

Fax completed form to: (855) 840-1678

If this is an URGENT request, please call (800) 882-4462

(800.88.CIGNA)

## Herceptin, Herzuma, Kanjinti, Ogivri, Ontruzant, Trazimera

(trastuzumab)

PHYSICIAN INFORMATION PATIENT INFORMATION \* Physician Name: \*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all \* DEA, NPI or TIN: Specialty: asterisked (\*) items on this form are completed.\* \* Patient Name: Office Contact Person: \* Cigna ID: \* Date of Birth: Office Phone: \* Patient Street Address: Office Fax: Office Street Address: City: State: Zip: City: State: Zip: Patient Phone: Urgency: Standard Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function) Medication Requested: Herceptin 150mg Herzuma 150mg Herzuma 420mg Kanjinti 150mg Kanjinti 420mg Ogivri 150mg Ogivri 420mg Ontruzant 150mg Ontruzant 420mg Trazimera 150mg Trazimera 420mg Dose: Frequency of therapy: Duration of therapy: Will this medication be given concurrently with other agents? If yes, please specify: ICD10: What is your patient's current weight? Is this a new start or continuation of therapy with the requested medication? If patient has been taking samples, please pick new start. continuation of therapy new start Start date: (if requested medication is Herceptin, Herzuma, or Ontruzant) The covered alternatives are: Kanjinti (trastuzumab-anns) [may require prior authorization] Ogivri (trastuzumab-dkst) [may require prior authorization], and Trazimera (trastuzumab-gypp) [may require prior authorization]. For the alternatives tried, please include medication name and strength, date(s) taken and for how long, and what the documented results were of taking each medication, including any intolerances or adverse reactions your patient experienced. For the alternatives NOT tried, please provide details why your patient can't try that medication. (if requested medication is Herceptin, Herzuma, or Ontruzant) For Kanjinti (trastuzumab-anns), which of the following applies to your patient? Patient has not tried this medication. Patient tried this medication, but it didn't work or didn't work well enough. Patient tried this medication, but had an allergic or adverse reaction. ☐ Other (if allergic/adverse reaction to Kanjinti) Is there documentation that this reaction was due to a formulation difference in the inactive ingredients between the requested medication and Kanjinti (trastuzumab-anns) (for example, difference in dyes, fillers, preservatives)? 🗌 Yes 🗌 No (if requested medication is Herceptin, Herzuma, or Ontruzant) For Ogivri (trastuzumb-dkst), which of the following applies to your patient? Patient has not tried this medication. Patient tried this medication, but it didn't work or didn't work well enough. Patient tried this medication, but had an allergic or adverse reaction. ☐ Other (if allergic/adverse reaction to Ogivri) Is there documentation that this reaction was due to a formulation difference in the inactive ingredients between the requested medication and Ogivri (trastuzumab-dkst) (for example, difference in dyes, fillers, preservatives)? ☐ Yes ☐ No

