

State of Oklahoma  
Oklahoma Health Care Authority  
**Lorbrena® (Lorlatinib) Prior Authorization Form**

Member Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Member ID#: \_\_\_\_\_

**Drug Information**

Pharmacy billing (NDC: \_\_\_\_\_) Start Date (or date of next dose): \_\_\_\_\_

Dose: \_\_\_\_\_ Regimen: \_\_\_\_\_

**Billing Provider Information**

Provider NPI: \_\_\_\_\_ Provider Name: \_\_\_\_\_

Provider Phone: \_\_\_\_\_ Provider Fax: \_\_\_\_\_

**Prescriber Information**

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_

Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

**Criteria**

**For Initial Authorization (Initial approval will be for the duration of 6 months):**

1. Diagnosis of non-small cell lung cancer (NSCLC)? Yes \_\_\_ No \_\_\_

A. If answer is 'yes' to question 1, please check all of the following that apply:

- Metastatic NSCLC
- Tumor expresses Anaplastic Lymphoma Kinase (ALK) translocation
- Lorlatinib will be used as a single-agent
- Lorlatinib will be used as second-line therapy following disease progression on alectinib or ceritinib
- Lorlatinib will be used as third-line or greater therapy following disease progression on crizotinib and one other ALK inhibitor (i.e., ceritinib or alectinib)

If answer is 'no' to question 1, please provide diagnosis: \_\_\_\_\_

Additional Information: \_\_\_\_\_

**For Continued Authorization:**

1. Date of last dose: \_\_\_\_\_

2. Does member have any evidence of progressive disease while on lorlatinib? Yes \_\_\_ No \_\_\_

3. Has the member experienced adverse drug reactions related to lorlatinib therapy? Yes \_\_\_ No \_\_\_

If yes, please specify adverse reactions: \_\_\_\_\_

Additional Information: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

**I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge.**

*Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.*

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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