# Medical Drug Clinical Criteria

Subject:	Polivy (polatuzumab ve	Polivy (polatuzumab vedotin-piiq)		
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Overview				

This document addresses the use of Polivy (polatuzumab vedotin-piiq). Polivy is a monoclonal antibody-drug conjugate (ADC) that consists of a humanized IgG1 antibody specific for CD79b and a small molecule, monomethyl auristatin E (MMAE), a microtubuledisrupting agent, and a protease cleavable linker. The anticancer activity is due to the binding of the ADC to CD79b-expressing cells, cleavage of MMAE component, and killing dividing cells by inhibiting cell division and inducing apoptosis. The target CD79b is a surface protein found exclusively on B-cells and Polivy is indicated to treat diffuse large B-cell lymphoma (DLBCL).

The FDA approved indications for Polivy include in combination with bendamustine and a rituximab product for the treatment of adult patients with relapsed or refractory diffuse large B-cell lymphoma, not otherwise specified, after at least two prior therapies. Accelerated approval was based on positive results from a phase 2 trial comparing Polivy plus bendamustine and rituximab (BR) to BR alone. Patients included in this study were not eligible for autologous hematopoietic stem cell transplantation (HSCT). Polivy is also FDA approved for previously untreated diffuse large B-cell lymphoma, not otherwise specified or high-grade B-cell lymphoma in combination with a rituximab product, cyclophosphamide, doxorubicin, and prednisone (R-CHP) for individuals who have an International Prognostic Index score of 2 or greater. The National Comprehensive Cancer Network<sup>®</sup> (NCCN) provides additional recommendations with a category 1 or 2A level of evidence for the use of Polivy. These include its use as first-line treatment for previously untreated DLBCL and high-grade B-cell lymphomas in combination with R-CHP (category 1); as well as second-line or subsequent therapy for relapsed/refractory DLBCL, high-grade B-cell lymphomas, HIV-related lymphomas, and B-Cell Post-Transplant lymphoproliferative disorders (category 2A). NCCN also recommends Polivy as a single agent or in combination with bendamustine or rituximab for relapsed/refractory B-cell lymphomas, or as a bridging option until a CAR T-cell product is available.

### **Definitions and Measures**

Hematopoietic stem cells: Primitive cells capable of replication and formation into mature blood cells in order to repopulate the bone marrow.

Line of Therapy:

- First-line therapy: The first or primary treatment for the diagnosis, which may include surgery, chemotherapy, radiation therapy or a combination of these therapies.
- Second-line therapy: Treatment given when initial treatment (first-line therapy) is not effective or there is disease progression.
- Third-line therapy: Treatment given when both initial (first-line therapy) and subsequent treatment (second-line therapy) are not effective or there is disease progression.

Monoclonal antibody: A protein developed in the laboratory that can locate and bind to specific substances in the body and on the surface of cancer cells.

Non-Hodgkin Lymphoma (NHL): A group of malignant solid tumors or lymphoid tissues.

Refractory Disease: Illness or disease that does not respond to treatment.

Relapse or recurrence: After a period of improvement, during which time a disease (for example, cancer) could not be detected, the return of signs and symptoms of illness or disease. For cancer, it may come back to the same place as the original (primary) tumor or to another place in the body.

**Clinical Criteria** 

When a drug is being reviewed for coverage under a member's medical benefit plan or is otherwise subject to clinical review (including prior authorization), the following criteria will be used to determine whether the drug meets any applicable medical necessity requirements for the intended/prescribed purpose.

#### Polivy (polatuzumab vedotin-piiq)

Requests for Polivy (polatuzumab vedotin-piiq) may be approved if the following criteria are met:

- I. Individual has a diagnosis of relapsed or refractory diffuse large B-cell lymphoma (DLBCL), not otherwise specified (including high-grade B-cell lymphomas); <u>HIV-related B-cell lymphoma</u>; or monomorphic post-transplant lymphoproliferative disorder (B-cell type) (Label, NCCN 2A); <u>AND</u>
- II. Individual is using as a single agent or in combination with in combination with bendamustine and/<u>or</u> a rituximab (including rituximab biosimilars) (Label, NCCN 2A); AND
- III. Individual has received at least one prior lines of therapy (NCCN 2A); AND
- IV. Individual is ineligible for autologous hematopoietic stem cell transplantation (HSCT);

