

## STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

## **DACOMITINIB**

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
DACOMITINIB	VIZIMPRO	45283		GPI-10	
				(2136001900)	

#### **GUIDELINES FOR USE**

- 1. Does the patient have a diagnosis of metastatic non-small cell lung cancer (NSCLC) and meet **ALL** of the following criteria?
  - The patient has epidermal growth factor receptor (EGFR) exon 19 deletion or exon 21 L858R substitution mutations as detected by an FDA-approved test
  - Vizimpro will be used as first-line treatment
  - Vizimpro will NOT be used concurrently with an epidermal growth factor receptor (EGFR) tyrosine kinase-inhibitor (e.g., Tarceva [erlotinib], Tagrisso [osimertinib], Iressa [gefitinib])

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #1 per day. If no, do not approve.

DENIAL TEXT: \*Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **DACOMITINIB** (**Vizimpro**) requires the following rule(s) be met for approval:

- A. You have metastatic non-small cell lung cancer (type of cancer that has spread) to other parts of the body)
- B. You have epidermal growth factor receptor (EGFR) exon 19 deletion or exon 21 L858R substitution mutations (types of gene mutations) as detected by an FDA (Food and Drug Administration)-approved test
- C. Vizimpro will be used as first-line treatment
- D. You will NOT be using Vizimpro concurrently (at the same time) with an epidermal growth factor receptor (EGFR) tyrosine kinase-inhibitor (such as Tarceva [erlotinib], Tagrisso [osimertinib], Iressa [gefitinib])

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

### **CONTINUED ON NEXT PAGE**

Copyright © 2022 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 5/27/2022 Page 1 of 2



# STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

### **DACOMITINIB**

### **RATIONALE**

For further information, please refer to the Prescribing Information and/or Drug Monograph for Vizimpro.

### **REFERENCES**

Vizimpro [Prescribing Information]. New York, NY: Pfizer Labs; December 2020.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 11/18

Commercial Effective: 07/01/22 Client Approval: 05/22 P&T Approval: 04/22

Copyright © 2022 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 5/27/2022 Page 2 of 2