

POLICY: Oncology (Injectable) – Nipent Utilization Management Medical Policy

- Nipent™ (pentostatin intravenous infusion – Hospira)

EFFECTIVE DATE: 2/1/2022

REVIEW DATE: 09/10/2025

COVERAGE CRITERIA FOR: All UCare Plans

OVERVIEW

Nipent, an adenosine deaminase inhibitor, is indicated for the treatment of untreated and alpha-interferon refractory **hairy cell leukemia** in patients with active disease, defined by clinically significant anemia, neutropenia, thrombocytopenia, or disease-related symptoms, as a single-agent.¹

Guidelines

Nipent is addressed in National Comprehensive Cancer Network guidelines:

- **Hairy Cell Leukemia:** Guidelines (version 1.2026 – August 11, 2025) recommend Nipent as preferred therapy as a single agent for initial therapy.² Nipent is also recommended as a single agent or in combination with a rituximab product (e.g., Rituxan, biosimilars) for less than a complete response to initial therapy or relapsed disease within 2 years of complete response following initial treatment with cladribine.^{2,3} Nipent is also recommended with a rituximab product for retreatment for relapse \geq 2 years following initial treatment and for relapse \geq 2 years following initial treatment with cladribine.
- **Graft-Versus-Host Disease:** Guidelines for Hematopoietic Cell Transplantation (version 2.2025 – June 3, 2025) recommend Nipent, in combination with corticosteroids, for acute or chronic graft-versus-host disease following no response to first-line therapy options (steroid-refractory).^{2,4} Nipent is also recommended as part of a conditioning regimen for hematopoietic cell transplant.
- **Primary Cutaneous Lymphoma:** Guidelines (version 3.2025 – June 10, 2025) recommend Nipent as a single agent for the subsequent treatment of disease refractory to multiple previous therapies.^{2,5}
- **T-Cell Lymphomas:** Guidelines (version 2.2025 – May 28, 2025) recommend Nipent as second-line therapy, as a single agent, for T-cell large granular lymphocytic leukemia, and in combination with Campath® (alemtuzumab intravenous infusion and subcutaneous injection) or as a single agent for T-cell prolymphocytic leukemia and hepatosplenic T-cell lymphoma.^{2,6}

Safety

Nipent has a Boxed Warning for dose-limiting severe renal, liver, pulmonary, and central nervous system toxicities when used at higher than recommended doses.¹ The use of Nipent in combination with fludarabine is not recommended due to severe or fatal pulmonary toxicity.

POLICY STATEMENT

Prior Authorization is recommended for medical benefit coverage of Nipent. Approval is recommended for those who meet the **Criteria** and **Dosing** for the listed indications. Extended approvals are allowed if the patient continues to meet the Criteria and Dosing. Requests for doses outside of the established dosing documented in this policy will be considered on a case-by-case basis by a clinician (i.e., Medical Director or Pharmacist). All approvals are provided for the duration noted below. In cases where the approval is

authorized in months, 1 month is equal to 30 days. Because of the specialized skills required for evaluation and diagnosis of patients treated with Nipent as well as the monitoring required for adverse events and long-term efficacy, approval requires Nipent to be prescribed by or in consultation with a physician who specializes in the condition being treated.

Automation: None.

RECOMMENDED AUTHORIZATION CRITERIA

Coverage of Nipent is recommended in those who meet one of the following criteria:

FDA-Approved Indication

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- 1. Hairy Cell Leukemia.** Approve for 6 months if the patient meets BOTH of the following (A and B):
 - Patient is \geq 18 years of age; AND
 - The medication is prescribed by or in consultation with an oncologist.

Dosing. Approve up to 4 mg/m^2 administered intravenously no more frequently than once every 2 weeks.

Other Uses with Supportive Evidence

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- 2. Graft-Versus-Host Disease.** Approve for 6 months if the patient meets ALL of the following (A, B, and C):
 - Patient has steroid-refractory disease; AND
 - The medication will be used in conjunction with systemic corticosteroids; AND

Note: Examples of corticosteroids include prednisone and methylprednisolone.

 - The medication is prescribed by or in consultation with an oncologist.

Dosing. Approve up to 1.5 mg/m^2 administered intravenously no more frequently than three times in each 14-day cycle.

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- 3. Hematopoietic Cell Transplantation.** Approve for 1 month if the patient meets ALL of the following (A, B, C, and D):
 - Patient is \geq 4 years of age; AND
 - The medication is used for reduced intensity conditioning; AND
 - The medication is used in combination with ONE of the following (i, ii, or iii):
 - Busulfan; OR
 - Busulfan plus cyclophosphamide; OR
 - Cyclophosphamide plus total body irradiation; AND
 - The medication is prescribed by or in consultation with an oncologist or a physician that specializes in hematopoietic cell transplantation.

Dosing. Approve up to 4 mg/m^2 administered intravenously twice prior to hematopoietic cell transplantation.

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- 4. Mycosis Fungoides/Sezary Syndrome.** Approve for 6 months if the patient meets ALL of the following (A, B, and C):
- A) Patient is \geq 18 years of age; AND
 - B) Patient has relapsed or refractory disease; AND
 - C) The medication is prescribed by or in consultation with an oncologist or dermatologist.

