

Clinical Policy: Lantidra (donislecel): Allogeneic Pancreatic Islet Cellular Therapy

Reference Number: CP.MP.250

Date of Last Revision: 07/26

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Description

This policy describes the medical necessity criteria for Lantidra (donislecel), an allogeneic pancreatic islet cellular therapy, used for the treatment of type 1 diabetes in those who are unable to reach target hemoglobin A1c (HbA1c).¹

***Note:** For criteria applicable to Medicare plans, please see MC.CP.MP.250 Lantidra (donislecel): Allogeneic pancreatic islet cellular therapy.*

Policy/Criteria

- I. It is the policy of health plans affiliated with Centene Corporation® that Lantidra (donislecel) is **medically necessary** when all of the following criteria are met:
 - A. Member/enrollee is ≥ 18 years of age;
 - B. Diagnosis of type 1 diabetes;
 - C. Member/enrollee is unable to approach target HbA1c due to current repeated episodes of severe hypoglycemia, despite intensive diabetes management and education;
 - D. Therapy will be used in conjunction with concomitant immunosuppression;
 - E. No more than three infusions will be given;
 - F. Member/enrollee does not have concomitant diseases or conditions, including pregnancy, that contraindicate the procedure for Lantidra infusion or immunosuppression.

Background

Type 1 diabetes is a chronic autoimmune disease, characterized by the destruction of pancreatic beta cells, causing insulin deficiency.^{2,3} Intensive diabetes management is the standard of care for type 1 diabetes, and includes coordinating frequent blood glucose monitoring and insulin replacement with meals and activity.³ Type 1 diabetes requires lifelong insulin therapy. However, some individuals experience significant challenges in achieving adequate glycemic control without increasing the risk of hypoglycemia, especially in those with hypoglycemia unawareness.² In these cases, insulin management can be complex, and Lantidra may be considered as a treatment option for appropriately selected patients.²

In June 2023, the U.S. Food and Drug Administration (FDA) approved Lantidra, an allogeneic pancreatic islet cellular therapy derived from deceased donor pancreatic islet cells, including insulin-producing beta cells.^{1,2} Lantidra is administered as a single infusion into the hepatic portal vein, where the transplanted islet cells engraft and produce enough insulin to regulate blood glucose levels, which may reduce or eliminate the need for additional insulin administration.² Lantidra is indicated for adult patients with type 1 diabetes who are unable to achieve target glycemic control due to recurrent severe hypoglycemic episodes, despite intensive diabetes management and education.^{1,2} To prevent rejection and maintain the viability of the transplanted islet cells, recipients must receive immunosuppressive therapy before receiving Lantidra and must continue long-term immunosuppression after the infusion.¹

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Two non-randomized, single-arm studies evaluated the safety and effectiveness of Lantidra and included a total of 30 enrollees with type 1 diabetes and hypoglycemic unawareness who received a minimum of one infusion and a maximum of three infusions. The studies found that 21 participants did not require insulin administration for one year or longer, 11 participants did not require insulin administration for one to five years, and 10 participants did not require insulin administration for over five years. Additionally, five of the total participants continued to require insulin administration without any resulting days of insulin independence.²

Coding Implications

This clinical policy references Current Procedural Terminology (CPT[®]). CPT[®] is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2025, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. 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.

HCP Codes	Description
J3590	Unclassified biologics

Reviews, Revisions, and Approvals	Review Date	Approval Date
Policy developed.	08/23	08/23
Added note to description regarding Medicare policy version. Removed maximum age requirement from Criteria I.A.	02/24	02/24
Annual review. References reviewed and updated. Reviewed by external specialist.	07/24	07/24
Annual review. References reviewed and updated.	07/25	07/25
Annual review. Background updated with no impact to criteria. Coding and description reviewed. References reviewed and updated. Reviewed by external specialist.	07/26	07/26

References

1. Lantidra [package insert]. Chicago, IL: CellTrans Inc.; 2023. Accessed June 08, 2026.
2. U.S. Food and Drug Administration. FDA Approves First Cellular Therapy to Treat Patients with Type 1 Diabetes. <https://www.fda.gov/news-events/press-announcements/fda-approves-first-cellular-therapy-treat-patients-type-1-diabetes>. Published June 28, 2023. Accessed June 04, 2026.
3. Hirsch IB. Glycemic management in adults with type 1 diabetes mellitus. UpToDate. www.uptodate.com. Published April 28, 2026. Accessed June 04, 2026.
4. Islet Transplantation in Type I Diabetic Patients Using the University of Illinois at Chicago (UIC) Protocol. National Institutes of Health. <https://clinicaltrials.gov/study/NCT03791567?intr=donislecel&rank=1>. Published March 16, 2022 (updated October 22, 2024). Accessed June 04, 2026.
5. National Institute for Health and Care Excellence (NICE). Type 1 diabetes in adults: diagnosis and management [NG17]. Published August 26, 2015. Updated August 17, 2022. Accessed June 08, 2026.

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6. Rickels MR, Ballou CM, Foster NC, et al. Islet Transplantation Versus Standard of Care for Type 1 Diabetes Complicated by Severe Hypoglycemia From the Collaborative Islet Transplant Registry and the T1D Exchange Registry. *Diabetes Care*. 2025;48(5):737-744. doi:10.2337/dc24-1915

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/enrollees. This clinical policy is not intended to recommend treatment for members/enrollees. Members/enrollees 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.

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providing services to members/enrollees and/or submitting claims for payment for such services.

Note: For Medicaid members/enrollees, 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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Note: For Medicare members/enrollees, to ensure consistency with the Medicare National Coverage Determinations (NCD) and Local Coverage Determinations (LCD), all applicable NCDs, LCDs, and Medicare Coverage Articles should be reviewed prior to applying the criteria set forth in this clinical policy. Refer to the CMS website at <http://www.cms.gov> for additional information.

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