

Prior Authorization Request Form for Erythropoiesis Stimulating Agents

FAX this completed form to (844) 205-3386

OR Mail requests to: PA Department | 5 River Park Place East, Suite 210 | Fresno, CA 93720

OR Prior authorization may be completed at https://www.covermymeds.com/main/prior-authorization-forms/

I. PROVIDER INFORMATION		II. MEMBER INFORMATION				
Prescriber Name:		Member Name:				
Prescriber Specialty:		Identifica	Identification #:			
Office Contact Name:		Group #:				
Group Name:		Date of Birth:				
Fax #:		Medication Allergies:				
Phone #:						
III. DRUG INFORMATION (One drug	g request per for	m)				
Drug name and strength: Dosage Interval (sig):		ig):		Qty. per Day:		
IV. REQUIRED DOCUMENTION (Detailed medical record documentation demonstrating evidence for each item must be submitted with prior authorization request)						
Specify diagnosis & diagnosis code releva	ant to this request:		Dx/Dx Code:			
Does the member have a history of a contraindication to the requested medication?			☐ Yes			
			□ No			
Requests for all non-preferred Erythropoiesis Stimulatic Agents: Does the member have a history of trial and failure contraindication or intolerance to the preferred Erythropoie Stimulating Agents? Refer to https://papdl.com/preferred-dr a list of preferred and non-preferred medications in this class.		of or esis r <u>ug-list</u> for	☐ Yes ☐ No	Medication Taken Previously (start and end date and dose):		
☐ If requesting for daily quantity exceeding daily limit (Refer to https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Quantity-Limits-and-Daily-Dose-Limits.aspx), please provide supporting information:						
specialist, nephrologist, surgeon, Has been evaluated and treated for folate deficiency, etc) One of the following: Has serum ferritin ≥ 100mcg Is receiving supplemental irous associated for a diagnosis of anemia associated Has pretreatment hemoglobited Is currently receiving myelosed Has pretreatment hemoglobited Has a serum erythropoietin less receiving a dose of zidovuction.	g specialist, (e.g., heretc) please indicate or other causes of a large of the cause of	matologist a speciali nemia (e.g ferrin satu dney disea otherapy, herapy an bers with l nL k	st consulted: st consulted: st, iron deficient aration ≥ 20% ase, has pretre both of the fo	etment hemoglobin <10g/dL ollowing: ated outcome is not cure all of the following:		
 Is receiving a dose of zidovudine ≤4200mg/week For a reduction of allogeneic blood transfusion in surgery patients, both of the following: Has pretreatment hemoglobin >10g/dL to ≤13g/dL 						

☐ Is undergoing elective, noncardiac, nonvascular surgery						
RENEWAL REQUIEST:						
One of the following:						
Experienced an increase in hemoglobin compared to baseline						
☐ Is prescribed an increased dose of the requested Erythropoiesis Stimulating Agents (I	SA) consistent with					
FDA-approved package labeling, nationally recognized compendia, or peer-reviewed	nedical literature					
One of the following:						
Has serum ferritin ≥ 100mcg/L and serum transferrin saturation ≥ 20%						
☐ Is receiving supplemental iron therapy						
 For a diagnosis of anemia associated with chronic renal disease, has one of the following: Hemoglobin ≤ 10g/dL for members not on dialysis 						
☐ Hemoglobin ≤ 10g/dL for members not on dialysis ☐ Hemoglobin ≤ 11g/dL for members on dialysis						
☐ For a diagnosis of anemia in cancer patients on chemotherapy, has hemoglobin ≤ 12g/dL						
For a diagnosis of anemia due to zidovudine in members with HIV infection, all of the following	ıg:					
☐ Has pretreatment hemoglobin <12g/dL	· ·					
☐ Has a serum erythropoietin level ≤500mUnits/mL						
☐ Is receiving a dose of zidovudine ≤4200mg/week						
IV. ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL INFORMATION :						
Appropriate clinical information to support the request on Provider Signature:	Date:					
the basis of medical necessity must be submitted.						

PA Department will respond via fax or phone within 24 hours.
Requests for prior authorization (PA) requests must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)