

**REVUFORJ
(revumenib)**

Pre - PA Allowance

None

Prior-Approval Requirements

Age 1 year of age or older

Diagnosis

Patient must have the following:

1. Relapsed or refractory acute leukemia

AND ALL of the following:

- a. Presence of lysine methyltransferase 2A gene (KMT2A) translocation in bone marrow cells
- b. Prescriber agrees to correct electrolyte abnormalities, including hypokalemia and hypomagnesemia, prior to treatment
- c. Prescriber agrees to monitor for signs and symptoms of differentiation syndrome
- d. Prescriber agrees to monitor for QTc interval prolongation
- e. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Revuforj and for 4 months after the last dose
- f. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Revuforj and for 4 months after the last dose

Prior - Approval Limits

Quantity 540 mg per day

Duration 12 months

Prior – Approval *Renewal* Requirements

Age 1 year of age or older

Diagnosis

Patient must have the following:



**BlueCross
BlueShield**

Federal Employee Program.

**REVUFORJ
(revumenib)**

1. Relapsed or refractory acute leukemia

AND ALL of the following:

- a. **NO** disease progression or unacceptable toxicity
- b. Prescriber agrees to monitor for signs and symptoms of differentiation syndrome
- c. Prescriber agrees to monitor for QTc interval prolongation
- d. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Revuforj and for 4 months after the last dose
- e. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Revuforj and for 4 months after the last dose

Prior - Approval *Renewal* Limits

Same as above