

## **Clinical Policy: Velaglucerase Alfa (VPRIV)**

Reference Number: CP.PHAR.163

Effective Date: 02.01.16

Last Review Date: 05.19

Line of Business: Commercial, HIM-Medical Benefit,  
Medicaid

[Coding Implications](#)  
[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

### **Description**

Velaglucerase alfa (VPRIV<sup>®</sup>) is a hydrolytic lysosomal glucocerebroside-specific enzyme.

### **FDA Approved Indication(s)**

VPRIV is indicated for long-term enzyme replacement therapy for patients with type 1 Gaucher disease (GD1).

### **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<sup>®</sup> that VPRIV is **medically necessary** when the following criteria are met:

#### **I. Initial Approval Criteria**

##### **A. Gaucher Disease** (must meet all):

1. Diagnosis of Type 1 (GD1) or Type 3 Gaucher disease (GD3) confirmed by one of the following (a or b):
  - a. Enzyme assay demonstrating a deficiency of beta-glucocerebrosidase (glucosidase) activity;
  - b. DNA testing;
2. Age  $\geq$  4 years;
3. Member is symptomatic (e.g., anemia, thrombocytopenia, bone disease, hepatomegaly, splenomegaly);
4. VPRIV is not prescribed concurrently with Elelyso<sup>®</sup> (taliglucerase alfa) or Cerezyme<sup>®</sup> (imiglucerase).

##### **Approval duration:**

**Medicaid/HIM** – 6 months

**Commercial** – 6 months or to the member's renewal date, whichever is longer

##### **B. 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 and CP.PMN.53 for Medicaid and HIM-Medical Benefit.

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#### II. Continued Therapy

##### A. Gaucher Disease (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 as evidenced by increased or stabilized platelet count or hemoglobin, reduced or stabilized spleen or liver volume, decreased bone pain;
3. VPRIV is not prescribed concurrently with Elelyso<sup>®</sup> (taliglucerase alfa) or Cerezyme<sup>®</sup> (imiglucerase).

##### Approval duration:

**Medicaid/HIM** – 12 months

**Commercial** – 6 months or to the member's renewal date, whichever is longer

##### B. Other diagnoses/indications (must meet 1 or 2):

1. Currently receiving medication via health plan benefit and documentation supports positive response to therapy.

**Approval duration: Duration of request or 6 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 and CP.PMN.53 for Medicaid and HIM-Medical Benefit.

#### 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 and CP.PMN.53 for Medicaid and HIM-Medical Benefit or evidence of coverage documents.

#### IV. Appendices/General Information

##### *Appendix A: Abbreviation/Acronym Key*

FDA: Food and Drug Administration

GD1: type 1 Gaucher disease

GD3: type 3 Gaucher disease

##### *Appendix B: Therapeutic Alternatives*

Not applicable

##### *Appendix C: Contraindications/Boxed Warnings*

- Contraindication(s): none reported.
- Boxed warning(s): none reported.

##### *Appendix D: General Information*

- Measures of Therapeutic Response: GD1 is a heterogeneous disorder which involves the visceral organs, bone marrow, and bone in almost all affected patients. Common conditions resulting from GD1 include anemia, thrombocytopenia, hepatomegaly,

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splenomegaly, and bone disease. Therefore, hemoglobin level, platelet count, liver volume, spleen volume, and bone pain are clinical parameters that can indicate therapeutic response to GD1 therapies. In some clinical trials, stability has been defined as the following thresholds of change from baseline: hemoglobin level < 1.5 g/dL decrease, platelet count < 25% decrease, liver volume < 20% increase, and spleen volume < 25% increase.

