AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-305-671-0200</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

Erythropoiesis Stimulating Agents (ESAs) *For Non-Dialysis Use*

This form is to be completed ONLY if the patient is self-administering

<u>Drug Requested</u>: (check one below)

	,		
□ Aranesp® (darbepoetin alfa	Epogen® (epoetin alfa)	☐ Mircera® (methoxy polyethylene glycol-epoetin beta)	
□ Procrit ® (epoetin alfa)	□ Retacrit [™] (epoetin alfa epbx)	1-	
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MEMBER & PRESCRIBI	ER INFORMATION: Auth	norization may be delayed if incomplete.	
Member Name:			
Member AvMed #: Date of Birth:			
Prescriber Name: Date:			
_			
Office Contact Name:			
Phone Number:	Fax Number:		
DEA OR NPI #:			
DRUG INFORMATION: Authorization may be delayed if incomplete.			
Drug Name/Form/Strength:			
osing Schedule: Length of Therapy:			
	ignosis: ICD Code, if applicable:		
Weight: Date:			
CLINICAL CRITERIA: 0	heck below all that apply. All ca	riteria must be met for approval. To	

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

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□ D	iagnosis: Anemia Due to Chronic Kidney Disease
<u>Initi</u>	al Authorization: 6 months
	Member has a documented diagnosis of anemia due to chronic kidney disease (CKD)
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days
	☐ Member must meet <u>ONE</u> of the following hemoglobin requirements:
	☐ Member is an adult with a hemoglobin level <10 g/dL
	☐ Member is a pediatric patient who is symptomatic with a hemoglobin level <11 g/dL
	☐ Member's serum ferritin $\geq 100 \text{ ng/mL (mcg/L)}$
	□ Member's transferrin saturation (TSAT) ≥ 20%
	Member is <u>NOT</u> receiving hemodialysis
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)
□ D	iagnosis: Anemia Due to Chronic Kidney Disease
To su	athorization: 6 months. Check below all that apply. All criteria must be checked for approval. pport each line checked, all documentation, including (lab results, diagnostics, and/or chart notes) be provided or request may be denied.
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days
	☐ Member's hemoglobin level $\leq 12 \text{ g/dL}$
	☐ Member's serum ferritin $\ge 100 \text{ ng/mL (mcg/L)}$
	□ Member's transferrin saturation (TSAT) \geq 20%
□ D	iagnosis: Anemia Due to Myelosuppressive Chemotherapy
Leng	<u>th of Authorization</u> : 6 months
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days
	☐ Member must meet <u>ONE</u> of the following hemoglobin requirements:
	☐ Member is an adult with a hemoglobin level <10 g/dL
	☐ Member is a pediatric patient who is symptomatic with a hemoglobin level <11 g/dL
	☐ Member's serum ferritin $\geq 100 \text{ ng/mL (mcg/L)}$
	☐ Member's transferrin saturation (TSAT) \geq 20%
	Member is being treated with myelosuppressive chemotherapy and provider has noted member's current treatment regimen:
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)

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	□ Diagnosis: Anemia Due to Myelodysplastic Syndrome (MDS)		
Initia	<u>ll Authorization</u> : 6 months		
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days		
	\square Member's hemoglobin level $< 10 \text{ g/dL}$		
	☐ Member's serum ferritin $\ge 100 \text{ ng/mL (mcg/L)}$		
	■ Member's transferrin saturation (TSAT) ≥ 20%		
	☐ Member's serum erythropoietin level ≤ 500 milliunits/mL		
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)		
□ Di	agnosis: Anemia Due to Myelodysplastic Syndrome (MDS)		
To sup	thorization: 12 months. Check below all that apply. All criteria must be checked for approval. port each line checked, all documentation, including (lab results, diagnostics, and/or chart notes) must wided or request may be denied.		
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days		
	☐ Member's hemoglobin level <12 g/dL		
	☐ Member's serum ferritin $\geq 100 \text{ ng/mL (mcg/L)}$		
	☐ Member's transferrin saturation $(TSAT) \ge 20\%$		
	☐ Member's serum erythropoietin level ≤ 500 milliunits/mL		
	agnosis: Anemia of Prematurity		
Leng	th of Authorization: 6 months		
	Documentation of <u>ALL</u> the following must be submitted:		
	☐ Medication will be used in combination with iron supplementation		
	☐ Member must meet <u>ONE</u> of the following:		
	☐ Member's birth weight <1500 grams		
	☐ Member's gestational age <33 weeks		
□ Di	agnosis: Anemia Due to Myelosuppressive Medication Regimen for HIV		
Initia	d Authorization: 6 months		
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days		
	☐ Member must meet ONE of the following hemoglobin requirements:		
	☐ Member is an adult with a hemoglobin level <10 g/dL		
	☐ Member is a pediatric patient who is symptomatic with a hemoglobin level <11 g/dL		
	☐ Member's serum ferritin $\geq 100 \text{ ng/mL (mcg/L)}$		

