

BlueShield. EPOETIN ALFA Federal Employee Program. PRIOR APPROVAL REQUEST

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

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.

Patient Information (required)			Provider Information (required)				
Date:			Provider Name:				
Patient Name:			Specialty:	NPI:			
Date of Birth:	Date of Birth: Sex: □Male □Female		Office Phone:	Office Fax:			
Street Address:			Office Street Address:				
City:	State:	Zip:	City:	State:	Zip:		
Patient ID: R	1 1		Physician Signature:	1			
PHYSICIAN COMPLETES							
For Standard Option patients Retacrit is a preferred product. Please consider prescribing the preferred product. Standard Option patients who switch to a preferred product can receive up to 2 fills without a copay in the benefit year.							
NOTE: Form must be completed in its entirety for processing							
Please select medication:	□Еро	gen (epoetin al	fa) □Pro	crit (epoetin alfa	a)		
**Check www.fepblue.org/formulary to c	confirm which medic	ation is part of the	patient's benefit				
Note: Approval cannot be given unless all lab values are provided for the diagnosis chosen							
Is this request for brand or generic? □Brand □Generic							
Procrit Request (Standard Optio	n): Would you lik	te to switch the p	patient to the preferred produ	ct, Retacrit?	es □No*		
*If NO, does the patient have an Retacrit? Please select answer be		ntraindication or	have they had an inadequate	e treatment respon	nse to		
☐Yes (please specify):							
□No: Is there a clinical reason for *If YES, please specify							
1. Is this medication being used in *If YES, please specify the m		•	ppoiesis stimulating agent (ES	SA)? □Yes* □	1 No		
2. What is the patient's diagnosis?							
☐Allogeneic bone marrow tran	splantation	□Anemia asso	ociated with Hepatitis C (HCV) treatment				
☐Myelodysplastic syndrome		□Anemia asso	ciated with Rheumatoid Arth	nritis (RA)/rheum	natic disease		
☐Anemia associated with chron a. What is the patient's serv		n nanograms per	milliliter (ng/mL)?	ng/mL			
b. Have both the serum ferr	ritin level and hen	noglobin level be	een obtained within the past t	three months?	Yes □No		
c. Has the patient been on t	his medication co	ntinuously for th	ne last 4 months, excluding s	samples? Select ar	iswer below:		
\square NO – this is INITIAT	ION of therapy, 1	please answer the	e following questions:				
i. Is the patient on di	alysis? <i>Please sele</i>	ct answer below:					
☐ Yes : What is the	e patient's *hemo	globin level in g	rams per deciliter (g/dL)?	g/dl	L		
			<i>ual to $10g/dL$</i> , will the dose to deciliter (g/dL) ? \square Yes		d until the		
\Box No : What is the	patient's *hemog	globin level in gr	rams per deciliter (g/dL)?	g/dL	1		
			<i>ual to $11g/dL$</i> , will the dose I deciliter (g/dL) ? \square Yes	be held or reduce INo	d until the		
			_	lowing question(s):		
☐ 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 is greater than $11g/dL$, will the dose be held or reduced until the hemoglobin level is less than or equal to 11 grams per deciliter (g/dL) ? \square Yes \square No							

PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL DIAGNOSES

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PAGE 2 - PHYSICIAN COMPLETES					
Patient Name:	DOB:	Patient ID: R			
☐ Anemia in patients scheduled to		•			
-	globin level in grams per deciliter	(g/dL): g/dL			
□Anemia secondary to chemothe	•				
a. Is the patient receiving cor	comitant myelosuppresive therap	y? UYes UNo			
b. Are there 2 or more addition	onal months of chemotherapy plan	nned for the patient? □Yes □No			
c. Will the prescriber agree to	discontinue use of this medication	on upon completion of the chemotherapy? □Yes □No			
1 0	hat transfusions are NOT an opticsk bacterial infections)? \(\text{\text{I}}\) Yes	on for treatment (i.e., end stage organ failure, chronic kidney \square No			
☐Anemia secondary to zidovudir	ne-treated Human Immunodeficie	ncy Virus (HIV) patients			
a. Are the patient's endogenou	s serum erythropoietin levels less th	han or equal to 500 milliunits per milliliter (mU/mL)? Yes	lNo		
☐Other diagnosis (please specify):			_		



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Message:

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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. Please only fax the completed form once as duplicate submissions may delay processing times.

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

CVS/caremark

