

JURISDICTION SPECIFIC MEDICARE PART B

HUMATE-P (antihemophilic factor/von Willebrand factor complex [human])

POLICY

I. COVERED USES

The indications below are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

- A. Treatment and prevention of bleeding in adults with hemophilia A
- B. Treatment of spontaneous and trauma-induced bleeding episodes due to von Willebrand disease (VWD)
- C. Prevention of excessive bleeding during and after surgery in patients with VWD. This applies to patients with severe VWD as well as patients with mild to moderate VWD where use of desmopressin (DDAVP) is known or suspected to be inadequate.

All other indications will be assessed on an individual basis. Submissions for indications other than those enumerated in this policy should be accompanied by supporting evidence from Medicare approved compendia.

II. DOCUMENTATION

The following documentation must be available in a legible format with patient identification information (e.g., complete name and dates of service) and signature of physician or non-physician practitioner responsible for and providing care to the member, upon request, for all submissions:

- A. The submitted medical record must support the use of the selected ICD-10-CM codes. The submitted CPT/HCPCS code must describe the service performed.
- B. The medical record documentation must support the medical necessity of the services as stated in this policy.

III. CRITERIA FOR APPROVAL

A. Hemophilia A (Hereditary Factor VIII Deficiency)

Authorization of 12 months may be granted for treatment of hemophilia A when the requested drug will be used for any of the following:

1. Primary prophylactic therapy when the member has less than 1 percent of normal factor (less than 0.1 IU/mL) with the aim of keeping the factor VIII level above 1 percent between doses.
2. Continuous prophylactic therapy when the member does not have severe disease (greater than 1 percent of normal factor levels) and the member experiences repeated episodes of spontaneous bleeding.
3. The requested drug will be used as on-demand treatment to control bleeding episodes.
4. The requested drug will be used for immune tolerance therapy.

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B. Von Willebrand Disease

Authorization of 12 months may be granted for von Willebrand disease.

IV. REFERENCES

1. Hemophilia Factor Products LCD (L35111) Version R16. Available at: <https://www.cms.gov/medicare-coverage-database/indexes/national-and-local-indexes.aspx>. Accessed December 15, 2021.
2. Billing and Coding: Hemophilia Factor Products (A56433) Version R7. Available at: <https://www.cms.gov/medicare-coverage-database/indexes/national-and-local-indexes.aspx>. Accessed December 15, 2021.
3. Humate-P [package insert]. Kankakee, IL: CSL Behring, LLC; June 2020.