<ul> <li>(if requested medication is Herceptin, Herzuma, or Ontruzant) For Trazimera (trastuzumab-qyyp), which of the following applies to your patient?</li> <li>Patient has not tried this medication.</li> <li>Patient tried this medication, but it didn't work or didn't work well enough.</li> <li>Patient tried this medication, but had an allergic or adverse reaction.</li> <li>Other</li> </ul>		
(if allergic/adverse reaction to Trazimera) Is there documentation that this reaction was due to a formulation difference in the inactive ingredients between the requested medication and Trazimera (trastuzumab-qyyp) (for example, difference in dyes, fillers, preservatives)?		
(if documentation that reaction due to formulation difference w/Kanjinti, Ogivri and/or Trazimera) Please provide details to support.		
Where will this medication be obtained?		
<ul> <li>Accredo Specialty Pharmacy**</li> <li>Prescriber's office stock (billing on a medical claim form)</li> <li>Other (please specify):</li> <li>Conter (please specify):</li> </ul>		
**Medication orders can be placed with Accredo via E-prescribe - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822   NCPDP 4436920), Fax 888.302.1028, or Verbal 866.759.1557		
Facility and/or doctor dispensing and administering medication:         Facility Name:       State:         Tax ID#:         Address (City, State, Zip Code):		
NOTE: Per some Cigna plans, infusion of medication MUST occur in the lowest cost, medically appropriate setting		
Is this infusion occurring in a facility affiliated with hospital outpatient setting?		
If yes- Is this patient a candidate for re-direction to an alternate setting (such as AIS, MDO, home) with assistance of a Specialty Care Option Case Manager?  Yes No (provide medical necessity rationale):		
Is your patient a candidate for home infusion?       Yes       No         Does the physician have an in-office infusion site?       Yes       No		
Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient?		
What is your patient's diagnosis?   biliary tract cancer   brain metastases   breast cancer   colorectal cancer (CRC)   endometrial carcinoma   gastric or gastroesophageal junction adenocarcinoma   leptomeningeal metastases from breast cancer   salivary gland tumors   other (please specify):		
Clinical Information         Does the patient have HER2-positive disease?         Yes		
(if CRC) Does the patient have the wild-type RAS gene (RAS-WT)?YesNo(if CRC) Does the patient have unresectable advanced or metastatic disease?YesNo(if CRC) Has the patient received other therapy for this diagnosis before requesting/using this medication?YesNo		
(if previously treated) Has the patient been treated with a human epidermal growth factor receptor-2 (HER2) inhibitor (like Enhertu, Nerlynx, Kadcyla, Perjeta, Tykerb, Vizimpro) for this diagnosis before starting therapy with this medication, one of its biosimilars (Herzuma, Kanjinti, Ogivri, Ontruzant, Trazimera) or Herceptin Hylecta? Notes: Please answer "no" if only switching from Herceptin to a biosim (Herzuma, Ogivri, Ontruzant, Kanjinti, or Trazimera) OR vice versa. Yes No		
(if previously treated) Has the patient previously been treated with an oxaliplatin-based therapy without irinotecan (Camptosar) for this diagnosis?		

(if no irinotecan therapy without oxaliplatin) Has the patient been treated with FOLFOXIRI (fluorouracil [Adrucil, 5FU] oxaliplatin, and irinotecan [Camptosar]) regimen for this diagnosis? (if no FOLFOXIRI) Has the patient previously been treated with a fluoropyrimidine (like capecitabine [Xeloda], floxuric [Adrucil, 5FU]) without irinotecan (Camptosar) or oxaliplatin for this diagnosis?	Yes 🗌 No 🗌	
(if gastric/GEJ adenocarcinoma) Does the patient have advanced or metastatic disease?	Yes 🗌 No 🗌	
(if gastric/GEJ adenocarcinoma) What is the patient's performance status (PS)? ☐ PS 0, 1 or 2 ☐ PS 3 or 4 ☐ Unknown		
(if biliary tract cancer or endometrial carcinoma) Does the patient have advanced or recurrent disease?	Yes 🗌 No 🗌	
(if endometrial carcinoma) Will the requested medication be taken in combination with carboplatin and paclitaxel (Ab	raxane)? Yes ∏ No ∏	
(if biliary tract cancer) Will the requested medication be taken in combination with pertuzumab (Perjeta) or tucatinib ( progression on or after system treatment?		
(if brain metastases) Is breast cancer the primary cancer?	Yes 🗌 No 🗌	
(if salivary gland tumors and requesting Herceptin) Does the patient have recurrent disease?	Yes 🗌 No 🗌	
(if salivary gland tumors and requesting Herceptin) Does the patient have distant metastases?	Yes 🗌 No 🗌	
(if salivary gland tumors and requesting Herceptin) What is the patient's performance status (PS)? ☐ PS 0, 1, 2 or 3 ☐ PS 4 ☐ Unknown		
Additional pertinent information (including disease stage, prior therapy, performance status, and names/doses/ad any agents to be used concurrently):	min schedule of	
Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.		
Prescriber Signature: Date: Da	inte in vour EUP	
Our standard response time for prescription drug coverage requests is 5 business days. If your request is urgent, it is important that		
you call us to expedite the request. View our Prescription Drug List and Coverage Policies online at cigna.com.		
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