

ERYTHROPOIESIS STIMULATING AGENTS (ESAs)

Prior Authorization Form

ARANESP, EPOGEN, OMONTYS, & PROCRIT

<input type="checkbox"/> Standard Request <input type="checkbox"/> Expedited Request	If you or your prescriber believe that waiting for a standard decision could seriously harm your life, health, or ability to regain maximum function, you can request an expedited decision. For state exchanges only: The above disclaimer applies for exigent circumstances. Expedited review may also be requested when you are undergoing a current course of treatment using a non-formulary drug.
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Demographics

Patient Information		Prescriber Information	
Patient Name:		Prescriber Name:	
DOB:	Age:	NPI#:	Specialty:
Health Plan ID#:		Phone:	Fax:
Pharmacy Name:	Pharmacy Phone:	Office Contact:	Direct Phone # or Ext:

Medication Information

Drug Requested:	Strength:	Directions:
Quantity Dispensed:	Day Supply:	<input type="checkbox"/> Generic <input type="checkbox"/> Brand Necessary

Generic equivalent drugs will be substituted for Brand name drugs unless you specifically indicate otherwise.

<input type="checkbox"/> New medication <input type="checkbox"/> Continuation of therapy	Start Date:	If this is continuation of therapy, please provide CHART DOCUMENTATION indicating the member showed improvement while on therapy.
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Billing Information

<input type="checkbox"/> Billed by PHARMACY dispensed to the member or provider for administration.	<input type="checkbox"/> Billed under MEDICAL benefit by provider. J CODE: _____ ICD-10 Code: _____	Place of Administration: <input type="checkbox"/> Physician's Office <input type="checkbox"/> Hospital/Clinic <input type="checkbox"/> Patient Home
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Clinical Information

Diagnosis:	Date Diagnosed:
<input type="checkbox"/> Anemia due to renal failure	If Yes, please complete the following: <input type="checkbox"/> Acute <input type="checkbox"/> Chronic
<input type="checkbox"/> Anemia due to End Stage Renal Disease (ESRD)	Indicate severity of renal failure: _____
<input type="checkbox"/> Non-myeloid Malignancy	Is member on renal dialysis: <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Anemia due to chemotherapy	Anemia due to iron or folate deficiency? <input type="checkbox"/> Yes <input type="checkbox"/> No Due to Hemolysis? <input type="checkbox"/> Yes <input type="checkbox"/> No Due to GI bleed? <input type="checkbox"/> Yes <input type="checkbox"/> No

Member Name:	DOB:	Health Plan ID:
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Please be sure to complete and include this page with the 1st page of this form.

<input type="checkbox"/> HIV positive	Is the patient receiving Zidovudine? <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Anemia due to other cause	If Yes, please submit chart documentation indicating rationale for therapy and supportive lab values.
Does member have uncontrolled Hypertension? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Is patient scheduled to undergo elective, noncardiac, or nonvascular surgery and at high risk for perioperative transfusions? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Specify Hgb level (g/dL): _____	Date of test: _____
Is patient currently receiving iron supplement therapy?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	Specify serum ferritin (mcg/L): <input type="checkbox"/> <100 <input type="checkbox"/> 100-200 <input type="checkbox"/> >200 Specify serum transferrin saturation: <input type="checkbox"/> <20% <input type="checkbox"/> >20%
Does patient have a known hypersensitivity to the product's active substance or excipients?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	If Yes, please specify reaction and rationale for decision to use product:
What is the medication's starting dose?	What is the medication's maintenance dose:

Authorization Reassessment

Reassessment Period	Change in Hgb (g/dL)	Hgb (g/dL) and Test Date	Change in Hct	Pharmacist Adherence Evaluation
1 Month				
6 Month				