

Program Number	2026 P 2265-5
Program	Prior Authorization/Medical Necessity
Medication	Tavneos® (avacopan)
P&T Approval Date	1/2022, 1/2023, 1/2024, 1/2025, 1/2026
Effective Date	4/1/2026

## 1. Background:

Tavneos (avacopan) is a complement 5a receptor (C5aR) antagonist indicated as an adjunctive treatment of adult patients with severe active anti-neutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis (granulomatosis with polyangiitis [GPA] and microscopic polyangiitis [MPA]) in combination with standard therapy including glucocorticoids. Tavneos does not eliminate glucocorticoid use.

## 2. Coverage Criteria <sup>a</sup>:

<p><b>A. Anti-Neutrophil Cytoplasmic Autoantibody (ANCA) - Associated Vasculitis</b></p> <p><b>1. <u>Initial Authorization</u></b></p> <p>a. <b>Tavneos</b> will be approved based on <b>all</b> of the following criteria:</p> <p>(1) Diagnosis of severe active ANCA-associated vasculitis</p> <p style="text-align: center;"><b>-AND-</b></p> <p>(2) Submission of medical records (e.g., chart notes, laboratory values, etc.) documenting the disease is <b>one</b> of the following types:</p> <p>(a) Granulomatosis with polyangiitis (GPA)</p> <p>(b) Microscopic polyangiitis (MPA)</p> <p style="text-align: center;"><b>-AND-</b></p> <p>(3) Patient is being treated with an initial immunosuppressive regimen to induce remission (i.e., rituximab, cyclophosphamide)</p> <p style="text-align: center;"><b>-AND-</b></p> <p>(4) Tavneos is being prescribed as adjunctive treatment in combination with standard therapy (e.g. prednisone, azathioprine, mycophenolate, methotrexate, rituximab, cyclophosphamide)</p> <p style="text-align: center;"><b>-AND-</b></p> <p>(5) Prescribed by or in consultation with <b>one</b> of the following:</p>
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- (a) Rheumatologist
- (b) Nephrologist
- (c) Pulmonologist
- (d) Vascular Medicine Specialist

**Authorization will be issued for 6 months.**

## 2. **Reauthorization**

a. **Tavneos** will be approved based on **all** of the following criteria:

- (1) Patient does not show evidence of progressive disease while on Tavneos therapy

**-AND-**

- (2) Tavneos is being prescribed as adjunctive treatment in combination with standard therapy (e.g. prednisone, azathioprine, mycophenolate, methotrexate, rituximab, cyclophosphamide)

**-AND-**

- (3) Prescribed by or in consultation with **one** of the following:

- (a) Rheumatologist
- (b) Nephrologist
- (c) Pulmonologist
- (d) Vascular Medicine Specialist

**Authorization will be issued for 12 months.**

<sup>a</sup> State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

## 3. **Additional Clinical Rules:**

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.
- Supply limits may be in place.



**4. References:**

1. Tavneos [package insert]. Cincinnati, OH: Thermo Fisher Scientific; June 2024.

Program	Prior Authorization/Medical Necessity - Tavneos® (avacopan)
<b>Change Control</b>	
1/2022	New program
1/2023	Annual review with no change to coverage criteria.
1/2024	Annual review with no changes.
1/2025	Annual review with no changes to coverage criteria. Updated reference.
1/2026	Annual review. Added “or in consultation with” for prescriber requirement in the initial criteria.