

**I. Requirements for Prior Authorization of Erythropoiesis Stimulating Agents****A. Prescriptions That Require Prior Authorization**

All prescriptions for Erythropoiesis Stimulating Agents must be prior authorized.

**B. Review of Documentation for Medical Necessity**

In evaluating a request for prior authorization of a prescription for an Erythropoiesis Stimulating Agent (ESA), the determination of whether the requested prescription is medically necessary will take into account whether the beneficiary:

1. Is prescribed the ESA for the treatment of a diagnosis that is indicated in the U.S. Food and Drug Administration (FDA)-approved package labeling OR a medically accepted indication; **AND**
2. Is prescribed the ESA by or in consultation with an appropriate specialist (e.g., gastroenterologist, hematologist/oncologist, infectious disease specialist, nephrologist, surgeon, etc.); **AND**
3. Does not have a contraindication to the prescribed ESA; **AND**
4. Is prescribed a dose and duration of therapy that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**
5. Has been evaluated and treated for other causes of anemia (e.g., iron deficiency, hemolysis, vitamin B12 deficiency, folate deficiency, etc.); **AND**
6. **One** of the following:
  - a. Has serum ferritin  $\geq 100$  mcg/L and serum transferrin saturation  $\geq 20\%$
  - b. Is receiving supplemental iron therapy;**AND**
7. For a diagnosis of anemia associated with chronic kidney disease, has pretreatment hemoglobin  $< 10$  g/dL; **AND**
8. For a diagnosis of anemia in cancer patients on chemotherapy, **both** of the following:
  - a. Has pretreatment hemoglobin  $< 10$  g/dL
  - b. Is currently receiving myelosuppressive chemotherapy and the anticipated outcome is not cure;**AND**
9. For a diagnosis of anemia due to zidovudine in beneficiaries with HIV infection, **all** of the following:
  - a. Has pretreatment hemoglobin  $< 10$  g/dL,
  - b. Has a serum erythropoietin level  $\leq 500$  mUnits/mL,
  - c. Is receiving a dose of zidovudine  $\leq 4200$  mg/week;**AND**
10. For a reduction of allogeneic blood transfusion in surgery patients, **both** of the following:
  - a. Has pretreatment hemoglobin  $> 10$  to  $\leq 13$  g/dL
  - b. Is undergoing elective, noncardiac, nonvascular surgery;**AND**

11. For a non-preferred ESA, has a history of therapeutic failure, contraindication, or intolerance of the preferred ESAs approved or medically accepted for the beneficiary's diagnosis.

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

FOR RENEWALS OF PRIOR AUTHORIZATION FOR ESAs: The determination of medical necessity of a request for renewal of a prior authorization for an ESA that was previously approved will take into account whether the beneficiary:

1. **One** of the following:
  - a. Experienced an increase in hemoglobin compared to baseline
  - b. Is prescribed an increased dose of the requested ESA consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature;**AND**
2. Does not have a contraindication to the prescribed ESA; **AND**
3. Is prescribed a dose and duration of therapy that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**
4. **One** of the following:
  - a. Has serum ferritin  $\geq 100$  mcg/L and serum transferrin saturation  $\geq 20\%$
  - b. Is receiving supplemental iron therapy;**AND**
5. For a diagnosis of anemia associated with chronic renal disease, has **one** of the following:
  - a. Hemoglobin  $\leq 10$  g/dL for beneficiaries not on dialysis
  - b. Hemoglobin  $\leq 11$  g/dL for beneficiaries on dialysis,**AND**
6. For a diagnosis of anemia in cancer patients on chemotherapy, has hemoglobin  $\leq 12$  g/dL; **AND**
7. For a diagnosis of anemia in zidovudine-treated HIV-infected patients, **all** of the following:
  - a. Has hemoglobin  $\leq 12$  g/dL,
  - b. Has a serum erythropoietin level  $\leq 500$  mUnits/mL,
  - c. Is receiving a dose of zidovudine  $\leq 4200$  mg/week;**AND**
8. For a non-preferred ESA, has a history of therapeutic failure, contraindication, or intolerance of the preferred ESAs approved or medically accepted for the beneficiary's diagnosis.

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

### C. Clinical Review Process

Prior authorization personnel will review the request for prior authorization and apply the clinical guidelines in Section B. above to assess the medical necessity of a prescription for a Erythropoiesis Stimulating Protein. If the guidelines in Section B. are met, the reviewer will prior authorize the prescription. If the guidelines are not met, the prior authorization request will be referred to a physician reviewer for a medical necessity determination. Such a request for prior authorization will be approved when, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary.

