

**TAZVERIK
(tazemetostat)**

Pre - PA Allowance

None

Prior-Approval Requirements

Diagnoses

The patient must have **ONE** of the following:

1. Metastatic or locally advanced epithelioid sarcoma
 - a. 16 years of age or older
 - b. **NOT** eligible for complete resection
2. Relapsed or refractory follicular lymphoma
 - a. 18 years of age or older
 - b. Patient must have **ONE** of the following:
 - i. Tumors are positive for an EZH2 mutation as detected by an FDA-approved test **AND** patient has received at least 2 prior systemic therapies
 - ii. Patient has no satisfactory alternative treatment options

AND ALL of the following:

- a. Prescriber agrees to monitor for the development of secondary malignancies
- b. Females of reproductive potential **only**: patient will be advised to use effective non-hormonal contraception during treatment with Tazverik and for 6 months after the final dose
- c. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Tazverik and for 3 months after the final dose

Prior - Approval Limits

Quantity 720 tablets per 90 days

Duration 12 months

Prior – Approval *Renewal* Requirements

Diagnoses



**BlueCross
BlueShield**

Federal Employee Program.

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1. Metastatic or locally advanced epithelioid sarcoma
 - a. 16 years of age or older
2. Relapsed or refractory follicular lymphoma
 - a. 18 years of age or older

AND ALL of the following:

- a. **NO** disease progression or unacceptable toxicity
- b. Prescriber agrees to monitor for the development of secondary malignancies
- c. Females of reproductive potential **only**: patient will be advised to use effective non-hormonal contraception during treatment with Tazverik and for 6 months after the final dose
- d. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Tazverik and for 3 months after the final dose

Prior - Approval *Renewal* Limits

Same as above