

Tretten (coagulation Factor XIII A-Subunit [recombinant]) Effective 01/01/2024

Plan	☐ MassHealth UPPL ☐ Commercial/Exchange	Dungung Time	⊠ Prior Authorization
Benefit	☑ Pharmacy Benefit☑ Medical Benefit	Program Type	☐ Quantity Limit ☐ Step Therapy
Specialty	This medication has been designated specialty and must be filled at a contracted		
Limitations	specialty pharmacy.		
Contact Information	Medical and Specialty Medications		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Tretten is indicated for routine prophylaxis of bleeding in patients with congenital factor XIII A-subunit deficiency. Tretten is not for use in patients with congenital factor XIII B-subunit deficiency. All other indications are considered experimental/investigational and not medically necessary.

Coverage Guidelines

Authorization may be granted for members new to the plan who are currently receiving treatment with the requested medication, excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted for prophylaxis of bleeding in members with congenital factor XIII A-subunit deficiency.

Continuation of Therapy

Authorization may be granted for continued treatment in members experiencing benefit from therapy (e.g., reduced frequency or severity of bleeds).

Limitations

Approvals will be granted for 12 months.

References

- 1. Tretten [package insert]. Plainsboro, NJ: Novo Nordisk Inc.; June 2020.
- National Hemophilia Foundation. MASAC recommendations concerning products licensed for the treatment of hemophilia and other bleeding disorders. Revised March 2022. MASAC Document #272. https://www.hemophilia.org/sites/default/files/document/files/272_Treatment.pdf. Accessed October 4, 2022.

Review History

12/13/2023: Reviewed at Dec P&T, switched from SGM to Custom. Effective 1/1/2024

