

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

TUCATINIB

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
TUCATINIB	TUKYSA	46459		GPI-10	
				(2153408000)	

GUIDELINES FOR USE

- 1. Does the patient have a diagnosis of advanced unresectable or metastatic HER2-positive breast cancer and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient has received one or more prior anti-HER2-based regimens (i.e., trastuzumab or trastuzumab with pertuzumab) in the metastatic setting
 - The requested medication will be used in combination with trastuzumab and capecitabine

If yes, approve for 12 months by GPID or GPI-14 for the requested strength with the following quantity limits:

Tukysa 50mg: #10 per day.Tukysa 150mg: #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 **TUCATINIB** (**Tukysa**) requires the following rule(s) be met for approval:

- A. You have advanced unresectable (cannot be removed with surgery) or metastatic (disease that has spread to other parts of the body) human epidermal growth factor receptor 2 (HER2: type of protein)-positive breast cancer
- B. You are 18 years of age or older
- C. You have previously received one or more anti-HER2-based treatment for metastatic disease (specifically either trastuzumab or trastuzumab with pertuzumab)
- D. The requested medication will be used in combination with trastuzumab and capecitabine

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 © 2020 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.

8/28/2020 Page 1 of 2



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

TUCATINIB

RATIONALE

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

REFERENCES

• Tukysa [Prescribing Information]. Bothell, WA: Seattle Genetics, Inc.; April 2020.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 08/20

Commercial Effective: 10/01/20 Client Approval: 08/20 P&T Approval: 07/20

8/28/2020 Page 2 of 2