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

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

Gleevec® (imatinib) Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)	Provider Information (required)
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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"/> Acute lymphoblastic leukemia (ALL) | <input type="checkbox"/> Dermatofibrosarcoma protuberans (DFSP) |
| <input type="checkbox"/> Aggressive systemic mastocytosis (ASM) | <input type="checkbox"/> Gastrointestinal stromal tumor (GIST) |
| <input type="checkbox"/> Chronic eosinophilic leukemia (CEL) | <input type="checkbox"/> Hypereosinophilic syndrome (HES) |
| <input type="checkbox"/> Chronic myelogenous/myeloid leukemia (CML) | <input type="checkbox"/> Myelodysplastic syndrome (MDS)/myeloproliferative disease (MPD) |
| <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____ | |

Clinical Information:

- Is the requested medication prescribed by or in consultation with a hematologist or oncologist? Yes No
- Is this a continuation of prior therapy? Yes No
- Has the patient used the requested medication in the past 120 days? Yes No
- For **brand Gleevec** requests: Does the patient have history of failure, contraindication, or intolerance to **generic imatinib**? Yes No

For acute lymphoblastic leukemia (ALL), also answer the following:

- Does the patient have Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ALL)? Yes No
- For patients < 18 years of age, is this a new diagnosis of Ph+ALL? Yes No

For aggressive systemic mastocytosis (ASM), also answer the following:

- Is the patient without the D816V c-Kit mutation or is the c-Kit mutational status unknown? Yes No

For chronic myelogenous/myeloid leukemia (CML), also answer the following:

- Does the patient have Philadelphia chromosome-positive chronic myelogenous/myeloid leukemia (Ph+CML)? Yes No
- Is the patient found to be Philadelphia chromosome positive or BCR-ABL positive as detected by bone marrow cytogenetics, FISH, or PCR? Yes No
- For patients < 18 years of age, is this a new diagnosis of Ph+CML in the chronic phase? Yes No

For dermatofibrosarcoma protuberans (DFSP), also answer the following:

- Does the patient have unresectable, recurrent, or metastatic disease? Yes No

For gastrointestinal stromal tumor (GIST), also answer the following:

- Does the patient have documented c-KIT (CD117) positive unresectable or metastatic malignant GIST? Yes No
- Has the patient had a resection of c-KIT (CD117) positive GIST? Yes No
- Will Gleevec (imatinib) be used as adjuvant therapy? Yes No

For myelodysplastic syndrome (MDS)/myeloproliferative disease (MPD), also answer the following:

- Is the disease associated with platelet-derived growth factor receptor (PDGFR) gene rearrangements? Yes 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: Gleevec-imatinib_CMS_2019Jan-W



Gleevec[®] (imatinib) Prior Authorization Request Form (Page 2 of 2)

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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.