



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.

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

If you have prior authorization questions, please call for assistance 385-425-5094.

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Form with fields: Date, Member Name, ID#, DOB, Gender, Physician, Office Phone, Office Fax, Office Contact, Height/Weight, HCPCS Code.

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with 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

Table with 4 columns: Questions, Yes, No, Comments/Notes. Contains 6 rows of questions regarding medication purchase, expedited review, provider type, hemoglobin levels, and medical indications.

REAUTHORIZATION			
1. Is the request for reauthorization of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Has the member responded to treatment, demonstrated by an improvement in the hematocrit and hemoglobin levels or a significant decrease in transfusion requirements?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Is current hemoglobin < 11g/dL OR > 10 to <12 g/dL?	<input type="checkbox"/>	<input type="checkbox"/>	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:			

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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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