



Herceptin Hylecta

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Referring Provider Info: Same as Requesting Provider
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: Same as Referring Provider Same as Requesting Provider
Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____ kg
Patient Height: _____ cm

Please indicate the place of service for the requested drug:
 Ambulatory Surgical Home Off Campus Outpatient Hospital
 On Campus Outpatient Hospital Office Pharmacy

Exception Criteria Questions:

- A. Is the product being requested for the treatment of one of the following indications?
- HER2-overexpressing breast cancer
 - HER2-overexpressing metastatic gastric or gastroesophageal junction adenocarcinoma
- Yes No *If No, Skip to Clinical Criteria Questions*
- B. The preferred products for your patient's health plan are Herzuma, Kanjinti, Ogivri, Ontruzant and Trazimera. Can the patient's treatment be switched to any of the preferred products?
- Yes – Herzuma, *Please obtain Form for preferred product and submit for corresponding PA.*
 - Yes – Kanjinti, *Please obtain Form for preferred product and submit for corresponding PA.*
 - Yes – Ogivri, *Please obtain Form for preferred product and submit for corresponding PA.*
 - Yes – Ontruzant, *Please obtain Form for preferred product and submit for corresponding PA.*
 - Yes – Trazimera, *Please obtain Form for preferred product and submit for corresponding PA.*
 - No
- C. Does the patient have a documented intolerable adverse event to at least three of the preferred products (Herzuma, Kanjinti, Ogivri, Ontruzant, or Trazimera)? **Action Required: If 'Yes', attach supporting chart note(s).**
- Yes No
- D. Was the documented intolerable adverse event an expected adverse event attributed to the active ingredient as described in the prescribing information? **Action Required: If 'No', attach supporting chart note(s).**
- Yes No

Criteria Questions:

1. What is the patient's diagnosis?
- Breast cancer
 - Other _____
2. What is the ICD-10 code? _____
3. Will the requested drug be used OR is currently being used in one of the following clinical settings?
- Neoadjuvant treatment
 - Adjuvant treatment
 - Treatment of recurrent, unresectable advanced, or metastatic disease
 - Other _____
4. How many months of treatment has the patient received? _____ months
5. Has the patient received trastuzumab therapy for 12 months (52 weeks) or greater? Yes No
6. Is the request for a continuation of therapy with the requested drug? Yes No *If No, skip to #8*
7. Is there evidence of unacceptable toxicity or disease progression while on the current regimen?
- Yes No *No further questions*
8. What is the patient's human epidermal growth factor receptor 2 (HER2) status? **ACTION REQUIRED: Please attach documentation of human epidermal growth factor receptor 2 (HER2) status.**
- Positive Negative Unknown
9. Will the requested drug be used as part of a complete treatment regimen? Yes No

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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**CVS Caremark Specialty Pharmacy • 2211 Sanders Road NBT-6 • Northbrook, IL 60062
Phone: 1-888-877-0518 • Fax: 1-855-330-1720 • www.caremark.com**

Step Therapy Override: Complete if Applicable for the state of Maryland.	Please Circle	
Is the requested drug being used to treat stage four advanced metastatic cancer?	Yes	No
Is the requested drug's use consistent with the FDA-approved indication or the National Comprehensive Cancer Network Drugs & Biologics Compendium indication for the treatment of stage four advanced metastatic cancer and is supported by peer-reviewed medical literature?	Yes	No
Is the requested drug being used for an FDA-approved indication OR an indication supported in the compendia of current literature (examples: AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?	Yes	No
Does the prescribed quantity fall within the manufacturer's published dosing guidelines or within dosing guidelines found in the compendia of current literature (examples: package insert, AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?	Yes	No
Do patient chart notes document the requested drug was ordered with a paid claim at the pharmacy, the pharmacy filled the prescription and delivered to the patient or other documentation that the requested drug was prescribed for the patient in the last 180 days?	Yes	No
Has the prescriber provided proof documented in the patient chart notes that in their opinion the requested drug is effective for the patient's condition?	Yes	No

Step Therapy Override: Complete if Applicable for the state of Virginia.	Please Circle	
Is the requested drug being used for an FDA-approved indication or an indication supported in the compendia of current literature (examples: AHFS, Micromedex, current accepted guidelines)?	Yes	No
Does the prescribed dose and quantity fall within the FDA-approved labeling or within dosing guidelines found in the compendia of current literature?	Yes	No
Is the request for a brand drug that has an AB-rated generic equivalent or interchangeable biological product available?	Yes	No
Has the patient had a trial and failure of the AB-rated generic equivalent or interchangeable biological product due to an adverse event (examples: rash, nausea, vomiting, anaphylaxis) that is thought to be due to an inactive ingredient?	Yes	No
Is the preferred drug contraindicated?	Yes	No
Is the preferred drug expected to be ineffective based on the known clinical characteristics of the patient and the prescription drug regimen?	Yes	No
Has the patient tried the preferred drug while on their current or previous health benefit plan and it was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event?	Yes	No
Is the patient currently receiving a positive therapeutic outcome with the requested drug for their medical condition?	Yes	No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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