

Prescriber Criteria Form

Jakafi 2024 PA Fax 723-A v1 010124.docx
 Jakafi (ruxolitinib)
 Coverage Determination

This fax machine is located in a secure location as required by HIPAA regulations.
 Complete/review information, sign and date. Fax signed forms to CVS Caremark at **1-855-633-7673**.
 Please contact CVS Caremark at **1-866-785-5714** with questions regarding the prior authorization process.
 When conditions are met, we will authorize the coverage of Jakafi (ruxolitinib).

Drug Name:
 Jakafi (ruxolitinib)

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|----------------------------|------------------------|-------------|
| Patient Name: | | |
| Patient ID: | | |
| Patient DOB: | Patient Phone: | |
| Prescriber Name: | | |
| Prescriber Address: | | |
| City: | State: | Zip: |
| Prescriber Phone: | Prescriber Fax: | |
| Diagnosis: | ICD Code(s): | |

| Please circle the appropriate answer for each question. | | | |
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| 1 | Does the patient have a diagnosis of myelofibrosis (e.g., lower-risk, intermediate-risk, high-risk, primary, post-polycythemia vera, post-essential thrombocythemia, accelerated phase myelofibrosis, or blast phase myelofibrosis/acute myeloid leukemia)? [If yes, then no further questions.] | Yes | No |
| 2 | Does the patient have a diagnosis of polycythemia vera (PV)? [If no, then skip to question 4.] | Yes | No |
| 3 | Has the patient had an inadequate response or intolerance to any of the following: A) hydroxyurea, B) interferon therapy? [No further questions.] | Yes | No |
| 4 | Does the patient have a diagnosis of steroid-refractory acute graft-versus-host disease or chronic graft-versus-host disease? [If yes, then no further questions.] | Yes | No |
| 5 | Does the patient have a diagnosis of acute lymphoblastic leukemia (ALL)? [If no, then skip to question 7.] | Yes | No |
| 6 | Does the patient have a cytokine receptor-like factor 2 (CRLF2) mutation or a mutation associated with activation of the Janus kinase/signal transducers and activators of transcription (JAK/STAT) pathway? [No further questions.] | Yes | No |

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|----|--|-----|----|
| 7 | Does the patient have a diagnosis of chronic myelomonocytic leukemia (CMML)-2? [If no, then skip to question 9.] | Yes | No |
| 8 | Will the requested drug be used in combination with a hypomethylating agent? [No further questions.] | Yes | No |
| 9 | Does the patient have a diagnosis of myelodysplastic syndrome/myeloproliferative neoplasm (MDS/MPN) with neutrophilia? [If no, then skip to question 12.] | Yes | No |
| 10 | Will the requested drug be used as a single agent? [If yes, then no further questions.] | Yes | No |
| 11 | Will the requested drug be used in combination with a hypomethylating agent? [No further questions.] | Yes | No |
| 12 | Does the patient have a diagnosis of essential thrombocythemia? [If no, then skip to question 14.] | Yes | No |
| 13 | Has the patient had an inadequate response or loss of response to any of the following: A) hydroxyurea, B) interferon therapy, C) anagrelide? [No further questions.] | Yes | No |
| 14 | Does the patient have a diagnosis of myeloid, lymphoid, or mixed lineage neoplasms with eosinophilia and janus kinase 2 (JAK2) rearrangement? [If no, then no further questions.] | Yes | No |
| 15 | Is the disease in chronic or blast phase? | Yes | No |

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| Comments: | |
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By signing this form, I attest that the information provided is accurate and true as of this date and that the documentation supporting this information is available for review if requested by the health plan.

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| Prescriber (or Authorized) Signature: _____ Date: _____ |
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