

Perjeta® (Pertuzumab) Prior Authorization Form

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Physician billing (HCPCS code: _____) Start Date (or date of next dose): _____

Dose: _____ Dosing Regimen: _____

Billing Provider Information

SoonerCare Provider ID: _____ Provider Name: _____

Provider Phone: _____ Provider Fax: _____

Prescriber Information

Prescriber NPI: _____ Prescriber Name: _____

Prescriber Phone: _____ Prescriber Fax: _____ Specialty: _____

Criteria

For Initial Authorization:

1. Is disease human epidermal receptor type 2 (HER2)-positive? Yes No
2. Please indicate the diagnosis and information:
 - Metastatic Breast Cancer**
 - A. Has member received prior anti-HER2 therapy or chemotherapy for metastatic disease? Yes No
 - B. Will pertuzumab be used in combination with trastuzumab and chemotherapy? Yes No
 - Neoadjuvant Treatment of Breast Cancer**
 - A. Is disease locally advanced, inflammatory, or early-stage breast cancer? Yes No
 - B. What is node status? Positive Negative
 - i. If tumor is node negative, is tumor >2cm in diameter? Yes No
 - C. Will pertuzumab be used in combination with trastuzumab and chemotherapy? Yes No
 - Adjuvant Treatment of Breast Cancer**
 - A. What is node status? Positive Negative
 - i. If tumor is node negative, indicate which of the following features are present. Please indicate all that apply:
 - tumor >1cm
 - tumor 0.5 to 1cm with histologic or nuclear grade 3
 - tumor 0.5 to 1cm with estrogen receptor (ER)/progesterone receptor (PR) negative
 - tumor 0.5 to 1cm and member age ≤35 years
 - B. Will pertuzumab be used in combination with trastuzumab and chemotherapy? Yes No
 - C. Will pertuzumab be used in combination with trastuzumab and docetaxel following doxorubicin/cyclophosphamide (AC)? Yes No
 - D. Will pertuzumab be used in combination with docetaxel/carboplatin/trastuzumab (TCH)? Yes No
 - E. Will pertuzumab be used in combination with trastuzumab following neoadjuvant therapy with paclitaxel or docetaxel and carboplatin/trastuzumab/pertuzumab? Yes No
 - Colorectal Cancer (CRC)**
 - A. Is disease RAS and BRAF mutation negative? Yes No
 - B. Will pertuzumab be used in combination with trastuzumab? Yes No
 - C. Will pertuzumab be used as first-line therapy? Yes No
 - i. Is the member a candidate for intensive therapy? Yes No
 - D. Will pertuzumab be used for the treatment of advanced or metastatic disease following disease progression? Yes No
 - If diagnosis is none of the above, please indicate diagnosis:** _____

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Criteria

Page 2 of 3—Please complete and return all pages. Failure to complete all pages will result in processing delays.

For Continued Authorization:

1. Date of last dose: _____
2. Does member have any evidence of progressive disease while on pertuzumab (when used for metastatic disease only)? Yes No
3. Has the member experienced any adverse drug reactions related to pertuzumab therapy? Yes No
 - a. If yes, please specify adverse reactions: _____

Prescriber Signature: _____ Date: _____

I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Failure to complete this form in full will result in processing delays.

Fax completed prior authorization request form to **888-601-8461** or submit Electronic Prior Authorization through CoverMyMeds® or SureScripts. All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at **AetnaBetterHealth.com/Oklahoma.**

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