

# STANDARD NON-SELF-ADMINISTERED DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

## **SUTIMLIMAB-JOME (INTERIM)**

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
SUTIMLIMAB-JOME	ENJAYMO	47799		GPI-10	
				(8580008530)	

## **GUIDELINES FOR USE**

- 1. Does the patient have a diagnosis of cold agglutinin disease (CAD) **AND** meet the following criterion?
  - The patient is 18 years of age or older

If yes, approve for a total of 12 months by HICL or GPI-10 as follows: Initial request:

- FIRST APPROVAL: Approve for 1 month with a quantity limit of #462 mL per 28 days
- SECOND APPROVAL: Approve for 11 months with a quantity limit of #308 per 28 days. (Start date is ONE WEEK BEFORE the end date of first approval)

## **Subsequent requests:**

• Approve for 12 months with a quantity limit of #308mL per 28 days.

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 **SUTIMLIMAB-JOME** (**Enjaymo**) requires the following rule(s) be met for approval:

- A. You have cold agglutinin disease (CAD: a rare type of blood condition)
- B. You are 18 years of age or older

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 Enjaymo.

#### REFERENCES

Enjaymo [Prescribing Information]. Waltham, MA: Bioverativ USA, Inc.; February 2022.

Library	Commercial	NSA
Yes	No	Yes

Part D Effective: N/A Created: 02/22

Commercial Effective: 02/18/22 Client Approval: 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.

Created: 2/14/2022 Page 1 of 1