

MED RX POLICY

POLICY: Oncology (Injectable) – Trastuzumab Products Med Rx Policy

- Herceptin[®] (trastuzumab intravenous infusion Genentech)
- Herceptin Hylecta[™] (trastuzumab and hyaluronidase-oysk subcutaneous injection Genentech)
- Herzuma® (trastuzumab-pkrb intravenous infusion Celltrion)
- Kanjinti[®] (trastuzumab-anns intravenous infusion Amgen)
- Ogivri® (trastuzumab-dkst intravenous infusion Mylan)
- Ontruzant® (trastuzumab-dttb intravenous infusion Merck)
- Trazimera® (trastuzumab-qyyp intravenous infusion Pfizer)

REVIEW DATE: 07/05/2023

OVERVIEW

Herzuma, Kanjinti, Ogivri, Ontruzant, and Trazimera are approved as biosimilars to Herceptin intravenous, indicating no clinically meaningful differences in safety and effectiveness and the same mechanism of action, route of administration, and dosage form as Herceptin intravenous.¹⁻⁷ However, minor differences in clinically inactive components are allowed. At this time, the trastuzumab biosimilars have only demonstrated biosimilarity, not interchangeability. Herceptin Hylecta is a combination of trastuzumab and hyaluronidase. It is a different formulation and dosage form of trastuzumab (not a Herceptin biosimilar). The hyaluronidase component helps increase the absorption rate of trastuzumab into the systemic circulation.

POLICY STATEMENT

This Med Rx program has been developed to encourage the use of the Preferred products. For all products (Preferred and Non-Preferred), the patient is required to meet the respective standard *Utilization Management Medical Policy* criteria. This program also directs the patient to try at least one Preferred Product prior to the approval of a Non-Preferred Product. Requests for Non-Preferred Products will be reviewed using the exception criteria (below). All approvals are provided for the duration noted in the respective standard *Utilization Management Medical Policy* criteria.

Automation: None.

Preferred Products: Kanjinti, Trazimera

Non-Preferred Products: Herceptin, Herceptin Hylecta, Herzuma, Ogivri, Ontruzant

RECOMMENDED EXCEPTION CRITERIA

Non-Preferred	Exception Criteria		
Products			
Herceptin	1. Approve if the patient meets BOTH of the following (A and B):		
Herzuma	A) Patient meets the standard Oncology (Injectable) - Trastuzum		
Ogivri	Products Utilization Management Medical Policy criteria; AND		
Ontruzant	B) Patient meets ONE of the following (i or ii):		
	i. Patient meets both of the following (a and b):		
	a) Patient has tried one of Kanjinti or Trazimera; AND		
	b) Patient cannot continue to use the Preferred Product due to a		
	formulation difference in the inactive ingredient(s) [e.g.,		
	differences in stabilizing agent, buffering agent, and/or		
	surfactant], which according to the prescriber, would result in a		
	significant allergy or serious adverse reaction; OR		
	ii. Patient is currently taking the requested agent.		
Herceptin Hylecta	1. Approve if the patient meets BOTH of the following (A <u>and</u> B):		
	A) Patient meets the standard Oncology (Injectable) – Herceptin Hylecta		
	Utilization Management Medical Policy criteria; AND		
	B) Patient meets ONE of the following (i, ii, <u>or</u> iii):		
	i. Patient has tried one of Kanjinti or Trazimera, but according to the		
	prescriber, cannot continue to use this product; OR		
	ii. Patient cannot continue on trastuzumab intravenous products due to		
	an inability to obtain or maintain intravenous access; OR		
	iii. Patient is currently taking the requested agent.		

REFERENCES

- 1. Herceptin® intravenous infusion [prescribing information]. South San Francisco, CA: Genentech; Februrary 2021.
- 2. Herzuma[®] intravenous infusion [prescribing information]. North Wales, PA: Teva; May 2019.
- 3. Kanjinti® intravenous infusion [prescribing information]. Thousand Oaks, CA: Amgen; October 2019.
- 4. Ogivri® intravenous infusion [prescribing information]. Steinhausen, Switzerland: Mylan; February 2021.
- 5. Trazimera[™] intravenous infusion [prescribing information]. New York, NY: Pfizer; November 2020.
- Herceptin Hylecta[™] subcutaneous injection [prescribing information]. South San Francisco, CA: Genentech; February 2019.
- 7. Ontuzant® intravenous infusion [prescribing information]. Whitehouse Station, NJ: Merck; March 2020.

HISTORY

Type of Revision	Summary of Changes	Review Date
Annual Revision	No criteria changes.	10/19/2022
Early Annual Revision	No criteria changes	07/05/2023