



**BlueCross
BlueShield**

Federal Employee Program

**EPOETIN ALFA
PRIOR APPROVAL REQUEST**

Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the physician portion and submit this completed form.

Send completed form to:
Service Benefit Plan
Prior Approval
P.O. Box 52080 MC 139
Phoenix, AZ 85072-2080
Attn: Clinical Services
Fax: **1-877-378-4727**

Patient Information (required)				Provider Information (required)		
Date:				Provider Name:		
Patient Name:				Specialty:		NPI:
Date of Birth:	Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female			Office Phone:		Office Fax:
Street Address:				Office Street Address:		
City:	State:	Zip:		City:	State:	Zip:
Patient ID:	<div style="border: 1px solid black; padding: 2px;"> R </div>			Physician Signature:		
PHYSICIAN COMPLETES						

Retacrit (epoetin alfa-epbx)

****Check www.fepblue.org/formulary to confirm which medication is part of the patient's benefit**

NOTE: Form must be completed in its **entirety for processing**

Note: Approval cannot be given unless all lab values are provided for the diagnosis chosen

Is this request for brand or generic? ☐ Brand ☐ Generic

1. Is Retacrit being used in combination with another erythropoiesis stimulating agent (ESA)? ☐ Yes* ☐ No

****If YES***, please specify the medication: _____

2. Has the patient been on Retacrit continuously for the last **4 months**, excluding samples? ☐ Yes ☐ No*

****If NO***, is Retacrit being requested as a change from Procrit to allow the member access to their copay benefit? ☐ Yes ☐ No

3. What is the patient's diagnosis?

- ☐ Allogeneic bone marrow transplantation
 ☐ Anemia associated with Hepatitis C (HCV) treatment
☐ Myelodysplastic syndrome
 ☐ Anemia associated with Rheumatoid Arthritis (RA)/rheumatic disease
☐ Anemia associated with chronic renal failure

a. What is the patient's serum ferritin level in nanograms per milliliter (ng/mL)? _____ ng/mL

b. Have both the serum ferritin level and hemoglobin been obtained within the past 3 months? ☐ Yes ☐ No

c. Has the patient been on Retacrit continuously for the last **4 months**, excluding samples? ***Please select answer below***

☐ **NO** – this is **INITIATION** of therapy, please answer the following questions:

i. Is the patient on dialysis? ***Please select answer below:***

☐ **Yes:** What is the patient's *hemoglobin level in grams per deciliter (g/dL)? _____ g/dL

****If hemoglobin level is greater than or equal to 10 g/dL***, will the dose be held or reduced until the hemoglobin level is less than 10 grams per deciliter (g/dL)? ☐ Yes ☐ No

☐ **No:** What is the patient's *hemoglobin level in grams per deciliter (g/dL)? _____ g/dL

****If hemoglobin level is greater than or equal to 11g/dL***, will the dose be held or reduced until the hemoglobin level is less than 11 grams per deciliter (g/dL)? ☐ Yes ☐ No

☐ **YES** – this is a PA renewal for **CONTINUATION** of therapy, please answer the following question(s):

i. What is the patient's *hemoglobin level in grams per deciliter (g/dL)? _____ g/dL

****If hemoglobin level greater than 11 g/dL***, will the dose be held or reduced until the hemoglobin level is less than or equal to 11 grams per deciliter (g/dL)? ☐ Yes ☐ No

☐ Anemia in patients scheduled to undergo elective, non-cardiac, nonvascular surgery

a. What is the patient's hemoglobin level in grams per deciliter (g/dL)? _____ g/dL

PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL DIAGNOSES

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PAGE 2 - PHYSICIAN COMPLETES

Patient Name: _____ DOB: _____ Patient ID: R _____

☐ Anemia secondary to chemotherapy

a. Is the patient receiving concomitant myelosuppressive therapy? ☐ Yes ☐ No

b. Are there 2 or more additional months of chemotherapy planned for the patient? ☐ Yes ☐ No

c. Does the prescriber agree to discontinue Retacrit upon completion of the chemotherapy? ☐ Yes ☐ No

d. Does the prescriber agree that transfusions are **NOT** an option for treatment (i.e., end stage organ failure, chronic kidney disease (CKD), and high risk bacterial infections)? ☐ Yes ☐ No

☐ Anemia secondary to zidovudine-treated Human Immunodeficiency Virus (HIV) patients

a. Are the patient's endogenous serum erythropoietin levels less than or equal to 500 milliunits per milliliter (mU/mL)? ☐ Yes ☐ No

☐ Other diagnosis (*please specify*): _____

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Message:

Attached is a Prior Authorization request form.

For your convenience, there are 3 ways to complete a Prior Authorization request:

Electronically Online (ePA) Results in 2-3 minutes FASTEST AND EASIEST	Now you can get responses to drug Prior Authorization requests securely online. Online submissions may receive instant responses and do not require faxing or phone calls. Requests can be made 24 hours a day, 7 days a week. For more information on electronic prior authorization (ePA) and to register, go to Caremark.com/ePA .
Phone (4-5 minutes for response)	The FEP Clinical Call Center can be reached at (877)-727-3784 between the hours of 7AM-9PM Eastern Time. A live representative will assist with the Prior Authorization, asking for the same information contained on the attached form. Please review the form and have your answers ready for faster service. The process over the phone takes on average between 4 and 5 minutes.
Fax (3-5 days for response)	Fax the attached form to (877)-378-4727 . Requests sent via fax will be processed and responded to within 5 business days. The form must be filled out completely, if there is any missing information the Prior Authorization request cannot be processed. <u>Please only fax the completed form once as duplicate submissions may delay processing times.</u>

**faster...
easier...
better...**

Introducing ePA! Online Prior Authorizations in minutes through **Caremark.com/ePA**. Sign up today!

CVS/caremark 