

UnitedHealthcare Pharmacy Clinical Pharmacy Programs

Program Number	2025 P 1474-2
Program	Prior Authorization/Notification
Medication	Alhemo® (concizumab-mtci)
P&T Approval Date	3/2025, 8/2025
Effective Date	10/1/2025

1. Background:

Alhemo (concizumab-mtci) is a tissue factor pathway inhibitor (TFPI) antagonist indicated for routine prophylaxis to prevent or reduce the frequency of bleeding episodes in adult and pediatric patients 12 years of age and older with:

- hemophilia A (congenital factor VIII deficiency) with or without FVIII inhibitors
- hemophilia B (congenital factor IX deficiency) with or without FIX inhibitors

2. Coverage Criteria^a:

A. Hemophilia A

1. Initial Authorization

- a. Alhemo will be approved based on all of the following criteria:
 - (1) Diagnosis of hemophilia A

-AND-

(2) Patient is 12 years of age or older

-AND-

(3) Prescribed for the prevention of bleeding episodes (i.e., routine prophylaxis)

Authorization of therapy will be issued for 12 months.

2. Reauthorization

- a. Alhemo will be approved based on the following criterion:
 - (1) Documentation of positive clinical response to Alhemo therapy

Authorization will be issued for 12 months.

B. Hemophilia B

1. Initial Authorization

a. Alhemo will be approved based on <u>all</u> of the following criteria:



(1) Diagnosis of hemophilia B

-AND-

(2) Patient is 12 years of age or older

-AND-

(3) Prescribed for the prevention of bleeding episodes (i.e., routine prophylaxis)

Authorization of therapy will be issued for 12 months.

2. Reauthorization

- a. **Alhemo** will be approved based on the following criterion:
 - (1) Documentation of positive clinical response to Alhemo therapy

Authorization will be issued for 12 months.

^a 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 reauthorization 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.
- Medical Necessity and supply limits may be in place.

4. References:

1. Alhemo® [package insert]. Plainsboro, NJ: Novo Nordisk Inc., July 2025.

Program	Prior Authorization/Notification - Hemlibra (emicizumab-kxwh)
Change Control	
3/2025	New program.
8/2025	Added coverage criteria for hemophilia A or B without inhibitors per updated FDA label.