

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

Prior authorization guidelines for **Erythropoiesis Stimulating Agents** are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<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  $\leq 13$  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 submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.**

**Prescriber Signature:**

**Date:**

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