

**INQOVI**  
**(decitabine and cedazuridine)**

## **Pre - PA Allowance**

None

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## **Prior-Approval Requirements**

**Age** 18 years of age or older

### **Diagnosis**

Patient must have the following:

1. Myelodysplastic syndromes (MDS), including:
  - a. De novo and secondary MDS
  - b. Chronic myelomonocytic leukemia (CMML)
  - c. Intermediate-1, intermediate-2, and high-risk International Prognostic Scoring System groups

**AND ALL** of the following:

- a. Prescriber agrees to monitor absolute neutrophil count (ANC) and platelets prior to initiating Inqovi and before each cycle and delay the next cycle resuming at the same or reduced dose as clinically indicated
- b. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Inqovi and for 6 months after the last dose
- c. Male patients with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Inqovi and for 3 months after the last dose

## **Prior - Approval Limits**

**Quantity** 15 tablets per 84 days

**Duration** 12 months

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## **Prior – Approval *Renewal* Requirements**

**Age** 18 years of age or older

### **Diagnosis**



**BlueCross  
BlueShield**

Federal Employee Program.

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Patient must have the following:

1. Myelodysplastic syndromes (MDS), including:
  - a. De novo and secondary MDS
  - b. Chronic myelomonocytic leukemia (CMML)
  - c. Intermediate-1, intermediate-2, and high-risk International Prognostic Scoring System groups

**AND ALL** of the following:

- a. **NO** disease progression or unacceptable toxicity
- b. Prescriber agrees to monitor absolute neutrophil count (ANC) and platelets before each cycle and delay the next cycle resuming at the same or reduced dose as clinically indicated
- c. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Inqovi and for 6 months after the last dose
- d. Male patients with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Inqovi and for 3 months after the last dose

**Prior - Approval *Renewal* Limits**

Same as above