

Market Applicability							
Market	DC	GA	KY	MD	NJ	NY	WA
Applicable	X	X	X	X	X	X	X

Jakafi (ruxolitinib)

Override(s)	Approval Duration
Prior Authorization Quantity Limit	1 year

Medications	Quantity Limit
Jakafi (ruxolitinib)	May be subject to quantity limit

APPROVAL CRITERIA

Requests for Jakafi (ruxolitinib) may be approved if the following are met:

- I. Individual has a diagnosis of low, intermediate, or high-risk myelofibrosis including any of the following (Label, NCCN 2A):
 - A. Primary myelofibrosis; **OR**
 - B. Post-polycythemia vera myelofibrosis; **OR**
 - C. Post-essential thrombocythemia myelofibrosis;
- OR**
- II. Individual has a diagnosis of polycythemia vera with an inadequate response to or intolerance to hydroxyurea or interferon therapy (Label, NCCN 2A);
- OR**
- III. Individual is 12 years and older; **AND**
 - IV. Individual has a diagnosis of steroid-refractory acute graft versus host disease (GVHD).

State Specific Mandates		
State name N/A	Date effective N/A	Mandate details (including specific bill if applicable) N/A

Key References:

This policy does not apply to health plans or member categories that do not have pharmacy benefits, nor does it apply to Medicare. Note that market specific restrictions or transition-of-care benefit limitations may apply.

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Applicable	X	X	X	X	X	X	X

1. DailyMed. Package inserts. U.S. National Library of Medicine, National Institutes of Health website. <http://dailymed.nlm.nih.gov/dailymed/about.cfm>. Accessed: June 4, 2019.
2. DrugPoints® System [electronic version]. Truven Health Analytics, Greenwood Village, CO. Updated periodically.
3. Lexi-Comp ONLINE™ with AHFS™, Hudson, Ohio: Lexi-Comp, Inc.; 2019; Updated periodically.
4. NCCN Clinical Practice Guidelines in Oncology™. © 2019 National Comprehensive Cancer Network, Inc. For additional information visit the NCCN website: <http://www.nccn.org/index.asp>. Accessed on June 4, 2019.
 - a. Myeloproliferative Neoplasms. V2.2019. Revised October 29, 2018.

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