	□ Member's transferrin saturation (TSAT) $\geq 20\%$
	☐ Member's serum erythropoietin level ≤ 500 milliunits/mL
	Member is being treated with an HIV medication regimen that includes zidovudine (≤ 4200mg/week)
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)
□ Di	iagnosis: Anemia Due to Myelosuppressive Medication Regimen for HIV
To sup	thorization: 6 months. Check below all that apply. All criteria must be checked for approval. pport each line checked, all documentation, including (lab results, diagnostics, and/or chart notes) be provided or request may be denied.
	Member continues to receive an HIV medication regimen that includes zidovudine (≤ 4200 mg/week)
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days: \Box Member's hemoglobin level \leq 12 g/dL
	☐ Member's serum ferritin ≥ 100 ng/mL (mcg/L)
	\square Member's transferrin saturation (TSAT) $\geq 20\%$
	☐ Member's serum erythropoietin level ≤ 500 milliunits/mL
□ Di	iagnosis: Anemia Due to Myelosuppressive Medication Regimen for Hepatitis C
<u>Initia</u>	al Authorization: 6 months
	Member has a documented diagnosis of anemia
	Member is being treated with a myelosuppressive regimen (e.g., ribavirin with interferon or peginterferon) for the treatment of Hepatitis C
	Provider must submit documentation of \underline{ALL} the following test results obtained within the last 30 days:
	☐ Member must meet <u>ONE</u> of the following hemoglobin requirements:
	☐ Member is an adult with a hemoglobin level <10 g/dL
	☐ Member is a pediatric patient who is symptomatic with a hemoglobin level <11 g/dL ☐ Member's sorrym familin > 100 ng/mL (mag/L)
	 □ Member's serum ferritin ≥ 100 ng/mL (mcg/L) □ Member's transferrin saturation (TSAT) ≥ 20%
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)
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□ Di	iagnosis: Anemia Due to Myelosuppressive Medication Regimen for Hepatitis C
To sup	thorization: 6 months. Check below all that apply. All criteria must be checked for approval. pport each line checked, all documentation, including (lab results, diagnostics, and/or chart notes)
	be provided or request may be denied.

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	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days
	☐ Member's hemoglobin level ≤12 g/dL
	□ Member's serum ferritin $\ge 100 \text{ ng/mL (mcg/L)}$
	□ Member's transferrin saturation (TSAT) \geq 20%
	iagnosis: Reduction of Allogenic Red Blood Cell Transfusions in Patients ndergoing Elective, Noncardiac, Nonvascular Surgery
Leng	th of Authorization: 3 months
	Requested drug will be used to decrease the need for blood transfusion in a surgery patient
	Member is scheduled to undergo surgery within the next three (3) months
	Provider must submit documentation of ALL the following test results obtained within the last 30 days
	☐ Member's hemoglobin level <13 g/dL
	☐ Member's serum ferritin $\ge 100 \text{ ng/mL (mcg/L)}$
	□ Member's transferrin saturation (TSAT) $\geq 20\%$
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)
□ D	iagnosis: All Other Indications
Leng	th of Authorization: 6 months
	Member's diagnosis of anemia and/or risk factors for development of anemia must be noted in submitted chart notes for medical necessity approval
	Provider must document requested length of therapy:
	Provider must submit documentation of <u>ALL</u> the following test results obtained within the last 30 days Member's current hemoglobin level:
	□ Member's serum ferritin $\ge 100 \text{ ng/mL (mcg/L)}$
	□ Member's transferrin saturation (TSAT) ≥ 20%
	☐ If applicable, any other test results to support medical necessity approval
	All other causes of anemia have been ruled out (e.g., iron, vitamin B12 or folate deficiency, hemolysis)
Med	ication being provided by Specialty Pharmacy – Proprium Rx

Use of samples to initiate therapy does not meet step edit/preauthorization criteria.

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.