

## PRIOR AUTHORIZATION REQUEST FORM **ARANESP®**

For authorization, please answer each question and fax this form PLUS chart notes back to Real Rx at 385-425-4052.

Fai	ilure to submit clinical documentation to support this request will re	esult in	a dism	nissal of the request.			
lf y	ou have prior authorization questions, please call for assistance 385-	-425-50	94.				
Dis	claimer: Prior authorization request forms are subject to change in accorda	nce wit	h Feder	al and State notice requirements.			
Dat	te: Member Name:		ID#:				
DO	B: Gender:	Physic		ian:			
Off	fice Phone: Office Fax:		Office	fice Contact:			
Height/Weight:			HCPCS Code:				
preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.  Product being requested:  Aranesp® (darbepoetin alfa)  Dosing/Frequency:							
If the request is for reauthorization, proceed to reauthorization section							
	Questions	Yes	No	Comments/Notes			
1.	Is the requested medication being purchased by the provider's						
	office and to be billed under the medical benefit ('buy-and-bill')?						
2.	Is this request for an <b>expedited</b> review?						
	By checking the "Yes" box to request an expedited review (24						
	hours), you are certifying that applying the standard review time						
	frame (72 hours) may place the member's life, health, or ability to						
<u> </u>	regain maximum function in serious jeopardy.						
პ.	Is the requesting provider a hematologist, oncologist, nephrologist, or in consultation with one?		Ц				
4.	Does documentation show that the member's hemoglobin is <10		П	Please provide documentation			
	g/dL and/or that the hematocrit is <30%?		_	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			
5.	Does the member have one of the following indications:			Please provide documentation			
	Anemia of chronic renal failure,			•			
	Anemia due to myelosuppressive chemotherapy with a						
	minimum of 8 additional weeks of planned chemotherapy,						
	<ul> <li>Myelodysplasia or myelodysplastic syndrome?</li> </ul>						
6.	Does the member have one of the following indications:						
	<ul> <li>Request will be used as a substitute for red blood cell</li> </ul>						
	transfusion in patients who require immediate correction of						
	anemia,						
	<ul> <li>Uncontrolled hypertension,</li> </ul>						
	<ul> <li>Pure Red Cell Aplasia (PRCA) that begins after treatment with</li> </ul>						
	erythropoietin drugs?						

REAUTHORIZATION							
1.	Is the request for reauthorization of therapy?						
2.	Has the member responded to treatment, demonstrated by an			Please provide documentation			
	improvement in the hematocrit and hemoglobin levels or a						
	significant decrease in transfusion requirements?						
3.	Is current hemoglobin < 11g/dL OR > 10 to <12 g/dL?			Please provide documentation			
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.							
Additional information:							
Physician's Signature:							

\*\* Failure to submit clinical documentation to support this request will result in a dismissal of the request.\*\*

Policy: PHARM-008

Origination Date: 05/24/2018 Reviewed/Revised Date: 01/17/2024 Next Review Date: 01/17/2025 Current Effective Date: 02/01/2024

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