**ERYTHROPOIESIS STIMULATING AGENTS PRIOR AUTHORIZATION FORM**

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		Total # of pgs: _____		Prescriber name:	
Name of office contact:				Specialty:	
Contact's phone number:				NPI:	State license #:
LTC facility contact/phone:				Street address:	
Beneficiary name:				Suite #:	City/State/Zip:
Beneficiary ID#:		DOB:	Phone:		Fax:

**CLINICAL INFORMATION**

Drug requested:		Strength & vial size:		<input type="checkbox"/> single-dose vial <input type="checkbox"/> multi-dose vial
Dose/directions:			Quantity:	Duration:
Diagnosis ( <i>submit documentation</i> ):			Dx code ( <i>required</i> ):	
<b>For non-preferred medication:</b> Does the beneficiary have a history of trial and failure of or contraindication or an intolerance to the preferred agents in this class that are approved or medically accepted for the beneficiary's diagnosis? Refer to <a href="https://papdl.com/preferred-drug-list">https://papdl.com/preferred-drug-list</a> for a list of preferred and non-preferred drugs in this class.			<input type="checkbox"/> Yes <i>Submit documentation.</i> <input type="checkbox"/> No	

**INITIAL requests**

Complete the section(s) below applicable to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

- ☐ Has transferrin or iron saturation  $\geq 20\%$  and ferritin  $\geq 100$  ng/mL
- ☐ Is receiving supplemental iron therapy
- ☐ Has adequately controlled blood pressure
- ☐ Was evaluated and treated for other causes of anemia (e.g., iron deficiency, hemolysis, vitamin B12 deficiency, folate deficiency, etc.)
- ☐ **For treatment of anemia associated with CHRONIC KIDNEY DISEASE:**
  - ☐ Has pretreatment hemoglobin  $< 10$  g/dL
- ☐ **For treatment of anemia in beneficiaries with CANCER RECEIVING CHEMOTHERAPY:**
  - ☐ Is currently receiving myelosuppressive chemotherapy
  - ☐ Is receiving chemotherapy with a non-curative intent
  - ☐ At initiation of therapy with an ESA, has an additional 2 or more months of planned chemotherapy
  - ☐ Has pretreatment hemoglobin  $< 10$  g/dL
- ☐ **For treatment of anemia in beneficiaries with HIV INFECTION RECEIVING ZIDOVUDINE:**
  - ☐ Has a serum erythropoietin level  $\leq 500$  mU/mL    ☐ Is taking zidovudine at a dose of  $\leq 4200$  mg/week
  - ☐ Has pretreatment hemoglobin  $< 10$  g/dL
- ☐ **For reduction of ALLOGENEIC BLOOD TRANSFUSIONS in beneficiaries undergoing SURGERY:**
  - ☐ Will be undergoing elective, non-cardiac, non-vascular surgery    ☐ Is not willing to donate autologous blood pre-operatively
  - ☐ Is at high risk for perioperative blood loss    ☐ Has pretreatment hemoglobin  $> 10$  g/dL and  $\leq 13$  g/dL

**RENEWAL requests**

Complete the section(s) below applicable to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

- ☐ Experienced an increase in hemoglobin compared to baseline
- ☐ Is prescribed an increased dose of the requested ESA
- ☐ Has transferrin or iron saturation  $\geq 20\%$  and ferritin  $\geq 100$  ng/mL
- ☐ Is receiving supplemental iron therapy
- ☐ Has adequately controlled blood pressure
- ☐ **For treatment of anemia associated with CHRONIC KIDNEY DISEASE:**
  - ☐ Is receiving dialysis and has a hemoglobin  $\leq 11$  g/dL
  - ☐ Is not receiving dialysis and has a hemoglobin  $\leq 10$  g/dL
- ☐ **For treatment of anemia in beneficiaries with CANCER RECEIVING CHEMOTHERAPY:**
  - ☐ Has a hemoglobin  $\leq 12$  g/dL
- ☐ **For treatment of anemia in beneficiaries with HIV INFECTION RECEIVING ZIDOVUDINE:**
  - ☐ Has a serum erythropoietin level  $\leq 500$  mU/mL    ☐ Is taking zidovudine at a dose of  $\leq 4200$  mg/week
  - ☐ Has a hemoglobin  $\leq 12$  g/dL

**PLEASE FAX COMPLETED FORM TO GATEWAY – PHARMACY DIVISION**

Prescriber Signature:	Date:
-----------------------	-------

**Confidentiality Notice:** The documents accompanying this telecopy may contain confidential information belonging to the sender. The information is intended only for the use of the individual named above. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or taking of any telecopy is strictly prohibited.