

Epogen, Procrit, Retacrit

Prior Authorization Request

Send completed form to: Case Review Unit CVS Caremark Prior Authorization Fax: 1-866-249-6155

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-866-814-5506**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name:	Date:
Patient's ID:	Patient's Date of Birth:
Physician's Name:	
Specialty:	NPI#:
Physician Office Telephone:	
Request Initiated For:	
☐ Continuation of therapy: Please complete the Therapy is complete: Please check box and Therapy is on hold or patient has medication	plete the following forms in entirety and fax to 866-249-6155. e following forms in entirety and fax to 866-249-6155. fax first page to 866-249-6155. a available: Please check box and fax first page to 866-249-6155. In when therapy resumes or when supply of medication is low.
 Which drug is being prescribed? ☐ Epog 	gen 🗆 Procrit 🗀 Retacrit 🗀 Other
2. What is the patient's diagnosis or reason for Anemia in chronic kidney disease (CKI Anemia due to myelosuppressive cheme Anemia in myelodysplastic syndrome (Presurgical use to reduce allogeneic blo Anemia in CHF Anemia in rheumatoid arthritis Anemia due to hepatitis C treatment Anemia due to zidovudine treatment in Anemia in patients whose religious beli Anemia in patients with primary myelosthrombocythemia myelofibrosis	otherapy MDS) od transfusions a patient with HIV infection fefs forbid blood transfusions fibrosis, post-polycythemia vera myelofibrosis, or post-essential
3. What is the ICD-10 code?	_
	vel? Exclude values due to recent transfusion uest): Hgb: g/dL Date of lab:
Current (i.e., within 30 days of request):	Hgb: g/dL Date of lab:

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Epogen, Procrit, Retacrit SGM - 10/2018.

Pre	escriber or Authorized Signature Date (mm/dd/yy)
inf	ttest that this information is accurate and true, and that documentation supporting this formation is available for review if requested by CVS Caremark or the benefit plan sponsor.
14.	What is the patient's pretreatment serum erythropoietin level? mU/mL ☐ Not available
Thr 13.	etion E: Anemia in Patients with Primary Myelofibrosis, Post-Polycythemia Vera Myelofibrosis, or Post-Essential rombocythemia Myelofibrosis-New Start ONLY Does the patient have symptomatic anemia? \(\textstyle \textstyle \textsty
	tion D: Presurgical Use to Reduce Allogeneic Blood Transfusions Is the patient scheduled to have an elective, noncardiac, nonvascular surgery? □ Yes □ No
	tion C: Anemia due to Hepatitis C Treatment Is the patient currently receiving treatment with ribavirin in combination with either interferon alfa or peginterferon alfa? □ Yes □ No
	tion B: Anemia due to Zidovudine Treatment in a Patient with HIV Infection Is the patient currently receiving treatment with zidovudine-containing medications? Yes No
9.	Is the intent of chemotherapy to cure the cancer (as opposed to palliative management or inducing remission)? \square Yes \square No
	tion A: Anemia due to Myelosuppressive Chemotherapy Does the patient have a diagnosis of a non-myeloid malignancy? □ Yes □ No
Cor	mplete the following section based on the patient's diagnosis, if applicable.
7.	How many weeks of ESA therapy has the patient completed? weeks; Document start date:
6.	At any time since the patient started ESA therapy, has the patient's Hgb increased by 1 g/dL or more? If Yes, skip to diagnosis section \square Yes \square No
5.	Has the patient received erythropoiesis stimulating agent (ESA) therapy in the previous month (within 30 days of request)? ☐ Yes ☐ No If No, skip to diagnosis section

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CVS Caremark Prior Authorization

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• Northbrook, IL 60062

Phone: 1-866-814-5506

• Fax: 1-866-249-6155

• www.caremark.com