

JAKAVI - Home - HCP

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## **JAKAVI® (ruxolitinib)**

JAKAVI® (ruxolitinib) is indicated for the treatment of disease-related splenomegaly or symptoms in adult patients with primary myelofibrosis (also known as chronic idiopathic myelofibrosis), post polycythaemia vera myelofibrosis or post essential thrombocythaemia myelofibrosis. JAKAVI® is also indicated for adult patients with polycythaemia vera who are resistant to or intolerant of hydroxyurea.<sup>1</sup>



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**JAKAVI® (ruxolitinib) in myelofibrosis**

**JAKAVI® (ruxolitinib) in myelofibrosis**

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**JAKAVI® (ruxolitinib) in polycythaemia vera**

**JAKAVI® (ruxolitinib) in polycythaemia vera**

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## Reference

1. JAKAVI® (ruxolitinib) Summary of Product Characteristics.

UK | November 2024 | FA-11208694

Adverse events should be reported. Reporting forms and information can be found at [www.mhra.gov.uk/yellowcard](http://www.mhra.gov.uk/yellowcard). Adverse events should also be reported to Novartis online through the pharmacovigilance intake (PVI) tool at [www.novartis.com/report](http://www.novartis.com/report), or alternatively email [medinfo.uk@novartis.com](mailto:medinfo.uk@novartis.com) or call 01276 698370.

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