

Rezurock (belumosudil)
Effective 10/01/2023

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 checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit		
Specialty Limitations	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
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
Exceptions	N/A		

Overview

Rezurock is indicated for the treatment of adult and pediatric patients 12 years and older with chronic graft-versus-host disease (cGVHD) after failure of at least two prior lines of systemic therapy.

Coverage Guidelines

Authorization may be granted for members new to General Brigham Health Plan who are currently receiving treatment with Rezurock excluding when the product is obtained as samples or via manufacturer's patient assistance programs.

OR

Authorization may be granted for members meeting ALL the following criteria:

1. Member has a diagnosis of chronic graft versus host disease (cGVHD)
2. Medical charts confirming member has failed two or more lines of systemic therapy
3. The member is at least 12 years of age

Continuation of Therapy

Reauthorization of 12 months may be granted for continued treatment when ALL the following criteria is met:

1. The member does not have evidence of unacceptable toxicity while on the current regimen
2. The member has not experienced clinically significant progression of cGVHD (i.e., progression that requires new systemic therapy) while on the current regimen

Limitations

Initial approvals and reauthorizations will be granted for 12 months.

References

1. Rezurock [package insert]. Bridgewater, NJ: Kadmon Pharmaceuticals; July 2022

Review History

08/9/2023 - Reviewed at August P&T, Effective 10/1/23

