

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

UMBRALISIB

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
UMBRALISIB TOSYLATE	UKONIQ	47104		GPI-10	
				(2153308040)	

GUIDELINES FOR USE

- 1. Does the patient have a diagnosis of relapsed or refractory marginal zone lymphoma (MZL) and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient has received at least one prior anti-CD20-based regimen

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #4 per day. If no, continue to #2.

- 2. Does the patient have a diagnosis of relapsed or refractory follicular lymphoma (FL) and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient has received at least three prior lines of systemic therapy

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #4 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 **UMBRALISIB** (Ukoniq) requires the following rule(s) be met for approval:

- A. You have relapsed or refractory marginal zone lymphoma or follicular lymphoma (types of immune system cancer that have returned or are not responding to treatment)
- B. You are 18 years of age or older
- C. If you have marginal zone lymphoma, approval also requires:
 - 1. You have received at least one prior anti-CD20-based regimen (type of cancer treatment)
- D. If you have follicular lymphoma, approval also requires:
 - You have received at least three prior lines of systemic therapy (treatment that travels throughout the body)

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 © 2021 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.

2/17/2021 Page 1 of 2



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

UMBRALISIB

RATIONALE

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

REFERENCES

Ukoniq [Prescribing Information]. Edison, NJ: TG Therapeutics, Inc.; February 2021.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 02/21

Commercial Effective: 03/01/21 Client Approval: 02/21 P&T Approval: 01/21

2/17/2021 Page 2 of 2