meets maintenance treatment parameters per indication.
4. All patients must be negative when evaluated for blood loss, hemolysis, and bone marrow fibrosis prior to initiation of therapy. Providers must assess and replete iron, folate, and Vitamin B12 prior to any treatment with ESA.
5. Patients cannot receive Iron Sucrose (VENOFER) and/or Vitamin B12 on the same day as ESA treatment. 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.



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

ADULT AMBULATORY INFUSION ORDER

**Epoetin Alfa-epbx (RETACRIT)**

**Injection**

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ACCOUNT NO.

MED. REC. NO.

NAME

BIRTHDATE

*Patient Identification*

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

**LABS:**

- Hemoglobin & Hematocrit, Routine, ONCE, every visit
- Ferritin, once clinic collect, comment as needed if not resulted in last 90 days, interval quarterly
- Iron and TIBC, once clinic collect, comment as needed if not resulted in last 90 days, interval quarterly

☐ Labs already drawn. Date: \_\_\_\_\_ (Labs scanned with orders)

**NURSING ORDERS:**

1. Patients cannot receive Iron Sucrose (VENOFER) and/or Vitamin B12 on same day as ESA treatment.
2. Do not obtain ferritin or transferrin saturation (TSAT) on the same day as ESA treatment.
3. OK to give erythropoiesis-stimulating agents on the same day as blood transfusions.
4. TREATMENT PARAMETERS
  - a. Initiation dose: hemoglobin and hematocrit must be obtained within 1 week. Hemoglobin must be less than 10 g/dL or hematocrit must be less than 30% prior to initiation.
  - b. Maintenance doses: hemoglobin must be obtained within 72 hours of each individual dose. Hemoglobin must be:
    - i. Chemotherapy induced anemia: Hgb less than 10 g/dL
    - ii. Anemia due to MDS: Hgb less than 12 g/dL
    - iii. Anemia due to CKD: Hgb less than or equal to 11 g/dL
    - iv. Other: Hgb less than \_\_\_\_\_ g/dL
  - c. Ferritin should be greater than or equal to 100 ng/mL and transferrin saturation should be greater than or equal to 20%.
  - d. Hold treatment and call provider if lab parameters are not met or if blood pressure is greater than 180 mm Hg systolic or 100 mm Hg diastolic.

**MEDICATIONS: (must check one if provider managed - opt out of pharmacy managed protocol)**

**Epoetin alfa-epbx (RETACRIT), subcutaneous, ONCE**

Initiate first dose within 1 week of obtaining baseline labs.

**PHARMACY MANAGED PROTOCOL / OPT OUT: (Must check one)**

- ☐ Pharmacist managed dosing protocol (**OHSU infusion centers only**). Do NOT indicate specific dose below, pharmacy to manage per institutional protocol.
- ☐ Provider managed dosing (indicated dosing below)

\*\*\*Fixed dose regimen\*\*\*

**Fixed dose regimens: (must check one)**

- ☐ 2,000 units
- ☐ 3,000 units
- ☐ 4,000 units
- ☐ 10,000 units
- ☐ 20,000 units
- ☐ 40,000 units



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**Epoetin Alfa-epbx (RETACRIT)  
Injection**  
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**Interval:**

- ☐ Once  
☐ Weekly x \_\_\_\_\_ weeks  
☐ \_\_\_\_\_ times per week x \_\_\_\_\_ week

