

**ERYTHROPOIESIS STIMULATING AGENTS PRIOR AUTHORIZATION FORM** (form effective 1/3/2022)

Prior authorization guidelines for Erythropoiesis Stimulating Agents are available on the DHS Pharmacy Services website at:

<https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>

<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 (submit documentation):	Dx code (required):		
<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 <input type="checkbox"/> No <i>Submit documentation.</i>		

**INITIAL requests**

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

- Is prescribed the ESA by or in consultation with a specialist (submit documentation of consultation if applicable)
- 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 at high risk for perioperative blood loss
  - Is not willing to donate autologous blood pre-operatively
  - Has pretreatment hemoglobin >10 g/dL and ≤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 ≥20% and ferritin ≥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 ≤11 g/dL
  - Is not receiving dialysis and has a hemoglobin ≤10 g/dL
- For treatment of anemia in beneficiaries with **CANCER RECEIVING CHEMOTHERAPY**:
  - Has a hemoglobin ≤12 g/dL
- For treatment of anemia in beneficiaries with **HIV INFECTION RECEIVING ZIDOVUDINE**:
  - Has a serum erythropoietin level ≤500 mU/mL
  - Is taking zidovudine at a dose of ≤4200 mg/week
  - Has a hemoglobin ≤12 g/dL

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION**

Prescriber Signature:

Date:

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