

Welireg (belzutifan) Effective 05/01/2022

Plan	 MassHealth UPPL Commercial/Exchange 	Due sure Ture	Prior Authorization
Benefit	 Pharmacy Benefit Medical Benefit 	Program Type	 Quantity Limit Step Therapy
Specialty	This medication has been designated specialty and must be filled at a contracted		
Limitations	specialty pharmacy.		
	Medical and Specialty Medications		
Contact Information	All Plans P	hone: 877-519-1908	Fax: 855-540-3693
	Non-Specialty Medications		
	All Plans P	hone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Welireg (belzutifan) is indicated for the treatment of adult patients with von Hippel-Lindau (VHL) disease who require therapy for associated renal cell carcinoma (RCC), central nervous system (CNS) hemangioblastomas, or pancreatic neuroendocrine tumors (pNET), not requiring immediate surgery.

Coverage Guidelines

Authorization may be reviewed for members new to the plan 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 if the member meets all following criteria and documentation has been submitted:

- 1. The member has a diagnosis of von Hippel-Lindau (VHL) disease associated renal cell carcinoma (RCC), central nervous system (CNS) hemangioblastomas, or pancreatic neuroendocrine tumors (pNET)
- 2. The member does not require immediate surgery
- 3. The member does not have metastatic disease
- 4. Medication will be used as a single agent

Continuation of Therapy

Reauthorization may be granted with physician documentation of no evidence of unacceptable toxicity or disease progression while on treatment.

Limitations

1. Initial approvals and reauthorizations will be granted for: 12 months

References

1. Welireg [package insert]. Whitehouse Station, NJ: Merck Sharp & Dohme Corp.; August 2021.

Review History

03/16/2022 – Created for March P&T Effective 05/01/2022.