



## Commercial/Healthcare Exchange PA Criteria Effective: November 13, 2019

**Prior Authorization:** Inrebic

**Products Affected:** Inrebic (fedratinib) oral capsules

**Medication Description:** Inrebic is an oral kinase inhibitor with activity against both wild-type and mutated Janus-associated kinase 2 (JAK2) and FMS-like tyrosine kinase 3 (FLT3). Abnormal JAK2 activation is associated with myelofibrosis and polycythemia vera.

**Covered Uses:** Treatment of adult patients with intermediate-2 or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis

**Exclusion Criteria:** N/A

**Required Medical Information:** Diagnosis

**Age Restrictions:** 18 years of age and older

**Prescriber Restrictions:** Prescribed by, or in consultation with a hematologist/oncologist.

**Coverage Duration:** 3 years

**Other Criteria:**

- A. Patient has a diagnosis of intermediate-2 or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocythemia) myelofibrosis; **AND**
- B. Patient is not currently using Jakafi (ruxolitinib), or will discontinue prior to initiation of Inrebic

**References:**

1. Product Information: Inrebic<sup>(R)</sup> oral capsules, fedratinib oral capsules. Celgene Corporation (per FDA), Summit, NJ, 2019.

**Policy Revision history**

Rev #	Type of Change	Summary of Change	Sections Affected	Date
1	New Policy	New Policy	All	11/13/2019

Last Res. 11.13.19