

2020 Procrit® (epoetin alfa) Prior Authorization Request

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(You must complete all pages.)

Fax completed form to: 1-800-408-2386

For urgent requests, please call: 1-800-414-2386

Coverage Criteria:

- Medication is covered on plan if determined not to be covered under Medicare Part A or Medicare Part B AND when being prescribed for anemia due to chronic kidney disease in patients not on dialysis, anemia due to myelosuppressive anticancer chemotherapy in patients with non-myeloid malignancies in which chemotherapy is not being given with a curative intent, anemia due to zidovudine therapy in an HIV-infected patient, reduction of allogenic red blood cell transfusion in patients undergoing elective, noncardiac, nonvalvular surgery, anemia due to myelodysplastic syndromes, anemia in congestive heart failure, anemia due to rheumatoid arthritis, anemia due to hepatitis C treatment (ribavirin in combination with either interferon alfa or peginterferon alfa), anemia due to primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis, anemia in patients whose religious beliefs forbid blood transfusions and anemia in cancer patients who are undergoing palliative treatment
- **For ALL REQUESTS (except surgery)**: patients must have a PRE-TREATMENT (no erythropoietin treatment in previous month) hemoglobin (Hgb) of less than 10 g/dL (or less than 9 g/dL for anemia in CHF only).
- For patients with primary myelofibrosis, post-polycythemia vera myelofibrosis and post-essential thrombocythemia:
 - Patient must have symptomatic anemia
 - For INITIAL therapy, PRE-TREATMENT (no erythropoietin treatment in previous month) serum erythropoietin levels must be less than 500 mU/mL
- For patients undergoing surgery:
 - Procedure must be elective, noncardiac, nonvascular surgery
 - AND
 - PRE-TREATMENT hemoglobin (Hgb) level is greater than 10 g/dL but not more than 13 g/dL
- For CONTINUATION OF THERAPY for ALL patients, except surgery:
 - There must be an increase in hemoglobin (Hgb) of at least 1 g/dL after at least 12 weeks of therapy **AND**
 - For anemia due to myelosuppressive cancer chemotherapy: Current hemoglobin (Hgb) must be less than 11 g/dL
 - For all other uses, except surgery and for anemia due to chemotherapy: Current hemoglobin (Hgb) must be less than or equal to 12 g/dL

Authorization duration: 16 weeks

Patient information		Prescriber informat	ion		
Patient name		Today's date		Physician specialty	
Patient insurance ID number		Physician name		NPI/DEA number	
Patient address, city, state, ZIP		Physician address, city, state, ZIP			
Patient home telephone number		M.D. office telephone number			
Gender ☐ Male ☐ Female	Patient date of birth	M.D. office fax number			
Diagnosis and medical inform	nation				
Medication requested ☐ Procrit					

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Diagnosis and medical information (continued):						
Diagnosis and medical information (continued): Diagnosis ☐ Anemia due to chronic kidney disease (CKD) ☐ Anemia due to end stage renal disease (ESRD) with DIALYSIS ☐ Anemia due to myelosuppressive anticancer chemotherapy in	 ☐ Anemia in congestive heart failure (CHF) ☐ Anemia in rheumatoid arthritis (RA) ☐ Anemia due to hepatitis C treatment (ribavirin in combination 					
patients with non-myeloid malignancies Anemia due to zidovudine therapy in an HIV-infected patient with endogenous serum erythropoietin levels less than or equal to 500 mU/mL.	with either interferon alfa or peginterferon alfa) Anemia in primary myelofibrosis Anemia in post-polycythemia vera myelofibrosis Anemia in post-essential thrombocythemia myelofibrosis					
 ☐ Elective, noncardiac, nonvascular surgery; to reduce the need for allogeneic red blood cell transfusions among patients with perioperative hemoglobin between 10-13 g/dL who are at high risk for perioperative blood loss ☐ Anemia due to myelodysplastic syndromes (MDS) 	 ☐ Anemia in patients whose religious beliefs forbid blood transfusions ☐ Cancer patients who are undergoing palliative treatment ☐ Other diagnosis/(ICD10): 					
Strength and route of administration	Quantity	Day supply	Expected length of therapy			
Please check all boxes that apply: Restart Renewal						
Patient's home (self-administered)	1. Where is medication being administered? ☐ Patient's home (self-administered) ☐ Office administered (office supplies drug) / J CODE:					
☐ Ambulatory infusion center (infusion center supplies drug) ☐ Office administered (office supplies drug)						
☐ Ambulatory infusion center (pharmacy supplies drug) ☐ Dialysis Center administered (dialysis center supplies drug) ☐ Other:						
2. Yes No Is the patient currently on dialysis or will the patient be starting dialysis soon? If yes, please answer the following:						
☐ Yes ☐ No Is the Procrit to be used for a d	ialysis-related condition	on?				
3. Patient is stable on current drug(s) and/or current quantity, a	nd therapy change v	vould likely result in	n adverse clinical outcome			
4. All covered Part D drugs on any tier of the plan's formulary would not be as effective for the enrollee as the requested formulary drug and/or would likely have adverse effects for the enrollee.						
5. Yes No For all uses EXCEPT surgical procedures, was t	he patient's PRE-TR	EATMENT (no eryth	ropoietin treatment in			
previous month) hemoglobin (Hgb) level less than 10 g/dL (or less than 9 g/dL for anemia in CHF only)?						
Hgb level: g/dL; Da	nte:					
6. For elective, noncardiac, nonvascular surgery use:						
☐ Yes ☐ No Does the patient have a PRE-TREATMENT hemoglobin (Hgb) level greater than 10 g/dL but not more than						
13 g/dL?						
Hgb level: g/dL; Da						
7. For a diagnosis of primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis:						
☐ Yes ☐ No Does the patient have symptomatic anemia?						
☐ Yes ☐ No Is/was the PRE-TREATMENT serum erythropoietin level LESS than 500 mU/mL? EPO level: mU/mL; Date:						
EPO level: mU/mL;	Date:					

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Please check all bo	xes that apply <i>(continued)</i> :					
	Is the patient receiving chemotherapy with curative intent?					
☐ Yes ☐ No	Does the patient have myeloid cancer (such as acute myeloid leukemia [AML] [CML])?	or chronic myeloid leukemia				
9. For RENEWALS, please provide most current hemoglobin (Hgb) level: g/dL and complete this section.						
☐ Yes ☐ No	Has the patient had a recent blood transfusion?					
☐ Yes ☐ No	If no recent blood transfusion, has there been an increase in hemoglobin (Hgb) of at least 1 g/dL after at least 12 weeks of therapy?					
☐ Yes ☐ No	For anemia due to myelosuppressive anticancer chemotherapy, does the patient have a current hemoglobin (Hgb) of less than 11 g/dL?					
☐ Yes ☐ No						
10. Other supporting information:						
*NOTE: Requests that are subject to prior authorization (or any other utilization management requirement), may require supporting information. Please attach supporting information, as necessary, for your request.						
I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that documentation supporting this information is available for review if requested by the health plan sponsor, or, if applicable, a state or federal regulatory agency. I understand that any person who knowingly makes or causes to be made a false record or statement that is material to a claim ultimately paid by the United States government or any state government may be subject to civil penalties and treble damages under both the federal and state False Claims Acts. See, e.g., 31 U.S.C. §§ 3729-3733. By signing this form, I represent that I have obtained patient consent as required under applicable state and federal law, including but not limited to the Health Information Portability and Accountability Act (HIPAA) and state re-disclosure laws related to HIV/AIDS.						
Prescriber signatu	re	Date				

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