

**Enhertu® (Fam-Trastuzumab Deruxtecan-nxki) Prior Authorization Form**

**Member Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Member ID#:** \_\_\_\_\_

**Drug Information**

**Physician billing (HCPCS code:** \_\_\_\_\_ **) Start Date (or date of next dose):** \_\_\_\_\_

**Dose:** \_\_\_\_\_ **Regimen:** \_\_\_\_\_

**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:**

1. Please indicate the diagnosis and information:

**Breast Cancer**

- A. Is diagnosis unresectable or metastatic breast cancer? Yes \_\_\_ No \_\_\_
- B. Is disease human epidermal growth factor receptor 2 (HER2)-positive? Yes \_\_\_ No \_\_\_
  - i. Has member received prior therapy in the metastatic, neoadjuvant, or adjuvant setting and developed disease recurrence during or within 6 months of completing therapy? Yes \_\_\_ No \_\_\_
  - ii. Has member received 1 or more prior anti-HER2-based regimens? Yes \_\_\_ No \_\_\_
- C. Is disease HER-2 low [immunohistochemistry (IHC) 1+ or IHC 2+/in situ hybridization (ISH)-]? Yes \_\_\_ No \_\_\_
  - i. Has member received prior chemotherapy in the metastatic setting or developed disease recurrence during or within 6 months of completing adjuvant chemotherapy? Yes \_\_\_ No \_\_\_

**Colorectal Cancer (CRC)**

- A. Is disease advanced or metastatic? Yes \_\_\_ No \_\_\_
- B. Has disease progressed on prior therapy? Yes \_\_\_ No \_\_\_
- C. Is disease HER2-amplified? Yes \_\_\_ No \_\_\_
- D. Is disease RAS and BRAF mutation negative? Yes \_\_\_ No \_\_\_
- E. Will Enhertu® be used as a single-agent? Yes \_\_\_ No \_\_\_

**Gastric or Gastroesophageal Junction (GEJ) Adenocarcinoma**

- A. Is disease locally advanced or metastatic? Yes \_\_\_ No \_\_\_
- B. Is disease HER2-positive? Yes \_\_\_ No \_\_\_
- C. Has member received at least 1 prior trastuzumab-based regimen? Yes \_\_\_ No \_\_\_

**Non-Small Cell Lung Cancer (NSCLC)**

- A. Is diagnosis unresectable or metastatic NSCLC? Yes \_\_\_ No \_\_\_
- B. Is disease HER2-positive? Yes \_\_\_ No \_\_\_
- C. Has member received prior systemic therapy? Yes \_\_\_ No \_\_\_

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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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**Enhertu® (Fam-Trastuzumab Deruxtecan-nxki) Prior Authorization Form**

**Member Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Member ID#:** \_\_\_\_\_

**Criteria**

**For Initial Authorization:** *(continued)*

1. Please indicate the diagnosis and information:

**Solid Tumor**

- A. Is diagnosis unresectable or metastatic human epidermal receptor type 2 (HER2)-positive immunohistochemistry (IHC) 3+ solid tumor? Yes \_\_\_ No \_\_\_
- B. Has member received prior systemic treatment with not satisfactory alternative treatment options? Yes \_\_\_ No \_\_\_

**If diagnosis is not listed above, please indicate diagnosis:** \_\_\_\_\_

Additional Information: \_\_\_\_\_

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**For Continued Authorization:**

- 1. Date of last dose: \_\_\_\_\_
- 2. Does member have any evidence of progressive disease while on Enhertu® therapy? Yes \_\_\_ No \_\_\_
- 3. Has member experienced any adverse drug reactions related to Enhertu® therapy? Yes \_\_\_ No \_\_\_

Additional Information: \_\_\_\_\_

\_\_\_\_\_  
\_\_\_\_\_

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**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 both pages of this form in full will result in processing delays.***

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