

Poteligeo (mogamulizumab-kpkc)
Effective 07/01/2025

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
Contact Information	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Poteligeo (mogamulizumab-kpkc) is a CC chemokine receptor type 4 (CCR4)-directed monoclonal antibody indicated for treatment of relapsed or refractory mycosis fungoides (MF) or Sézary syndrome (SS) after at least one prior systemic therapy.

All other indications are considered experimental/investigational and are not a covered benefit.

Coverage Guidelines

Authorization may be granted for members new to the plan within the past 90 days who are currently receiving treatment with the requested medication, excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

Or

Authorization may be granted when all the following criteria are met, and clinical documentation has been submitted:

- Member has one of the following diagnoses:
 - Relapsed or refractory mycosis fungoides
 - Sézary syndrome
- Prescriber is an oncologist or hematologist
- Member has received at least one prior systemic therapy (see Appendix), which resulted in an inadequate response
- Dosing is appropriate based on member's current weight

Limitations

Approvals will be granted for 12 months.

Appendix

First-line systemic therapies for mycosis fungoides or Sézary syndrome

- Extracorporeal phototherapy

2. Oral retinoids (bexarotene, tretinoin capsules, isotretinoin capsules)
3. Interferons (Pegasys/Intron-A)
4. HDAC inhibitors (vorinostat, romidepsin)
5. Methotrexate or pralatrexate
6. Adcetris (brentuximab)
7. Cyclophosphamide

References

1. Avilés A, Nambo MJ, Neri N, et al. Interferon and low dose methotrexate improve outcome in refractory mycosis fungoides/Sézary syndrome. *Cancer Biother Radiopharm* 2007; 22:836
2. Booken N, Weiss C, Utikal J, et al. Combination therapy with extracorporeal photopheresis, interferon-alpha, PUVA and topical corticosteroids in the management of Sézary syndrome. *J Dtsch Dermatol Ges* 2010; 8:428
3. Burg G, Dummer R. Historical perspective on the use of retinoids in cutaneous T-cell lymphoma (CTCL). *Clin Lymphoma* 2000; 1 Suppl 1:S41
4. Horwitz SM, Kim YH, Foss F, et al. Identification of an active, well-tolerated dose of pralatrexate in patients with relapsed or refractory cutaneous T-cell lymphoma. *Blood* 2012; 119:4115.
5. Kim YH, Bagot M, Pinter-Brown L, et al; MAVORIC Investigators. Mogamulizumab versus vorinostat in previously treated cutaneous T-cell lymphoma (MAVORIC): an international, open-label, randomized, controlled phase 3 trial [published online August 9, 2018]
6. Kim YH, Demierre MF, Kim EJ, et al. Clinically meaningful reduction in pruritus in patients with cutaneous T-cell lymphoma treated with romidepsin. *Leuk Lymphoma* 2013; 54:284
7. Poteligeo (mogamulizumab-kpkc) [prescribing information]. Princeton, NJ: Kyowa Kirin, Inc; January 2025.
8. Scarisbrick JJ, Child FJ, Clift A, et al. A trial of fludarabine and cyclophosphamide combination chemotherapy in the treatment of advanced refractory primary cutaneous T-cell lymphoma. *Br J Dermatol* 2001; 144:1010
9. Stephen S, Morrissey KA, Benoit BM, et al. Inhibition of cell-mediated immunity by the histone deacetylase inhibitor vorinostat: implications for therapy of cutaneous T-cell lymphoma. *Am J Hematol* 2012; 87:226

Review History

04/17/2019 – Reviewed

05/20/2020 – Reviewed and Updated May P&T; updated overview; added started and stabilized statement. Effective 8/1/20.

06/11/2025 – Reviewed and Updated at June P&T. Updated language for members who are new to the Plan. Effective 07/01/2025.