Epoetin								
Indication	Weight	Dose level 0 (Starting Dose)	Dose Decrease		Dose Increase			
			Dose level -1	Dose level -2	Dose level +1	Dose level +2	Adjunctive agent	Notes
<b>MDS</b>	≥ 60 kg (or flat dose)	40,000 units weekly	30,000 units weekly	22,000 units weekly	50,000 units weekly	60,000 units weekly	By week 12 if no response, contact provider to add GCSF 300 mcg 1-3x per week	By week 16 if no increase in Hgb by 1.5 or reach target of 10-12 g/dL or decrease in transfusion needs discontinue
	< 60 kg	24,000 units weekly	18,000 units weekly	14,000 units weekly	40,000 units weekly	60,000 units weekly		
<b>Chemo induced</b>	≥ 60 kg (or flat dose)	40,000 units weekly	30,000 units weekly	22,000 units weekly	60,000 units weekly			By week 8 if no improvement in Hgb, maintain lowest dose to avoid transfusions, if no improvement in transfusion requirements discontinue
	< 60 kg	24,000 units weekly	18,000 units weekly	14,000 units weekly	40,000 units weekly			
<b>CKD (no HD)</b>	≥ 60 kg (or flat dose)	20,000 units every 2 weeks	14,000 units every 2 weeks	10,000 units every 2 weeks	24,000 units every 2 weeks	30,000 units every 2 weeks		By week 12 if no improvement in Hgb, maintain lowest dose to avoid transfusions, if no improvement in transfusion requirements discontinue
	< 60 kg	10,000 units every 2 weeks	8,000 units every 2 weeks	6,000 units every 2 weeks	12,000 units every 2 weeks	16,000 units every 2 weeks		



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**HYPERSENSITIVITY MEDICATIONS:**

1. NURSING COMMUNICATION – If hypersensitivity or infusion reactions develop, temporarily hold the infusion and notify provider immediately. Administer emergency medications per the Treatment Algorithm for Acute Infusion Reaction (OHSU HC-PAT-133-GUD, HMC C-132). Refer to algorithm for symptom monitoring and continuously assess as grade of severity may progress.
2. diphenhydramine (BENADRYL) injection, 25-50 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
3. EPINEPHrine HCl (ADRENALIN) injection, 0.3 mg, intramuscular, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
4. hydrocortisone sodium succinate (SOLU-CORTEF) injection, 100 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
5. famotidine (PEPCID) injection, 20 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction

**By signing below, I represent the following:**

I am responsible for the care of the patient (*who is identified at the top of this form*);

I hold an active, unrestricted license to practice medicine in: ☐ Oregon ☐ \_\_\_\_\_ (*check box that corresponds with state where you provide care to patient and where you are currently licensed. Specify state if not Oregon*);

**My physician license Number is # \_\_\_\_\_ (MUST BE COMPLETED TO BE A VALID PRESCRIPTION);** and I am acting within my scope of practice and authorized by law to order Infusion of the medication described above for the patient identified on this form.

**Provider signature:** \_\_\_\_\_ **Date/Time:** \_\_\_\_\_

**Printed Name:** \_\_\_\_\_ **Phone:** \_\_\_\_\_ **Fax:** \_\_\_\_\_

**INFUSION REFERRAL TEAM**

Phone (providers only)  
(971) 262-9645

Fax completed orders to  
(503) 346-8058

Infusion orders located at:  
[www.ohsuksnight.com/infusionorders](http://www.ohsuksnight.com/infusionorders)

☒ **Please indicate the patient's preferred clinic location below**

- |   |   |
|---|---|
| <input type="checkbox"/> <b>BEAVERTON</b><br>OHSU Knight Cancer Institute   | 15700 SW Greystone Court<br>Beaverton OR 97006                                |
| <input type="checkbox"/> <b>NW PORTLAND</b><br>Legacy Good Samaritan campus | Medical Office Building 3 – Suite 150<br>1130 NW 22nd Ave, Portland OR 97210  |
| <input type="checkbox"/> <b>GRESHAM</b><br>Legacy Mount Hood campus         | Medical Office Building 3 – Suite 140<br>24988 SE Stark, Gresham OR 97030     |
| <input type="checkbox"/> <b>TUALATIN</b><br>Legacy Meridian Park campus     | Medical Office Building 2 – Suite 140<br>19260 SW 65th Ave, Tualatin OR 97062 |