CENTENE® Corporation

# Clinical Policy: Betibeglogene Autotemcel

Reference Number: CP.PHAR.545
Effective Date: FDA Approval Date
Last Review Date: 08.22
Line of Business: Commercial, HIM, Medicaid

Coding Implications
Revision Log

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



## Description

Betibeglogene autotemcel is a genetically modified autologous CD34+ cell enriched population that contains hematopoietic stem cells transduced with a lentiviral vector encoding the $\beta^{A\text{-}T87Q}$-globin gene.

## FDA Approved Indication(s) [Pending]

Betibeglogene autotemcel is proposed for the treatment of transfusion-dependent β-thalassemia (TDT).

## 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 betibeglogene autotemcel is **medically necessary** when the following criteria are met:

**I. Initial Approval Criteria***

**Criteria will mirror the clinical information from the prescribing information once FDA-approved.*
**Only for initial treatment dose; subsequent doses will not be covered.*

**A. Transfusion-Dependent β-Thalassemia** (must meet all)**:**

1. Diagnosis of TDT with genetic confirmation (*see Appendix E*);*
2. Prescribed by or in consultation with a hematologist;
3. Member meets one of the following (a or b):*
   a. Age ≥ 5 years and ≤ 50 years;
   b. If age < 5 years, member meets both of the following (i and ii):
      i. Weight ≥ 6 kg;
      ii. Provider submits medical rationale that member is anticipated to be able to provide at least the minimum number of cells required to initiate the manufacturing process;
4. Documentation of one of the following in the two previous years (a or b):*
   a. Receipt of ≥ 8 transfusions of packed red blood cells (pRBC) per year (*see Appendix D*);
   b. Receipt of ≥ 100 mL/kg pRBC per year (*see Appendix D*);
5. Member is eligible for an allogeneic hematopoietic stem cell transplantation (HSCT);*
6. Member has not received prior allogeneic HSCT or gene therapy;*

7. Member does not have advanced liver disease (*see Appendix D*);*
8. Member is not positive for the presence of HIV type 1 or 2, hepatitis B virus, or hepatitis C virus;*
9. Member does not have any prior or current malignancy;*
10. Dose does not exceed 20 x $10^6$ CD34+ cells per kg.*

**Approval duration: 3 months (one time infusion per lifetime)**

**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, HIM.PA.154 for health insurance marketplace, and CP.PMN.53 for Medicaid.

## II. Continued Therapy*

**Criteria will mirror the clinical information from the prescribing information once FDA-approved.*

**A. Transfusion-Dependent β-Thalassemia**

1. Re-authorization is not permitted.

**Approval duration: Not applicable**

**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, HIM.PA.154 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 policies – CP.CPA.09 for commercial, HIM.PA.154 for health insurance marketplace, and CP.PMN.53 for Medicaid or evidence of coverage documents.

## IV. Appendices/General Information

*Appendix A: Abbreviation/Acronym Key*

FDA: Food and Drug Administration
HIV: human immunodeficiency virus
HLA: human leukocyte antigen
HSCT: hematopoietic stem cell transplantation
pRBC: packed red blood cells
TDT: transfusion-dependent β-thalassemia

*Appendix B: Therapeutic Alternatives*

Not applicable

*Appendix C: Contraindications/Boxed Warnings [Pending]*

- Contraindication(s): pending
- Boxed warning(s): pending

*Appendix D: General Information*

- Conversion of RBC units from mL: 1 RBC unit in this criteria refers to a quantity of pRBC approximately 200-350 mL.
  - Sites who use transfusion bags within this range, or ≥ 350 mL, the conversion in units should be done by dividing the volume transfused to the patient by 350 mL.
  - Sites who use transfusion bags < 200 mL, the conversion in units should be done by dividing the volume transfused to the patient by 200 mL.
- Examples of advanced liver disease include, but are not limited to, the following:
  - Cirrhosis
  - Active hepatitis
  - Bridging fibrosis
  - Fatty liver disease

*Appendix E: Genetic Confirmation of TDT*

| Beta Thalassemia Genotype Examples |
|---|
| $\beta^0/\beta^0$ |
| $\beta^0/\beta^+$ |
| $\beta^+$ IVS1-110/$\beta^+$ IVS1-110 |
| $B^E/\beta^0$ |

## V. Dosage and Administration [Pending]

| Indication | Dosing Regimen | Maximum Dose |
|---|---|---|
| TDT* | Pending | Pending |

## VI. Product Availability [Pending]

Pending

## VII. References

1. ClinicalTrials.gov. A study evaluating the efficacy and safety of the Lentiglobin® BB305 drug product in subjects with transfusion-dependent β-thalassemia, who do not have a β0/β0 genotype. Last updated June 25, 2021. Available at: https://clinicaltrials.gov/ct2/show/NCT02906202. Accessed June 26, 2021.
2. ClinicalTrials.gov. A study evaluating the efficacy and safety of the Lentiglobin® BB305 drug product in subjects with transfusion-dependent β-thalassemia. Last updated June 24, 2021. Available at: https://clinicaltrials.gov/ct2/show/NCT03207009. Accessed June 26, 2021.
3. Porter JB, Thompson AA, Walters MC, et al. Improvement in erythropoiesis in patients with transfusion dependent β-thalassemia following treatment with betibeglogene autotemcel (LentiGlobin for β-thalassemia) in the phase 3 HGB-207 study. EHA 2020 Virtual Congress Abstract: S296.
4. Cappellini MD, Farmakis D, Porter J, et al. Guidelines for the management of transfusion dependent thalassemia (TDT) 4th Edition. Thalassemia International Federation (2021). Available at: https://thalassaemia.org.cy/publications/tif-publications/guidelines-for-the-management-of-transfusion-dependent-thalassaemia-4th-edition-2021/. Accessed May 3, 2022.

**Coding Implications [Pending]**
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 |
|---|---|
| Pending | Pending |

| Reviews, Revisions, and Approvals | Date | P&T Approval Date |
|---|---|---|
| Policy created pre-emptively. | 07.13.21 | 08.21 |
| 3Q 2022 annual review: no significant changes as drug is not yet FDA-approved; references reviewed and updated. | 05.03.22 | 08.22 |

**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.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

This clinical policy is the property of the Health Plan. Unauthorized copying, use, and distribution of this clinical policy or any information contained herein are strictly prohibited. Providers, members and their representatives are bound to the terms and conditions expressed herein through the terms of their contracts. Where no such contract exists, providers, members 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.