

Rybrevant[®] (amivantamab-vmjw) Effective 04/01/2022

Plan	 ☑ MassHealth □ MH UPPL □Commercial/Exchange 		Program Type	☑ Prior Authorization ☐ Quantity Limit
Benefit	□ Pharmacy Benefit⊠ Medical Benefit (NLX)		0 //	□ Step Therapy
Specialty Limitations	N/A			
	Specialty Medications			
	All Plans	Ph	one: 866-814-5506	Fax: 866-249-6155
	Non-Specialty Medications			
Contact Information	MassHealth	Ph	Phone: 877-433-7643 Fax: 866-255-7569	
	Commercial	Ph	Phone: 800-294-5979 Fax: 888-836-0730	
	Exchange	Ph	one: 855-582-2022	Fax: 855-245-2134
	Medical Specialty Medications (NLX)			
	All Plans	Ph	one: 844-345-2803	Fax: 844-851-0882
Exceptions	N/A			

Overview

Rybrevant is indicated for the treatment of locally advanced or metastatic non–small cell lung cancer (NSCLC) in adults with epidermal growth factor receptor (EGFR) exon 20 insertion mutations (as detected by an approved test) with disease progression on or after platinum-based chemotherapy.

Coverage Guidelines

Authorization may be reviewed for members new to AllWays Health Partners who are currently receiving treatment with Rybrevant 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 has a diagnosis of advanced or metastatic non-small cell lung cancer (NSCLC)
- 2. The prescriber specialty is an oncologist or medication is written in consultation with an oncologist
- 3. Appropriate dosing
- 4. Cancer has EGFR exon 20 insertion mutation
- 5. Member meets ONE Of the following:
 - Inadequate response or adverse reaction to at least one platinum-based chemotherapy regimen (e.g., carboplatin, oxaliplatin, cisplatin)
 - Contraindication to ALL platinum-based chemotherapy (e.g., carboplatin, oxaliplatin, cisplatin)

399 Revolution Drive, Suite 810, Somerville, MA 02145 | allwayshealthpartners.org



Continuation of Therapy

Reauthorization will be granted when physician provides attestation of positive response to therapy and member has not shown signs of excessive toxicity.

Limitations

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

References

1. Rybrevant (amivantamab) [prescribing information]. Horsham, PA: Janssen Biotech Inc; July 2021.

Review History

01/19/2022 - Reviewed and Created Jan P&T. Effective 04/01/2022.

Disclaimer

AllWays Health Partners complies with applicable federal civil rights laws and does not discriminate or exclude people on the basis of race, color, national origin, age, disability, or sex.