

Member Name: {{MEMFIRST}} {{MEMLAST}} DOB: {{MEMBERDOB}} PA Number: {{PANUMBER}}



>{{PANUMCODE}}

## Tarceva

### 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 1-866-249-6155.** If you have questions regarding the prior authorization, please contact CVS Caremark at 1-866-814-5506. 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:** {{MEMFIRST}} {{MEMLAST}} **Date:** {{TODAY}}  
**Patient's ID:** {{MEMBERID}} **Patient's Date of Birth:** {{MEMBERDOB}}  
**Physician's Name:** {{PHYFIRST}} {{PHYLAST}}  
**Specialty:** \_\_\_\_\_, **NPI#:** \_\_\_\_\_  
**Physician Office Telephone:** {{PHYSICIANPHONE}} **Physician Office Fax:** {{PHYSICIANFAX}}  
**Request Initiated For:** {{DRUGNAME}}

- What is the patient's diagnosis?  
 Non-small cell lung cancer (including brain metastases from non-small cell lung cancer)  
 Pancreatic cancer  
 Chordoma  
 Renal cell carcinoma  
 Other \_\_\_\_\_
- What is the ICD-10 code? \_\_\_\_\_
- Is the patient currently receiving treatment with the requested medication?  
 Yes  No *If No, skip to #8*
- If the diagnosis is Non-small cell lung cancer (including brain metastases from non-small cell lung cancer), has the patient experienced either unacceptable toxicity or disease progression while on the current regimen?*  
 Yes, unacceptable toxicity  Yes, disease progression  No *No further questions*
- If the diagnosis is Pancreatic cancer, Chordoma, Renal cell carcinoma, is there evidence of unacceptable toxicity or disease progression while on the current regimen? If Yes, no further questions*  Yes  No
- Is the disease T790M negative?  Yes  No
- Has the patient experienced unacceptable toxicity or disease progression while on the current regimen?  
 Yes  No *No further questions*
- Will Tarceva be used as any of the following?  
 A single agent  
 In combination with ramucirumab or bevacizumab  
 In combination with gemcitabine  
 In combination with bevacizumab
- What is the clinical setting in which the requested drug will be used?  
 Recurrent  Advanced  Metastatic  Locally advanced  Unresectable  
 Relapsed  Stage IV

**Send completed form to: Case Review Unit, CVS Caremark Prior Authorization Fax: 1-866-249-6155**

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Tarceva [erlotinib] SGM - 8/2023.

CVS Caremark Prior Authorization • 1300 E. Campbell Road • Richardson, TX 75081

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*Complete the following section based on the patient's diagnosis, if applicable.*

Section A: Non-Small Cell Lung Cancer (Including Brain Metastases from Non-Small Cell Lung Cancer)

10. Does the patient have sensitizing epidermal growth factor receptor (EGFR) mutation-positive disease?

***ACTION REQUIRED: If Yes, attach EGFR mutation test results.***  Yes  No  Unknown

Section B: Renal Cell Carcinoma

11. Does the disease have non-clear cell histology?  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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