Clinical Policy: Factor VIII/von Willebrand Factor Complex (Human - Alphanate, Humate-P, Wilate)  
Reference Number: CP.PHAR.216  
Effective Date: 05.01.16  
Last Review Date: 02.20  
Line of Business: Commercial, HIM, Medicaid

See Important Reminder at the end of this policy for important regulatory and legal information.

Description
The following are factor VIII/von Willebrand factor complexes (human) requiring prior authorization: Alphanate®, Humate®-P, and Wilate®.

FDA Approved Indication(s)
Factor VIII/von Willebrand factor complexes are indicated for:

- Hemophilia A
  - Alphanate: Control and prevention of bleeding episodes and perioperative management in adult and pediatric patients with factor VIII deficiency due to hemophilia A
  - Humate-P: Treatment and prevention of bleeding in adults with hemophilia A (classical hemophilia)
  - Wilate:
    - Control and prevention of bleeding episodes
    - Routine prophylaxis to reduce the frequency of bleeding episodes

- Von Willebrand disease (VWD) in children and adults:
  - Alphanate: Surgical and/or invasive procedures in patients in whom desmopressin (DDAVP) is either ineffective or contraindicated
  - Humate-P:
    - Treatment of spontaneous and trauma-induced bleeding episodes
    - Prevention of excessive bleeding during and after surgery. This applies to patients with severe VWD as well as patients with mild to moderate VWD where use of DDAVP is known or suspected to be inadequate
  - Wilate:
    - On-demand treatment and control of bleeding episodes
    - Perioperative management of bleeding

Limitation(s) of use:
- Alphanate is not indicated for patients with severe VWD (type 3) undergoing major surgery.

Policy/Criteria
Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of health plans affiliated with Centene Corporation® that Alphanate, Humate-P, and Wilate are medically necessary when the following criteria are met:
I. Initial Approval Criteria
   A. Congenital Hemophilia A (must meet all):
      1. Diagnosis of congenital hemophilia A (factor VIII deficiency);
      2. Prescribed by or in consultation with a hematologist;
      3. Request is for one of the following uses (a, b, or c):
         a. Control or prevention of bleeding episodes;
         b. Perioperative management (Alphanate only);
         c. Routine prophylaxis to reduce the frequency of bleeding episodes (Wilate only);
      4. If factor VIII coagulant activity levels are > 5%, failure of desmopressin acetate, unless contraindicated, clinically significant adverse effects are experienced, or an appropriate formulation of desmopressin acetate is unavailable;
      5. Dose does not exceed the FDA approved maximum recommended dose for the relevant indication.
   
   Approval duration: 3 months

   B. Von Willebrand Disease (must meet all):
      1. Diagnosis of VWD (types 1, 2, or 3);
      2. Prescribed by or in consultation with a hematologist;
      3. Request is for one of the following uses (a or b):
         a. Treatment of bleeding episodes (Humate-P and Wilate only);
         b. Perioperative management;
      4. Dose does not exceed the FDA approved maximum recommended dose for the relevant indication.
   
   Approval duration: 3 months

   C. Other diagnoses/indications
      1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid.

II. Continued Therapy
   A. All Indications in Section I (must meet all):
      1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
      2. Member is responding positively to therapy;
      3. If request is for a dose increase, new dose does not exceed the FDA approved maximum recommended dose for the relevant indication.
   
   Approval duration: 3 months

   B. Other diagnoses/indications (must meet 1 or 2):
      1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.
   
   Approval duration: Duration of request or 3 months (whichever is less); or
2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid.

III. Diagnoses/Indications for which coverage is NOT authorized:

A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policy – CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid, or evidence of coverage documents.

IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key

DDAVP: desmopressin acetate
FDA: Food and Drug Administration
VWD: von Willebrand disease

Appendix B: Therapeutic Alternatives

This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent and may require prior authorization.

<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Dosing Regimen</th>
<th>Dose Limit/ Maximum Dose</th>
</tr>
</thead>
</table>
| desmopressin acetate (Stimate® nasal spray; generic injection solution) | When Factor VIII coagulant activity levels are > 5%  
Injection: 0.3 mcg/kg IV every 48 hours  
Nasal spray:  
< 50 kg: 1 spray intranasally in one nostril only; may repeat based on laboratory response and clinical condition  
≥ 50 kg: 1 spray intranasally in each nostril; may repeat based on laboratory response and clinical condition | Injection: 0.3 mcg/kg IV every 48 hours  
Nasal spray: 1 spray intranasally in each nostril |

Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.

Appendix C: Contraindications/Boxed Warnings

- Contraindication(s): patients with known hypersensitivity reactions, including anaphylactic or severe systemic reaction, to human plasma-derived products, any ingredient in the formulation, or components of the container
- Boxed warning(s): none reported
## V. Dosage and Administration

