

# STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

### **PALBOCICLIB**

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
PALBOCICLIB	IBRANCE	41725		GPI-10	
				(2153106000)	

#### **GUIDELINES FOR USE**

- 1. Does the patient have a diagnosis of advanced or metastatic breast cancer and meet **ALL** the following criteria?
  - The patient's cancer is hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative

If yes, continue to #2.

If no, do not approve.

**DENIAL TEXT:** See the denial text at the end of the guideline.

- 2. Will Ibrance be used in combination with an aromatase inhibitor (e.g., anastrozole, letrozole, exemestane) **AND** the patient meets the following criterion?
  - The patient has NOT received prior endocrine-based therapy (e.g., letrozole, anastrozole, tamoxifen)

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #21 per 28 days. If no, continue to #3.

- 3. Will Ibrance be used in combination with Faslodex (fulvestrant) **AND** the patient meets the following criterion?
  - The patient has experienced disease progression following endocrine therapy (e.g., letrozole, anastrozole, tamoxifen)

If yes, approve for 12 months by HICL or GPI-10 with a quantity limit of #21 per 28 days. If no, do not approve.

**DENIAL TEXT:** See the denial text at the end of the guideline.

#### CONTINUED ON NEXT PAGE

Copyright © 2023 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: 5/23/2023 Page 1 of 2



# STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

### **PALBOCICLIB**

## **GUIDELINES FOR USE (CONTINUED)**

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 **PALBOCICLIB** (**Ibrance**) requires the following rule(s) be met for approval:

- A. You have advanced or metastatic breast cancer (cancer that has worsened or has spread to other parts of the body)
- B. Your cancer is hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative (a type of protein)
- C. You meet ONE of the following:
  - 1. Ibrance will be used in combination with an aromatase inhibitor (such as anastrozole, letrozole, exemestane) AND you have not received prior endocrine (hormone)-based therapy (such as letrozole, anastrozole, tamoxifen)
  - 2. Ibrance will be used in combination with Faslodex (fulvestrant) AND your disease has worsened after endocrine (hormone) therapy (such as letrozole, anastrozole, tamoxifen)

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 Ibrance.

### REFERENCES

Ibrance [Prescribing Information]. New York, NY: Pfizer Laboratories. December 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 05/15

Commercial Effective: 07/01/23 Client Approval: 05/23 P&T Approval: 04/23

Copyright © 2023 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: 5/23/2023 Page 2 of 2