#### OR

- Individual has a diagnosis of relapsed or refractory diffuse large B-cell lymphoma (DLBCL), not otherwise specified (including high-grade B-cell lymphomas); <u>HIV-related B-cell lymphoma; or monomorphic post-transplant lymphoproliferative disorder (Bcell type);</u> AND
- Individual is using as a bridging option (typically 1 or more cycles as necessary) until CAR T-cell product is available (NCCN 2A);

#### OR

- VII. Individual has a diagnosis of previously untreated diffuse large B-cell lymphoma (DLBCL), not otherwise specified (including high-grade B-cell lymphomas); AND
- VIII. Individual is using in combination with a rituximab product (including rituximab biosimilars), cyclophosphamide, doxorubicin, and prednisone (Pola-R-CHP); AND
- IX. Individual has international prognostic index for diffuse large B-cell Lymphoma (IPI) 2 or higher.

Requests for Polivy (polatuzumab vedotin-piiq) may not be approved when the above criteria are not met and for all other indications.

# Coding

The following codes for treatments and procedures applicable to this document are included below for informational purposes. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member coverage or provider reimbursement policy. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

## HCPCS

J9309 Injection, polatuzumab vedotin-piiq, 1 mg [Polivy]

#### **ICD-10** Diagnosis

B20	Human Immunodeficiency Virus (HIV) disease (when specified as HIV-related B-cell lymphoma)
C82.00-C82.99	Follicular Lymphoma
C83.30-C83.39	Diffuse large B-cell lymphoma
C85.10-C85.29	Unspecified B-cell lymphoma/ Mediastinal (thymic) large B-cell lymphoma
D47.Z1	Post-transplant lymphoproliferative disorder (PTLD)

# **Document History**

Revised: 05/17/2024

- Document History:
  - 05/17/2024 Annual Review: Add additional types of B-cell lymphoma per NCCN; add option for single agent use or combination with rituximab or bendamustine only per NCCN. Coding Reviewed: Added ICD-10-CM B20, D47.Z1.
  - 11/17/2023 Select Review: Remove duplicative criteria and clarify criteria for Pola-R-CHP regimen. Coding Reviewed: No changes.
  - 08/18/2023 Select Review: Reword POLA-R-CHP criteria. Coding Reviewed: No changes.
  - 05/19/2023 Annual Review: add DLBCL stage 2 mesenteric and fda approval for untreated DLBCL. Coding Reviewed: No changes.

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- 05/20/2022 Annual Review: Clarify prior therapy requirement and add use as bridging therapy to CAR-T per NCCN. Coding Reviewed: No changes.
- 05/21/2021 Annual Review: No changes. Coding Reviewed: No changes.
- 05/15/2020 Annual Review: Clarify use in combination with rituximab to include biosimilar. Coding reviewed: No changes.
- 08/16/2019 Annual Review: Add clinical criteria for new agent Polivy. Coding Reviewed: Added HCPCS codes J9999, C9399. All Diagnosis codes allowed. Coding change update: Delete J9999 and C9399 for Polivy. Add J9309 for Polivy effective 1/1/2020. Deleted All Diagnoses and added C82.00-C82.99, C83.30-C83.39, C85.10-C85.29.

# References

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  DrugPoints® System [electronic version]. Truven Health Analytics, Greenwood Village, CO. Updated periodically.
- Lexi-Comp ONLINET<sup>™</sup> with AHFS<sup>™</sup>, Hudson, Ohio: Lexi-Comp, Inc.; 2024; Updated periodically. Morschhauser F, Flinn IW, Advani R, et al. Polatuzumab vedotin or pinatuzumab vedotin plus rituximab in patients with relapsed or 3. 4. refractory non-Hodgkin lymphoma: final results from a phase 2 randomised study (ROMULUS). Lancet Haematol. 2019
  - May;6(5):e254-e265. doi: 10.1016/S2352-3026(19)30026-2. Epub 2019 Mar 29.
- Sehn LH, Herrera AF, Flowers CR, et al. Polatuzumab Vedotin in Relapsed or Refractory Diffuse Lage B-Cell Lymphoma. J Clin 5. Oncol 2020; 38:155-165.
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Federal and state laws or requirements, contract language, and Plan utilization management programs or polices may take precedence over the application of this clinical criteria.

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