

Polivy (polatuzumab vedotin-piiq)
 Effective 06/01/2020

Plan	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Benefit	<input type="checkbox"/> Pharmacy Benefit <input checked="" type="checkbox"/> Medical Benefit		
Specialty Limitations	N/A		
Contact Information	Medical and Specialty Medications		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
Exceptions	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029

Overview

Polatuzumab vedotin is an antibody drug conjugate (ADC) directed at CD79b which consists of 3 components: 1) a CD79b-specific humanized IgG1 antibody; 2) a microtubule-disrupting agent, monomethylauristatin E (MMAE); and 3) a protease cleavable linker (which covalently conjugates MMAE to the polatuzumab antibody). The conjugate binds to CD79b (B-cell specific cell surface protein commonly expressed in mature B cell lymphomas) and forms a complex which is internalized within the cell and releases MMAE. MMAE binds to the tubules and disrupts the cellular microtubule network, inducing cell cycle arrest (G2/M phase) and apoptosis.

Coverage Guidelines

Authorization may be granted for members who are currently receiving treatment with Polivy excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted for members when all the following criteria are met, and documentation is provided:

1. The member is \geq 18 years of age
2. The member has a diagnosis of relapsed or refractory diffuse large B-cell lymphoma (DLBCL)
3. Provider specialty is oncology and/or hematology or medication is being prescribed in consultation with an oncologist/hematologist
4. The member has had adverse reaction, inadequate response, or contraindication to two systemic therapies for DLBCL

Continuation of Therapy

Reauthorization requires physician documentation of improvement of member's condition.

Limitations

Authorizations will be approved for 12 months

References

1. Polivy [package insert]. South San Francisco, CA: Genentech, Inc.; June 2019.
2. National Comprehensive Cancer Network. NCCN Clinical Practice Guidelines in Oncology: B-Cell Lymphomas. Version 4.2019. <https://www.nccn.org>. Accessed June 19, 2019.
3. The NCCN Drugs & Biologics Compendium® © 2019 National Comprehensive Cancer Network, Inc. <https://www.nccn.org>. Accessed June 19, 2019.

Review History

01/23/2020 – Reviewed P&T Mtg (effective 6/1/20).