- Enzyme replacement therapy such as Cerezyme may have beneficial palliative effects in Type 2 disease, but does not alter the outcome and is not generally used.
- According to the European consensus guidelines revised recommendations on the management of neuronopathic Gaucher disease by Vellodi et al: (1) there is clear evidence in most patients that enzyme replacement therapy (ERT) ameliorates systemic involvement in non-neuronopathic (Type 1) as well as chronic neuronopathic Gaucher disease (Type 3), enhancing quality of life; (2) There is no evidence that ERT has reversed, stabilized or slowed the progression of neurological involvement; (3) In patients with established acute neuronopathic Gaucher disease (Type 2), enzyme replacement therapy has had little effect on the progressively downhill course. It has merely resulted in prolongation of pain and suffering.
- There is currently insufficient clinical evidence that supports the combination use of enzyme replacement therapy with Zavesca® (miglustat).

**V. Dosage and Administration**

Indication	Dosing Regimen	Maximum Dose
Gaucher Disease	<p><u>Patients naïve to enzyme replacement therapy:</u> 60 units/kg IV every other week</p> <p><u>Patients being treated with stable imiglucerase dosages:</u> Switch to VPRIV at previous imiglucerase dose 2 weeks after last imiglucerase dose</p>	Individualized

**VI. Product Availability**

Single-use vial: 400 units

**VII. References**

1. VPRIV Prescribing Information. Lexington, MA: Shire Human Genetic Therapies, Inc.; April 2015. Available at <http://www.vpriv.com>. Accessed February 27, 2019.
2. Charrow J, Andersson HC, Kaplan P. Enzyme replacement therapy and monitoring for children with type 1 Gaucher disease: consensus recommendations. J Pediatr. 2004; 144: 112-20.
3. Hollak, CEM, Weinreb NJ. The attenuated/late onset lysosomal storage disorders: therapeutic goals and indications for enzyme replacement treatment in Gaucher and Fabry disease. Best Pract Res Clin Endocrinol Metab. 2015; 29: 205-218.
4. Pastores GM, Weinreb NJ, Aerts H, et al. Therapeutic goals in the treatment of Gaucher disease. Semin Hematol. 2004; 41(suppl 5): 4-14.

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5. Andersson HC, Charrow J, Kaplan P, et al. Individualization of long-term enzyme replacement therapy for Gaucher disease. *Genet Med.* 2005; 7(2): 105-110.
6. Altarescu G, Hill S, Wiggs E, et al. The efficacy of enzyme replacement therapy in patients with chronic neuronopathic Gaucher`s disease. *J Pediatr.* 2001;138:539-547.
7. Vellodi A, Tytki-Szymanska A, Davies E, et al. Management of neuronopathic Gaucher disease: Revised recommendations. *J Inherit Metab Dis.* 2009;32:660-664.

### 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.

HCPCS Codes	Description
J3385	Injection, velaglucerase alfa, 100 units

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy split from CP.PHAR.48. Policy converted to new template.	01.16	02.16
Age restriction removed; DNA testing added to diagnostic methods. Allergy history is removed as the drug can be continued in some cases. ERT monotherapy added; Positive response added; Background section converted to new template.	12.16	02.17
Added age restriction; Added requirement for presence of symptoms. Added examples of what can constitute a positive response to therapy. Added ERT monotherapy requirement for re-auth requests in addition to the initial criteria; Added appendix B.	08.24.17	11.17
2Q 2018 annual review: no significant changes from previously approved corporate policy; policies combined for Medicaid and Commercial lines of business; HIM added; Commercial: split from “CP.CPA.241 Gaucher Disease Therapy – Enzyme Replacement” into individual Elelyso commercial policy; added age limit; removed maximum dose limit, as dosing is individualized per patient response to therapy; added the requirement that VPRIV is not be used concurrently with other enzyme replacement therapies; added specific examples of positive response to therapy, for reauthorization. Medicaid: added coverage for Type 3 Gaucher disease, as covered under the Commercial policy; references reviewed and updated.	02.26.18	05.18
2Q 2019 annual review: no significant changes; references reviewed and updated.	02.27.19	05.19

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#### **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.

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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.

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and their representatives agree to be bound by such terms and conditions by providing services to members and/or submitting claims for payment for such services.

**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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