



## MEDICATION POLICY

**Generic Name:** Epoetin Alfa

**Therapeutic Class or Brand Name:** Erythropoietins

**Applicable Drugs** (if Therapeutic Class):

Preferred: Procrit®

Non-Preferred: Epogen®

**Date of Origin:** 2/1/13

**Date Last Reviewed/Revised:** 12/1/17

**GPI Code:** 8240102000

### **Prior Authorization Criteria (may be considered medically necessary when criteria I through V are met):**

- I. Documented diagnosis of one of the following conditions A through D AND must meet criteria listed under applicable diagnosis:
  - A. Anemia due to Chronic Kidney Disease in patients on dialysis and patients not on dialysis, and criterion 1 is met:
    1. Minimum age requirement: 1 month old.
  - B. Anemia due to Zidovudine in HIV-infected patients and criterion 1 is met:
    1. Minimum age requirement: 8 months old.
  - C. Anemia due to the effects of concomitant myelosuppressive chemotherapy, and upon initiation, there is a minimum of two additional months of planned chemotherapy, and criterion 1 is met:
    1. Minimum age requirement: 5 years old.
  - D. Reduction of allogeneic RBC transfusions in patients undergoing elective, nonvascular, noncardiac surgery and criterion 1 is met (approve one time only):
    1. Minimum age requirement: 18 years old.
- II. Prescribing authority limited to hematologist, oncologist, nephrologist, gastroenterologist, and infectious disease specialist or based upon a consult with one of these specialists.
- III. Documentation showing that the patient does not have any GI bleeding.
- IV. Documentation that current hemoglobin is less than 10 g/dL.

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- V. Non-preferred products require a documented failure, intolerance, or contraindication to the preferred product(s).

### Exclusion Criteria:

- Patients with cancer receiving hormonal agents, biologic products, or radiotherapy, unless also receiving concomitant myelosuppressive chemotherapy.
- Patients with cancer receiving myelosuppressive chemotherapy when the anticipated outcome is cure.
- Patients with cancer receiving myelosuppressive chemotherapy when the anemia can be managed by transfusion.
- Patients scheduled for surgery who are willing to donate autologous blood.
- Patients undergoing cardiac or vascular surgery.
- As a substitute for RBC transfusions in patients who require immediate correction of anemia.
- Patients with uncontrolled hypertension.
- Patients with Pure Red Cell Aplasia (PRCA) that begins after treatment with erythropoietin protein drugs.

### Other Criteria:

- N/A

### Quantity/Days Supply Restrictions:

- The quantity is limited to a maximum of a 30 day supply per fill.

### Approval Length:

- **Authorization:** 6 months (unless otherwise stated under Prior Authorization Criteria section).
- **Re-Authorization:** An updated letter of medical necessity or progress notes showing no GI bleeding and hemoglobin less than 11 g/dL.

### Appendix:

N/A

### References:

1. <http://www.fda.gov/Drugs/DrugSafety/ucm259639.htm>.

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2. Medi-Span.
3. <http://www.procrit.com/sites/all/themes/procrit/resources/ProcritBooklet.pdf>
4. [http://pi.amgen.com/united\\_states/epogen/epogen\\_pi\\_hcp\\_english.pdf](http://pi.amgen.com/united_states/epogen/epogen_pi_hcp_english.pdf)

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<b>Historical Tracking Of Changes Made To Policy</b>	
12/1/2017	<ol style="list-style-type: none"> <li>1. <b>Added</b> “Patients with cancer receiving myelosuppressive chemotherapy when the anemia can be managed by transfusion” <b>under Exclusion Criteria.</b></li> </ol>
9/22/2016	<ol style="list-style-type: none"> <li>1. <b>Changed</b> “III. No GI bleeding” to “III. Documentation showing that the patient does not have any GI bleeding” <b>under Prior Authorization Criteria.</b></li> <li>2. <b>Changed</b> “IV. Hemoglobin less than 10 g/dL” to “IV. Documentation that current hemoglobin is less than 10 g/dL” <b>under Prior Authorization Criteria.</b></li> <li>3. <b>Changed</b> “V. Non-preferred Epogen® requires a documented failure, intolerance, or contraindication to the preferred product Procrit®” to “V. Non-preferred products require a documented failure, intolerance, or contraindication to the preferred product(s)” <b>under Prior Authorization Criteria.</b></li> <li>4. <b>Removed</b> “<a href="https://medicaid.utah.gov/pharmacy/priorauthorization/pdf/Erythropoetins.pdf">https://medicaid.utah.gov/pharmacy/priorauthorization/pdf/Erythropoetins.pdf</a>” <b>from References</b> (link no longer valid).</li> </ol>
3/4/2015	<ol style="list-style-type: none"> <li>1. <b>Changed Applicable Drugs from</b> “Procrit® and Epogen®” to “Preferred: Procrit®; Non-Preferred: Epogen®”.</li> <li>2. <b>Deleted duplicate GPI</b> “8240102000”.</li> <li>3. <b>Added</b> “Non-preferred Epogen® requires a documented failure, intolerance, or contraindication to the preferred product Procrit®” <b>under Prior Authorization Criteria.</b></li> <li>4. <b>Changed</b> “N/A” to “The quantity is limited to a maximum of a 30 day supply per fill” <b>under Quantity/Days Supply Restrictions.</b></li> <li>5. <b>Updated</b> “<a href="http://www.health.utah.gov/medicaid/pharmacy/priorauthorization/pdf/Erythropoetins.pdf">http://www.health.utah.gov/medicaid/pharmacy/priorauthorization/pdf/Erythropoetins.pdf</a>” to “<a href="https://medicaid.utah.gov/pharmacy/priorauthorization/pdf/Erythropoetins.pdf">https://medicaid.utah.gov/pharmacy/priorauthorization/pdf/Erythropoetins.pdf</a>” <b>under References.</b></li> </ol>
11/11/2013	<ol style="list-style-type: none"> <li>1. <b>Adapted policy to new format.</b></li> <li>2. <b>Added GPI Codes.</b></li> <li>3. <b>Changed Re-Authorization from</b>  “No GI bleeding and Hemoglobin less than 11 g/dL”  <b>to</b>  “An updated letter of medical necessity or progress notes showing no GI bleeding and hemoglobin less than 11 g/dL”.</li> <li>4. <b>Updated references</b> to include Medi-Span.</li> </ol>

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