



OptumRx has partnered with CoverMyMeds to receive prior authorization requests, saving you time and often delivering real-time determinations.

Visit [go.covermymeds.com/OptumRx](http://go.covermymeds.com/OptumRx) to begin using this free service.

Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

## Sutent<sup>®</sup> Prior Authorization Request Form (Page 1 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting <b>brand</b>			Directions for Use:		
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>					
Clinical Information (required)					
<b>Select the diagnosis below:</b>					
<input type="checkbox"/> Gastrointestinal stromal tumor (GIST)					
<input type="checkbox"/> Islet cell tumors					
<input type="checkbox"/> Pancreatic neuroendocrine tumors (pNET)					
<input type="checkbox"/> Renal cell carcinoma (RCC)					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
<b>Clinical Information:</b>					
Is this request for continuation of prior Sutent therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient used Sutent within the past 120 days? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select if Sutent is prescribed by or in consultation with one of the following specialists:					
<input type="checkbox"/> Oncologist					
<input type="checkbox"/> Neuro-oncologist					
<b>For gastrointestinal stromal tumor (GIST), also answer the following:</b>					
Has the patient had trial and failure, contraindication, or intolerance to imatinib mesylate (generic Gleevec)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient experienced disease progression on, or contraindication or intolerance to Gleevec (imatinib)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>For pancreatic neuroendocrine tumors (pNET), also answer the following:</b>					
Does the patient have progressive, well-differentiated pNET? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have unresectable disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have locally advanced disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have metastatic disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>For renal cell carcinoma (RCC), also answer the following:</b>					
Select if the following applies to the patient's disease:					
<input type="checkbox"/> Advanced					
<input type="checkbox"/> Metastatic (stage IV disease)					
<input type="checkbox"/> Medically or surgically unresectable tumor					
<input type="checkbox"/> Relapsed					
Will Sutent be used as adjuvant therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is the patient at high risk of recurrent RCC following nephrectomy? <input type="checkbox"/> Yes <input type="checkbox"/> No					

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

Office use only: Sutent\_CMS\_2019Jan-W



## Sutent<sup>®</sup> Prior Authorization Request Form (Page 2 of 2)

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**Quantity Limit:**

What is the quantity requested per DAY? \_\_\_\_\_

**What is the reason for exceeding the plan limitations?**

- Titration or loading-dose purposes
- Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)
- Requested strength/dose is not commercially available
- There is a medically necessary justification why the patient cannot use a higher commercially available strength to achieve the same dosage and remain within the same dosing frequency. **Please specify:** \_\_\_\_\_
- Other: \_\_\_\_\_

**Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?**

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Please note:

This request may be denied unless all required information is received.  
If the patient is not able to meet the above standard prior authorization requirements, please call 1-800-711-4555.  
For urgent or expedited requests please call 1-800-711-4555.  
This form may be used for non-urgent requests and faxed to 1-800-527-0531.