



Epogen, Procrit, Retacrit

Fax completed form to: (855) 840-1678
If this is an URGENT request, please call (800) 882-4462
(800.88.CIGNA)

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on this form are completed.*		
Specialty:	* DEA, NPI or TIN:				
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:	* Date of Birth:	
Office Fax:			* Patient Street Address:		
Office Street Address:			City:	State:	Zip:
City:	State:	Zip:	Patient Phone:		
Urgency: <input type="checkbox"/> Standard <input type="checkbox"/> Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)					
Medication requested: <input type="checkbox"/> Epogen <input type="checkbox"/> Procrit <input type="checkbox"/> Retacrit <input type="checkbox"/> Other (please specify): Strength: _____ Dosing schedule: _____ J-Code: _____ ICD10: _____ Number of Injections per month: _____ Expected duration: _____ Patient's weight: _____					
Where will this medication be obtained? <input type="checkbox"/> Accredo Specialty Pharmacy** <input type="checkbox"/> Home Health / Home Infusion vendor <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Physician's office stock (billing on a medical claim form) <input type="checkbox"/> Retail pharmacy <input type="checkbox"/> Other (please specify): **Cigna's nationally preferred specialty pharmacy					
**Medication orders can be placed with Accredo via E-prescribe - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822 NCPDP 4436920), Fax 888.302.1028, or Verbal 866.759.1557					
Facility and/or doctor dispensing and administering medication: Facility Name: _____ State: _____ Tax ID#: _____ Address (City, State, Zip Code): _____					
Where will this drug be administered? <input type="checkbox"/> Patient's Home <input type="checkbox"/> Physician's Office <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Other (please specify): _____					
NOTE: Per some Cigna plans, infusion of medication MUST occur in the least intensive, medically appropriate setting. Is this patient a candidate for re-direction to an alternate setting (such as alternate infusion site, physician's office, home) with assistance of a Specialty Care Options Case Manager? <input type="checkbox"/> Yes <input type="checkbox"/> No (provide medical necessity rationale):					
Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient? <input type="checkbox"/> Yes <input type="checkbox"/> No					
What is the patient's diagnosis? <input type="checkbox"/> Anemia in a Patient with Chronic Kidney Disease who is ON Dialysis <input type="checkbox"/> Anemia in a Patient with Chronic Kidney Disease who is NOT on Dialysis <input type="checkbox"/> Anemia in a Patient with Cancer due to Myelosuppressive Cancer Chemotherapy <input type="checkbox"/> Anemia Associated with Cancer in a Patient NOT Receiving Myelosuppressive Cancer Chemotherapy <input type="checkbox"/> Anemia Associated with Acute Myelogenous Leukemias (AML), Chronic Myelogenous Leukemias (CML), or other Myeloid Cancers <input type="checkbox"/> Anemia Associated with Radiotherapy in Cancer <input type="checkbox"/> Anemia Associated with Myelodysplastic Syndrome (MDS) <input type="checkbox"/> Anemia Associated with Myelofibrosis <input type="checkbox"/> Anemia in a Patient with Human Immunodeficiency Virus who is Receiving Zidovudine <input type="checkbox"/> Reduction of Allogeneic Red Blood Cell Transfusions in a Patient Undergoing Surgery <input type="checkbox"/> To Enhance Athletic Performance <input type="checkbox"/> Anemia due to Acute Blood Loss <input type="checkbox"/> Non-Anemic Patient (Hemoglobin greater than 13.0 g/dL) Prior to Surgery <input type="checkbox"/> Other					

(if other) Please provide the patient's diagnosis or reason for treatment.

Clinical Information:

(if CKD NOT on Dialysis) Is this initial therapy or is the patient currently receiving an Erythropoiesis-Stimulating Agent? Note: Examples of erythropoiesis-stimulating agents include an epoetin alfa product (for example, Epogen, Procrit, or Retacrit), a darbepoetin alfa product (for example, Aranesp), or a methoxy polyethylene glycol-epoetin beta product (for example, Mircera).

