Medimpact

# STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

#### MARIBAVIR

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
MARIBAVIR	LIVTENCITY	47687		GPI-10 (1220005000)	

## **GUIDELINES FOR USE**

- 1. Does the patient have a diagnosis of post-transplant cytomegalovirus (CMV) infection and meet **ALL** of the following criteria?
  - The patient is 12 years of age or older
  - The patient is refractory to prior therapy with ganciclovir, valganciclovir, cidofovir or foscarnet

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 **MARIBAVIR (Livtencity)** requires the following rule(s) be met for approval:

- A. You have a post-transplant cytomegalovirus (CMV) infection (a type of viral infection)
- B. You are 12 years of age or older
- C. You are refractory to prior therapy with ganciclovir, valganciclovir, cidofovir or foscarnet

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 Livtencity.

#### REFERENCES

 Livtencity [Prescribing Information]. Lexington, MA: Takeda Pharmaceuticals America, Inc.; November 2021.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Commercial Effective:12/10/21 Created: 12/21 Client Approval:

P&T Approval: 10/21

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.