

HEALTHY U MEDICAID

PRIOR AUTHORIZATION REQUEST FORM

TAVNEOS®

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department at 888-509-8142.

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 Pharmacy Customer Service for assistance at 855-856-5694

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

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:

Height/Weight:

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: Tavneos® (avacopan)

Dosing/Frequency: _____

If the request is for reauthorization, proceed to reauthorization section.

Questions	Yes	No	Comments/Notes
SEVERE ANTINEUTROPHIL CYTOPLASMIC AUTOANTIBODY (ANCA)-ASSOCIATED VASCULITIS (AAV)			
1. Is the member ≥ 18 years of age?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
2. Is the request made by, or in consultation with, a rheumatologist?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Does the member have a diagnosis of either of the following: <ul style="list-style-type: none"> • Active antineutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis (AAV) due to granulomatosis with polyangiitis (GPA); or • Microscopic polyangiitis (MPA) 	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Does documentation show positive anti-proteinase 3 (PR3) OR anti-myeloperoxidase (MPO) ANCA-associated vasculitis?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
5. Does the member have a current eGFR ≥ 15 mL/min/1.73m ² ?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
6. Does the member currently require dialysis or have a kidney transplant, and has received plasma exchange in the past 12 weeks?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
7. Does the member have a baseline Birmingham Vasculitis Activity Score (BVAS; version 3) with either of the following: <ul style="list-style-type: none"> • At least one or more major items • At least three or more non-major items • At least two renal items of hematuria and proteinuria 	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

8. Has the member had at least a 3-month trial and failure of glucocorticoid therapy at the maximally indicated doses, unless contraindicated or a clinically significant intolerance is experienced?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
9. Does documentation show concurrent therapy with cyclophosphamide or rituximab?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
10. Does documentation show baseline Hepatitis B (HBV) prior to initiating therapy?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
REAUTHORIZATION			
1. Is the requesting for reauthorization of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Has the member shown a $\geq 50\%$ reduction of BVAS score from baseline?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
3. Does documentation show continued liver function monitoring performed by the provider?	<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 Signature:			

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Policy: PHARM-HU-132
 Origination Date: 02/02/2022
 Reviewed/Revised Date: 03/15/2023
 Next Review Date: 03/15/2024
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