SPECIALTY GUIDELINE MANAGEMENT

TAVNEOS (avacopan)

POLICY

I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met, and the member has no exclusions to the prescribed therapy.

FDA-Approved Indication

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.

All other indications are considered experimental/investigational and not covered benefits.

II. DOCUMENTATION

Submission of the following information is necessary to initiate the prior authorization review:

- A. Initial requests:
 - 1. Chart notes or medical records showing positive serum assay for Proteinase-3 (PR3) or myeloperoxidase (MPO) antibody
 - 2. Chart notes or medical records of pre-treatment Birmingham Vasculitis Activity Score (BVAS) version 3
 - 3. Chart notes or medical records of at least one major item, or at least three non-major items, or at least two renal items of hematuria and proteinuria present on BVAS version 3 (refer to Appendix)
- B. Continuation requests:
 - 1. Chart notes or medical records showing stabilization or improvement in the BVAS version 3

III. CRITERIA FOR INITIAL APPROVAL

Anti-neutrophil cytoplasmic autoantibody (ANCA)-associated vasculitis (granulomatosis with polyangiitis [GPA] and microscopic polyangiitis [MPA])

Authorization of 6 months may be granted for treatment of severe active ANCA-associated vasculitis (GPA and MPA) when all of the following are met:

- A. Tavneos will be used in combination with standard therapy (e.g., cyclophosphamide, azathioprine, mycophenolate, rituximab)
- B. The member is positive for anti-PR3 or anti-MPO antibody

C. Pre-treatment Birmingham Vasculitis Activity Score (BVAS) version 3 demonstrates the presence of at least one major item, or at least three non-major items, or at least two renal items of hematuria and proteinuria (refer to Appendix)

IV. CONTINUATION OF THERAPY

Authorization of 6 months may be granted for continued treatment for severe active ANCA-associated vasculitis (GPA and MPA) in members who achieve or maintain positive clinical response as evidenced by stabilization or improvement in the BVAS and the member has not received more than 12 months of therapy with Tavneos.

V. QUANTITY LIMIT

• Tavneos 10mg capsules: 6 capsules a day

VI. APPENDIX

Birmingham Vasculitis Activity Score (version 3)

*Major items are indicated in bold italics

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General	Cardiovascular	
Myalgia	 Loss of pulses 	
Arthralgia/ arthritis	Valvular heart disease	
• Fever ≥ 38 °C	 Pericarditis 	
 Weight loss ≥ 2 kg 	Ischemic cardiac pain	
	 Cardiomyopathy 	
	Congestive cardiac failure	
Cutaneous	Abdominal	
Infarct	 Peritonitis 	
Purpura	Bloody diarrhea	
• Ulcer	Ischemic abdominal pain	
• Gangrene		
Other skin vasculitis		
Mucous membranes/ eyes	Renal	
Mouth ulcers	Hypertension	
Genital ulcers	• Proteinuria $> 1 + \text{ or } > 0.2 \text{ g/g}$ creatinine	
Adnexal inflammation	• Hematuria ≥ 10 RBCs/hpf	
Significant proptosis	• Serum creatinine 125-249 μmol/L (1.41-	
Scleritis/ Episcleritis	2.82 mg/dL)	
Conjunctivitis/ Blepharitis/ Keratitis	• Serum creatinine 250-499 μmol/L (2.83-	
Blurred vision	5.64 mg/dL)	
Sudden vision loss	• Serum creatinine ≥ 500 μmol/L (5.65	
• Uveitis	mg/dL)	
Retinal changes (vasculitis/	• Rise in serum creatinine > 30% or fall in	
thrombosis/ exudate/ hemorrhage	creatinine clearance > 25%	

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	Reviewed: 01/2022, 3/2023, 2/2024	
	Scope: Medicaid	
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Ear Nose & Throat	Nervous System
Bloody nasal discharge/ crusts/ ulcers/	Headache
granulomata	Meningitis
 Paranasal sinus involvement 	Seizures (not hypertensive)
Subglottic stenosis	Cerebrovascular accident
 Conductive hearing loss 	Organic confusion
• Sensorineural hearing loss	Spinal cord lesion
	• Cranial nerve palsy
	Sensory peripheral neuropathy
	Mononeuritis multiplex
Chest	Other
• Wheeze	• RBC casts and/or glomerulonephritis
Nodules or cavities	
Pleural effusion/ pleurisy	
Infiltrate	
Endobronchial involvement	
Massive hemoptysis/ alveolar	
hemorrhage	
Respiratory failure	

VII. REFERENCES

- 1. Tavneos [package insert]. Cincinnati, OH: ChemoCentryx, Inc.; February 2022.
- 2. American College of Rheumatology. 2021 American college of rheumatology/vasculitis foundation guideline for the management of antineutrophil cytoplasmic antibody-associated vasculitis. *Arthritis & Rheumatology*. Https://www.vasculitisfoundation.org/wp-content/uploads/2021/07/2021-ACR-VF-Guideline-for-Management-of-ANCA-Associated-Vasculitis.pdf. Accessed October 21, 2021.
- 3. Geetha D, Jefferson JA. ANCA-Associated vasculitis: Core curriculum 2020. *Am J Kidney Dis.* 75(1): 124-137.
- 4. Jayne DRW, Merkel PA, Schall TJ et al. Avacopan for the treatment of ANCA-associated vasculitis [supplemental appendix]. N Engl J Med. 2021; 384:599-609. DOI: 10.1056/NEJMoa2023386. https://www.nejm.org/doi/suppl/10.1056/NEJMoa2023386/suppl_file/nejmoa2023386_appendix. pdf. Accessed October 28, 2021.