

Fax completed form to: 1-866-868-0858  
Questions, please call: 1-866-316-6049

**Important Information About Prescription Drug Coverage**

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To: \_\_\_\_\_ From: \_\_\_\_\_

Fax: \_\_\_\_\_ Pages: \_\_\_\_\_

Re: Sutent® (sutinib): Prior Authorization Request Form: Please respond.

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- Please complete the attached Sutent® (sutinib) Prior Authorization Request Form
- To expedite the review process please complete all requested fields.
- Completed forms should be faxed to: 1-866-868-0858. It is not necessary to fax this cover page.

**Please note:** By signing the attached form you are certifying that the treatment requested is medically necessary. No contraindications are present and precautions have been considered. You will be supervising the patient's treatment. Supporting documentation is available in the patient's record. Signature on this form attests to the fact that the corresponding ICD-9 code has been documented in the patients chart and submitted on a CMS- 1500 form.

**Information about this drug**

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**Sutent® (sutinib)**

Sutent® will be approved only for FDA approved uses. Sutent® should be used cautiously in patients with pre-existing cardiac histories as reductions in LVEF and/or elevation in blood pressure have been observed.

You can make an expedited request by indicating this at the top of the attached form. If you request an expedited review, and sign the attached form, you certify that applying the 72 hour standard review time frame may seriously jeopardize the life or health of the member or the member's ability to regain maximum function.

**Information on the attached form is protected health information and subject to all privacy and security regulations under HIPAA.**

# Prior Authorization Request Form

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## Patient Information

Name \_\_\_\_\_  
Member ID \_\_\_\_\_  
Medicare ID \_\_\_\_\_  
Date of birth \_\_\_\_\_ Sex: M / F  
Address \_\_\_\_\_  
City \_\_\_\_\_  
State \_\_\_\_\_ ZIP \_\_\_\_\_  
Phone \_\_\_\_\_  
Nursing home resident? YES / NO  
Home care patient? YES / NO

## Prescriber and Pharmacy Information

Name \_\_\_\_\_  
Specialty \_\_\_\_\_  
DEA \_\_\_\_\_  
NPI \_\_\_\_\_  
Address \_\_\_\_\_  
City \_\_\_\_\_  
State \_\_\_\_\_ ZIP \_\_\_\_\_  
Phone \_\_\_\_\_ Fax \_\_\_\_\_  
Pharmacy name \_\_\_\_\_  
NCPDP \_\_\_\_\_  
NPI \_\_\_\_\_  
Phone \_\_\_\_\_ Fax \_\_\_\_\_

## All items below this line are for Physician Use Only

### Information for Requested Drug

Strength: \_\_\_\_\_ Dosage form: \_\_\_\_\_ Qty per 30 days: \_\_\_\_\_ Drug is (circle one): Newly prescribed/Refill  
Directions: \_\_\_\_\_ Diagnosis: \_\_\_\_\_ ICD-9 Code: \_\_\_\_\_

Standard Reviews will be completed in under 72 hours. An expedited review is available if you certify that a standard review timeframe will seriously jeopardize the health of your patient. To request an expedited review, simply indicate this at the top of this page.

### Sutent® (sutinib) Criteria

- 1 Please indicate the patient's diagnosis. Select or specify the correct ICD-9 code. This question must be completed  
 first line treatment of advanced renal cell carcinoma (ICD-9: 189)  
 gastrointestinal stromal tumor (GIST) after disease progression on or intolerance to imatinib mesylate (Gleevec®) (ICD-9: 171.8)  
 Other (diagnosis and ICD-9 code): \_\_\_\_\_
  - 2 Has the patient previously failed treatment with chemotherapy, immunotherapy, and/or surgical intervention?  
 Yes  No  
 Please list therapeutic modalities previously attempted: \_\_\_\_\_
- Please note that Sutent® dose adjustments may be required in patients taking CYP3A4 inducers (i.e., phenytoin, carbamazepine, phenobarbital, St. John's Wort) or inhibitors (i.e., ketoconazole, clarithromycin, protease inhibitors, grapefruit juice).

Medical Justification: (Attach additional page if necessary): \_\_\_\_\_  
\_\_\_\_\_

Prescriber's signature: \_\_\_\_\_ Date: \_\_\_\_\_