

**APPLICANT** (stamp or sticker acceptable)      **PATIENT NHI:** .....      **REFERRER** Reg No: .....

Reg No: .....      First Names: .....      First Names: .....

Name: .....      Surname: .....      Surname: .....

Address: .....      DOB: .....      Address: .....

.....      Address: .....      .....

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Fax Number: .....      Fax Number: .....

**Ruxolitinib**

**Initial application**  
Applications only from a haematologist. Approvals valid for 12 months.  
**Prerequisites**(tick boxes where appropriate)

The patient has primary myelofibrosis or post-polycythemia vera myelofibrosis or post-essential thrombocythemia myelofibrosis  
**and**

A classification of risk of intermediate-2 or high-risk myelofibrosis according to either the International Prognostic Scoring System (IPSS), Dynamic International Prognostic Scoring System (DIPSS), or the Age-Adjusted DIPSS  
**or**

A classification of risk of intermediate-1 myelofibrosis according to either the International Prognostic Scoring System (IPSS), Dynamic International Prognostic Scoring System (DIPSS), or the Age-Adjusted DIPSS  
**and**  
 Patient has severe disease-related symptoms that are resistant, refractory or intolerant to available therapy

**and**  
 A maximum dose of 20 mg twice daily is to be given

**Renewal**  
Current approval Number (if known):.....  
Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 12 months.  
**Prerequisites**(tick boxes where appropriate)

The treatment remains appropriate and the patient is benefiting from treatment  
**and**  
 A maximum dose of 20 mg twice daily is to be given

I confirm the above details are correct and that in signing this form I understand I may be audited.

Signed: ..... Date: .....  
Post application to Ministry of Health, Private Bag 3015, Wanganui – email: [customerservice@health.govt.nz](mailto:customerservice@health.govt.nz)