



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

Stivarga® Prior Authorization Request Form (Page 1 of 2)

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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 brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below: <input type="checkbox"/> Colorectal cancer <input type="checkbox"/> Gastrointestinal stromal tumor (GIST) <input type="checkbox"/> Hepatocellular carcinoma <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical Information: Is this request for continuation of prior Stivarga therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient used Stivarga within the past 120 days? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if Stivarga is prescribed by or in consultation with one of the following specialists: <input type="checkbox"/> Hepatologist <input type="checkbox"/> Gastroenterologist <input type="checkbox"/> Oncologist					
For colorectal cancer, also answer the following: Does the patient have advanced disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have metastatic disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if the patient has had trial and failure, contraindication, or intolerance to the following: <input type="checkbox"/> Anti-VEGF therapy (e.g. Avastin [bevacizumab]) <input type="checkbox"/> FOLFOXIRI (fluorouracil, leucovorin, oxaliplatin, and irinotecan) <input type="checkbox"/> Fluoropyrimidine-based chemotherapy <input type="checkbox"/> Irinotecan-based chemotherapy <input type="checkbox"/> Oxaliplatin-based chemotherapy Has the disease progressed through all available regimens? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a RAS mutation? <input type="checkbox"/> Yes <input type="checkbox"/> No If the patient is RAS wild-type (no RAS mutation), has the patient had trial and failure, contraindication, or intolerance to an anti-EGFR therapy [e.g., Vectibix (panitumumab), Erbitux (cetuximab)]? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For gastrointestinal stromal tumor (GIST), also answer the following: Select if the following applies to the patient's disease: <input type="checkbox"/> Locally advanced <input type="checkbox"/> Metastatic <input type="checkbox"/> Progressive <input type="checkbox"/> Unresectable Select if the patient has had trial and failure, contraindication, or intolerance to the following: <input type="checkbox"/> Brand Gleevec <input type="checkbox"/> Imatinib mesylate (generic Gleevec) <input type="checkbox"/> Sutent (sunitinib malate)					



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For hepatocellular carcinoma, also answer the following:

Has the patient had trial and failure or intolerance to Nexavar (sorafenib tosylate)? Yes No

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?

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.