

XALKORI
(crizotinib)**Pre - PA Allowance**

None

Prior-Approval Requirements**Age** 18 years of age and older**Diagnoses**Patient must have **ONE** of the following:

1. Recurrent or metastatic non-small cell lung cancer (NSCLC)
 - a. 18 years of age or older
 - b. Patient must have **ONE** of the following:
 - i. Tumor is positive for ALK mutation as determined by an FDA-approved test
 - ii. Tumor is positive for ROS-1 mutation, as determined by an FDA-approved test
 - iii. Tumor has MET amplification or MET exon 14 skipping mutation
2. Inflammatory myofibroblastic tumor (IMT)
 - a. 18 years of age or older
 - b. Tumor is positive for ALK mutation
3. Unresectable, recurrent, or refractory inflammatory myofibroblastic tumor (IMT)
 - a. 1 year of age or older
 - b. Tumor is positive for ALK mutation
4. Relapsed or refractory, systemic anaplastic large cell lymphoma (ALCL)
 - a. 1 to 21 years of age
 - b. Tumor is positive for ALK mutation

AND ALL of the following for **ALL** indications:

1. Ophthalmology examination at baseline and periodically throughout treatment

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2. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Xalkori and for at least 45 days after the last dose
3. Males with female partners of reproductive potential **only**: patient will be advised to use condoms during treatment with Xalkori and for at least 90 days after the last dose

Prior - Approval Limits

Quantity 360 capsules per 90 days **OR**
720 oral pellets per 90 days

Duration 12 months

Prior – Approval *Renewal* Requirements

Age 18 years of age and older

Diagnoses

Patient must have **ONE** of the following:

1. Recurrent or metastatic non-small cell lung cancer (NSCLC)
 - a. 18 years of age or older
2. Inflammatory myofibroblastic tumor (IMT)
 - a. 18 years of age or older
3. Unresectable, recurrent, or refractory inflammatory myofibroblastic tumor (IMT)
 - a. 1 year of age or older
4. Relapsed or refractory, systemic anaplastic large cell lymphoma (ALCL)
 - a. 1 to 21 years of age

AND ALL of the following for **ALL** indications:

1. **NO** symptoms indicative of treatment-related pneumonitis
2. Ophthalmology examinations are done periodically throughout treatment
3. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Xalkori and for at least 45 days following the last dose



Federal Employee Program.

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4. Males with female partners of reproductive potential **only**: patient will be advised to use condoms during treatment with Xalkori and for at least 90 days after the last dose

Prior – Approval *Renewal* Limits

Same as above