<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
</tr>
</thead>
</table>
| Factor VIII/von Willebrand factor complex (Alphanate) | Hemophilia A - control and prevention of bleeding episodes | Minor episodes: 15 IU/kg IV every 12 hours  
Moderate episodes: 25 IU/kg IV every 12 hours  
Major episodes: 40-50 IU/kg IV initially followed by 25 IU/kg IV every 12 hours | 100 IU/kg/day |
| Factor VIII/von Willebrand factor complex (Humate-P) | Hemophilia A - control and prevention of bleeding episodes | Minor episodes: 15 IU/kg IV loading dose followed by half of the loading dose given once or twice daily if needed  
Moderate episodes: 25 IU/kg IV loading dose followed by 15 IU/kg IV every 8-12 hours  
Major episodes: 40-50 IU/kg IV initially followed by 20-25 IU/kg IV every 8 hours | 75 IU/kg/day |
| Factor VIII/von Willebrand factor complex (Alphanate) | Hemophilia A – perioperative management | Pre-operative: 40-50 IU/kg IV once as a single dose  
Post-operative: 30-50 IU/kg IV every 12 hours | 100 IU/kg/day |
| Factor VIII/von Willebrand factor complex (Humate-P) | VWD – control and prevention of bleeding episodes | Type 1 VWD, mild disease  
Minor or major episodes: 40-60 IU/kg IV loading | 240 IU/kg/day |
<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Factor VIII/von Willebrand factor complex (Wilate)</td>
<td>Hemophilia A - control and prevention of bleeding episodes</td>
<td>Minor or moderate episodes: 30-40 IU/kg IV every 12-24 hours</td>
<td>150 IU/kg/day</td>
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<tr>
<td></td>
<td></td>
<td>Major episodes: 35-50 IU/kg IV every 12-24 hours</td>
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<td></td>
<td>Life-threatening episodes: 35-50 IU/kg IV every 8-24 hours</td>
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<tr>
<td>Factor VIII/von Willebrand factor complex (Wilate)</td>
<td>Hemophilia A – routine prophylaxis</td>
<td>20-40 IU/kg IV every 2 to 3 days</td>
<td>40 IU/kg/day</td>
</tr>
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</table>
### CLINICAL POLICY

#### Factor VIII/von Willebrand Factor Complex

<table>
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<tr>
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<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
</tr>
</thead>
</table>
| Factor VIII/von Willebrand factor complex (Wilate) | VWD – control and prevention of bleeding episodes | Minor episodes: 20-40 IU/kg IV loading dose followed by 20-30 IU/kg every 12-24 hours  
Major episodes: 40-60 IU/kg IV loading dose followed by 20-40 IU/kg every 12-24 hours | 60 IU/kg/day |
| Factor VIII/von Willebrand factor complex (Wilate) | VWD – perioperative management                  | Minor surgeries (including tooth extraction): 30-60 IU/kg IV loading dose followed by 15-30 IU/kg every 12-24 hours  
Major surgeries: 40-60 IU/kg IV loading dose followed by 20-40 IU/kg every 12-24 hours | 60 IU/kg/day |

### VI. Product Availability

<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Availability</th>
</tr>
</thead>
<tbody>
<tr>
<td>Factor VIII/von Willebrand factor complex (Alphanate)</td>
<td>Vial: 250, 500, 1,000, 1,500 IU and 2,000 IU FVIII</td>
</tr>
<tr>
<td>Factor VIII/von Willebrand factor complex (Humate-P)</td>
<td>Vial: 250/600, 500/1,200, 1,000/2,400 IU FVIII/VWF:RCo</td>
</tr>
<tr>
<td>Factor VIII/von Willebrand factor complex (Wilate)</td>
<td>Vial: 500/500, 1,000/1,000 IU FVIII/VWF:RCo</td>
</tr>
</tbody>
</table>

### VII. References

Coding Implications

Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

<table>
<thead>
<tr>
<th>HCPCS Codes</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>J7183</td>
<td>Injection, von Willebrand factor complex (human), Wilate, 1 IU vWF:RCo</td>
</tr>
<tr>
<td>J7186</td>
<td>Injection, antihemophilic factor VIII/von Willebrand factor complex (human), per factor VIII i.u. (Alphanate)</td>
</tr>
<tr>
<td>J7187</td>
<td>Injection, von Willebrand factor complex (Humate-P), per IU VWF:RCo</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Reviews, Revisions, and Approvals</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Policy split from CP.PHAR.12.Blood Factors and converted to new template. Removed requests for documentation. Removed indication for prophylaxis after 2 joint bleeds/approval period 6 months as there is no FDA approved indication for long-term prophylaxis. Approval period is edited to be 3 months initial and one 3-month re-auth as, in some circumstances, treatment could be necessary for up to six months (e.g., intracranial hemorrhage per Alphanate PI). Reviewed by specialist.</td>
<td>04.01.16</td>
<td>05.16</td>
</tr>
<tr>
<td>Removed “major surgery” restriction for Alphanate. Required trial of desmopressin is edited to avoid necessity of testing for coagulation factors. Safety information removed. Uses and approval periods across all blood factor policies worded consistently. Efficacy statement added to renewal criteria. Hemophilies are specified as “congenital” versus “acquired” across blood factor policies where indicated. Reviewed by specialist-hematology/ internal medicine</td>
<td>04.01.17</td>
<td>05.17</td>
</tr>
<tr>
<td>1Q18 annual review: - Converted to new template -No significant changes - References reviewed and updated.</td>
<td>11.27.17</td>
<td>02.18</td>
</tr>
<tr>
<td>1Q 2019 annual review: added HIM-Medical Benefit; no significant changes; references reviewed and updated.</td>
<td>09.26.18</td>
<td>02.19</td>
</tr>
<tr>
<td>1Q 2020 annual review: no significant changes; added HIM line of business; RT4 policy update addition of hemophilia A indication for</td>
<td>11.27.19</td>
<td>02.20</td>
</tr>
</tbody>
</table>
Reviews, Revisions, and Approvals | Date | P&T Approval Date
---|---|---
Wilate, mirroring previously approved hemophilia A coverage policies for other FVIII products; references reviewed and updated | | 03.13.20
Added Commercial line of business | | 03.13.20

**Important Reminder**
This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.
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Note: For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

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