

**State of Oklahoma**  
**SoonerCare**  
**Inrebic<sup>®</sup> (Fedratinib) Prior Authorization Form**

**Member Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Member ID#:** \_\_\_\_\_

**Drug Information**

**Pharmacy billing (NDC:** \_\_\_\_\_ **) Start Date (or date of next dose):** \_\_\_\_\_

**Dose:** \_\_\_\_\_ **Regimen:** \_\_\_\_\_

**Pharmacy Information**

**Pharmacy NPI:** \_\_\_\_\_ **Pharmacy Name:** \_\_\_\_\_

**Pharmacy Phone:** \_\_\_\_\_ **Pharmacy Fax:** \_\_\_\_\_

**Prescriber Information**

**Prescriber NPI:** \_\_\_\_\_ **Prescriber Name:** \_\_\_\_\_

**Prescriber Phone:** \_\_\_\_\_ **Prescriber Fax:** \_\_\_\_\_ **Specialty:** \_\_\_\_\_

**Criteria**

**For Initial Authorization:**

1. Please indicate the diagnosis and information:

**Myelofibrosis**

A. Is disease intermediate-2 or high-risk primary or secondary (post polycythemia vera or post-essential thrombocythemia)? Yes \_\_\_ No \_\_\_

**If answer is none of the above, please indicate diagnosis:** \_\_\_\_\_

Additional Information: \_\_\_\_\_

**For Continued Authorization:**

1. Date of last dose: \_\_\_\_\_

2. Does patient have any evidence of progressive disease while on fedratinib therapy? Yes \_\_\_ No \_\_\_

3. Has the member experienced any adverse drug reactions related to fedratinib therapy? Yes \_\_\_ No \_\_\_

*If yes, please specify adverse reactions:* \_\_\_\_\_

Additional Information: \_\_\_\_\_

**Prescriber Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

***I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge.***

*Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.*

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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