

State of Oklahoma
Oklahoma Health Care Authority
Tykerb® (Lapatinib) Prior Authorization Form

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

Drug Information

Pharmacy billing (NDC: _____)
Dose: _____ Regimen: _____ Start Date: _____

Billing Provider Information

Provider NPI: _____ Provider Name: _____
Provider Phone: _____ Provider Fax: _____

Prescriber Information

Prescriber NPI: _____ Prescriber Name: _____
Prescriber Phone: _____ Prescriber Fax: _____ Specialty: _____

Criteria

For Initial Authorization (Initial approval will be for the duration of 6 months):

1. Diagnosis of metastatic or recurrent breast cancer? Yes ___ No ___
2. If answer is 'no' from previous question, please indicate diagnosis: _____
3. Please indicate requested information:
Yes ___ No ___ Positive expression of Human Epidermal Receptor Type 2 (HER2)?
Yes ___ No ___ Using in combination with one of the following: Herceptin® (trastuzumab),
Xeloda® (capecitabine), or an aromatase inhibitor, such as Aromasin®
(exemestane), Femara® (letrozole), or Arimidex® (anastrozole)?
4. Please provide regimen details of combination treatment: _____

Additional Information: _____

For Continued Authorization:

1. Does member have any evidence of progressive disease while on lapatinib?
Yes ___ No ___
 2. Has the member experienced any adverse drug reactions related to lapatinib therapy?
Yes ___ No ___
- If yes, please specify adverse reactions: _____

Additional Information: _____

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.

Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy
Pharmacy Management Consultants
Product Based Prior Authorization Unit

Fax: 1-800-224-4014
Phone: 1-800-522-0114 Option 4

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