

# STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

### **DUVELISIB**

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
DUVELISIB	COPIKTRA	45269		GPI-10	
				(2153803000)	

#### **GUIDELINES FOR USE**

- 1. Does the patient have a diagnosis of relapsed or refractory chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) and meet **ALL** of the following criteria?
  - The patient is 18 years of age or older
  - The patient has received at least two prior therapies for CLL or SLL

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #2 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 **DUVELISIB** (Copiktra) requires the following rule(s) be met for approval:

- A. You have ONE of the following diagnoses:
  - 1. Relapsed or refractory chronic lymphocytic leukemia (CLL: a type of blood cancer that has returned after treatment or does not fully respond to treatment)
  - 2. Small lymphocytic lymphoma (SLL: a type of blood cancer)
- B. You are 18 years of age or older
- C. You have received at least two prior therapies for chronic lymphocytic leukemia or small lymphocytic lymphoma

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.

## **RATIONALE**

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

## **REFERENCES**

Copiktra [Prescribing Information]. Needham, MA: Verastem, Inc.; December 2021.

Library	Commercial	NSA
Yes	Yes	No

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

Commercial Effective: 04/01/22 Client Approval: 03/22 P&T Approval: 10/18

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