Dosing. Approve up to 5 mg/m² administered intravenously no more frequently than three times in each 21-day cycle.

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- 5. T-Cell Lymphoma.** Approve for 6 months if the patient meets ALL of the following (A, B, and C):
- Note: Examples of T-cell lymphoma include hepatosplenic T-cell lymphoma, T-cell prolymphocytic leukemia, and T-cell large granular lymphocytic leukemia.
- A) Patient is \geq 18 years of age; AND
 - B) Patient meets ONE of the following (i or ii):
 - i. Patient has hepatosplenic T-cell lymphoma or T-cell prolymphocytic leukemia; OR
 - ii. Patient meets BOTH of the following (a and b):
 - a) Patient has T-cell large granular lymphocytic leukemia; AND
 - b) Patient has progressive or refractory disease; AND
 - C) The medication is prescribed by or in consultation with an oncologist.

Dosing. Approve up to 4 mg/m² administered intravenously no more frequently than once weekly.

CONDITIONS NOT RECOMMENDED FOR APPROVAL

Coverage of Nipent is not recommended in the following situations:

- 1.** Coverage is not recommended for circumstances not listed in the Recommended Authorization Criteria. Criteria will be updated as new published data are available.

REFERENCES

1. Nipent intravenous infusion [prescribing information]. Lake Forest, IL: Hospira; October 2019.
2. The NCCN Drugs and Biologics Compendium. © 2025 National Comprehensive Cancer Network. Available at: <http://www.nccn.org>. Accessed on September 3, 2025. Search term: pentostatin.
3. The NCCN Hairy Cell Leukemia Clinical Practice Guidelines in Oncology (version 1.2026 – August 11, 2025). © 2025 National Comprehensive Cancer Network. Available at: <http://www.nccn.org>. Accessed on September 3, 2025.
4. The NCCN Hematopoietic Cell Transplantation Clinical Practice Guidelines in Oncology (version 2.2025 – June 3, 2025). © 2025 National Comprehensive Cancer Network. Available at: <http://www.nccn.org>. Accessed on September 3, 2025.
5. The NCCN Primary Cutaneous Lymphomas Clinical Practice Guidelines in Oncology (version 3.2025 – June 10, 2025). © 2025 National Comprehensive Cancer Network. Available at: <http://www.nccn.org>. Accessed on September 3, 2025.
6. The NCCN T-Cell Lymphomas Clinical Practice Guidelines in Oncology (version 2.2025 – May 28, 2025). © 2025 National Comprehensive Cancer Network. Available at: <http://www.nccn.org>. Accessed on September 3, 2025.
7. Ragon BK, Mehta RS, Gulbis AM, et al. Pentostatin therapy for steroid-refractory acute graft versus host disease: Identifying those who may benefit. *Bone Marrow Transplant.* 2018;53:315-325.
8. Tsimberidou AM, Giles F, Duvic M, et al. Phase II study of pentostatin in advanced T-cell lymphoid malignancies. *Cancer.* 2004;100:342-349.
9. Ravandi F, Aribi A, O'Brien S, et al. Phase II study of alemtuzumab in combination with pentostatin in patients with T-cell neoplasms. *J Clin Oncol.* 2009;27:5425-5430.
10. Dimitrova D, Kanakry JA. Beyond fludarabine: Pentostatin plus cyclophosphamide are well-tolerated alternative in reduced intensity conditioning. *Bone Marrow Transplant.* 2020;57:1837-1838.
11. Dimitrova D, Gea-Banacloche J, Steinberg SM, et al. Prospective study of a novel, radiation-free, reduced-intensity bone marrow transplantation platform for primary immunodeficiency diseases. *Biol Blood Marrow Transplant.* 2020;26:94-106.

HISTORY

Type of Revision	Summary of Changes	Review Date
Annual Revision	No criteria changes.	09/21/2022
Annual Revision	Hematopoietic Cell Transplantation: Added new condition of approval. T-Cell Prolymphocytic Leukemia: Medication used as a single agent added as new option for approval.	09/20/2023
Annual Revision	No criteria changes.	09/11/2024
UCare P&T Review	Policy reviewed and approved by UCare P&T committee. Annual review process	09/16/2024
Annual Revision	Hairy Cell Leukemia: The requirements that the medication is used as a single agent and medication is used in combination with rituximab were removed. Mycosis Fungoides/Sezary Syndrome: The requirement that the medication will be used as a single agent was removed. T-Cell Lymphoma: T-cell large granular lymphocytic leukemia, T-cell prolymphocytic leukemia, and hepatosplenic T-cell lymphoma were moved to be addressed under T-Cell lymphoma. For T-cell large granular lymphocytic leukemia, the requirement that the medication will be used as a single agent was removed. For T-cell prolymphocytic leukemia and hepatosplenic T-cell lymphoma, the requirements that the medication is used as a single agent and that medication is used in combination with Campath were removed.	09/10/2025
UCare P&T Review	Policy reviewed and approved by UCare P&T committee. Annual review process	09/15/2025