

**PYRUKYND  
(mitapivat)**

**Pre - PA Allowance**

None

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**Prior-Approval Requirements**

**Age** 18 years of age and older

**Diagnosis**

Patient must have the following:

1. Hemolytic anemia with pyruvate kinase (PK) deficiency
  - a. Confirmed by a PK deficiency test OR a mutation in the PKLR gene

**AND ALL** of the following:

1. Patient has **ONE** of the following:
  - a. Hemoglobin  $\leq$  10 g/dL
  - b. Six or more RBC transfusion episodes in the last 52 weeks (1 year)
2. **NO** moderate to severe hepatic impairment (Child-Pugh Class B or C)
3. Prescriber agrees to dose the patient based on hemoglobin levels and transfusion requirements
4. Prescriber agrees to discontinue treatment with Pyrukynd if no clinical benefit is observed by 24 weeks
5. Prescriber agrees to taper the Pyrukynd dose when discontinuing therapy to reduce the risk of acute hemolysis

**Prior - Approval Limits**

**Quantity** 168 tablets per 84 days

**Duration** 6 months

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**Prior – Approval *Renewal* Requirements**

**Age** 18 years of age and older

**Diagnosis**

Patient must have the following:

1. Hemolytic anemia with pyruvate kinase (PK) deficiency



**BlueCross  
BlueShield**

Federal Employee Program.

## **PYRUKYND (mitapivat)**

**AND ALL** of the following:

1. Patient has clinical benefit from therapy as defined by **ONE** of the following:
  - a. Increase in hemoglobin level
  - b. Reduction in need for RBC transfusion
2. **NO** moderate to severe hepatic impairment (Child-Pugh Class B or C)
3. Prescriber agrees to dose the patient based on hemoglobin levels and transfusion requirements
4. Prescriber agrees to taper the Pyrukynd dose when discontinuing therapy to reduce the risk of acute hemolysis

### **Prior - Approval *Renewal* Limits**

**Quantity**      168 tablets per 84 days

**Duration**      12 months