

Bosulif
Prior Authorization Request

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

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.

Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____ **NPI#:** _____
Specialty: _____ **Physician Office Telephone:** _____ **Physician Office Fax:** _____
Request Initiated For: _____

1. What is the patient's diagnosis?
 Chronic myeloid leukemia (CML)
 Other _____
2. What is the ICD-10 code? _____
3. Prior to starting treatment for CML, was cytogenetic (conventional or FISH) and/or molecular testing (PCR) performed to detect the Philadelphia chromosome or the BCR-ABL gene? Yes No
4. Were the cells Philadelphia chromosome positive and/or BCR-ABL positive? **ACTION REQUIRED: Attach cytogenetic and/or molecular testing (documentation is NOT required for patients who have been previously approved for requested drug through CVS/caremark SGM prior authorization process).** Yes No
5. Has the patient received a hematopoietic stem cell transplant (HSCT) for CML?
If Yes, no further questions Yes No
6. Please indicate if any of the following apply.
Indicate ALL that apply or mark "None of the above" in the appropriate section.
 A) Received prior therapy with:
 imatinib (Gleevec) nilotinib (Tasigna) dasatinib (Sprycel) ponatinib (Iclusig)
 None of the above
 B) Experienced resistance to:
 imatinib (Gleevec) nilotinib (Tasigna) dasatinib (Sprycel) ponatinib (Iclusig)
 None of the above
 C) Experience toxicity or intolerance to:
 imatinib (Gleevec) nilotinib (Tasigna) dasatinib (Sprycel) ponatinib (Iclusig)
 None of the above
7. What is the CML phase? Chronic phase Accelerated phase Blast phase
8. Is the request for new start or continuation of Bosulif therapy?
 New start Continuation, *skip to #11 (if applicable)*
9. Was T315I mutation testing performed? Yes No

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. Bosulif SGM - 11/2016.

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10. Is the patient positive for T315I mutation? ***ACTION REQUIRED: Attach T315I mutation test result.***
 Yes No

Complete the following question if patient has chronic CML phase.

11. How long has the patient been receiving Bosulif? _____ months
12. *If the patient has received at least 12 months*, does the patient show evidence of disease progression?
 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)**