- Initial therapy
- Currently receiving an Erythropoiesis-Stimulating Agent
- None of the above

(if Myelosuppressive Chemo, MDS, Myelofibrosis, zidovudine HIV) Is this initial therapy or is the patient currently receiving an Erythropoiesis-Stimulating Agent? Note: Examples of erythropoiesis-stimulating agents include an epoetin alfa product (for example, Epogen, Procrit, or Retacrit) or a darbepoetin alfa product (for example, Aranesp).

- Initial therapy
- Currently receiving an Erythropoiesis-Stimulating Agent
- None of the above

(if CURRENTLY receiving CKD NOT on Dialysis, Myelosuppressive Chemo, MDS, Myelofibrosis, Zidovudine HIV) Which of the following best applies to your patient's hemoglobin?

- hemoglobin is 12 g/dL or less
- hemoglobin is 12.1 g/dL or higher

(if CKD NOT on Dialysis, Myelosuppressive Chemo, MDS, Myelofibrosis, Zidovudine HIV, Transfusions) Is the patient currently receiving iron therapy? Yes No

(if no) According to the prescriber, does the patient have adequate iron stores? Yes No

(if INITIAL, CKD NOT on Dialysis, 17 yr or younger) Which of the following best applies to your patient's hemoglobin?

- hemoglobin is 11 g/dL or less
- hemoglobin is 11.1 g/dL or higher

(if INITIAL, CKD NOT on Dialysis, 18 yr or older) Which of the following best applies to your patient's hemoglobin?

- hemoglobin is 9.9 g/dL or less
- hemoglobin is 10 g/dL or higher

(if Myelosuppressive Chemo) Is the patient currently receiving myelosuppressive chemotherapy? Yes No

(if yes) According to the prescriber, is the myelosuppressive chemotherapy considered non-curative? Yes No

(if INITIAL, Myelosuppressive Chemo) Which of the following best applies to your patient's hemoglobin?

- hemoglobin is 9.9 g/dL or less
- hemoglobin is 10 g/dL or higher

(if INITIAL, MDS/Myelofibrosis/Zidovudine HIV) Which of the following best applies to your patient's hemoglobin?

- hemoglobin is less than 10.0 g/dL
- hemoglobin is 10.1 g/dL or higher

(if not met) Which of the following best applies to your patient's serum erythropoietin level?

- serum erythropoietin level is 500 mU/ml or less
- serum erythropoietin level is 500.1 mU/ml or higher

(if MDS, Myelofibrosis) Is the requested medication being prescribed by (or in consultation with) a hematologist or oncologist? Yes No

if Myelofibrosis, currently receiving) According to the prescriber, has the patient responded to therapy which is defined as a hemoglobin of at least 10 g/dL? Yes No

(if no) Is your patient's current hemoglobin at least 2 g/dL higher than their pretreatment hemoglobin? Yes No

(if Zidovudine HIV) Is the patient currently receiving zidovudine therapy? Yes No

(if Transfusions) Which of the following best applies to your patient's hemoglobin?

- hemoglobin is 13 g/dL or less
- hemoglobin is 13.1 g/dL or higher

(if Transfusions) Is your patient scheduled for elective surgery? Yes No

(if yes) Is your patient scheduled for vascular or cardiac surgery? Yes No

(if Transfusions) Is your patient willing or able to donate autologous blood prior to surgery?

- Willing or able
 NOT willing or able

(if requesting Epogen) For Procrit [may require prior authorization], which of the following applies to your patient?

- Patient has not tried Procrit
 Patient tried Procrit, but it didn't work or didn't work well enough
 Patient tried Procrit, but had a significant allergy or serious adverse reaction
 Other

(if allergy or adverse reaction) Was this reaction due to a formulation difference in the inactive ingredients between Epogen and Procrit (for example, differences in stabilizing agent, buffering agent, and/or surfactant)? Yes No

(if yes) Please provide details to support.

Additional Pertinent Information:

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature: _____ **Date:** _____

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