

# Alunbrig® (Brigatinib) 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

Pharmacy NPI: \_\_\_\_\_ Pharmacy Name: \_\_\_\_\_

Pharmacy Phone: \_\_\_\_\_ Pharmacy 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. Please indicate diagnosis and information

**Non-small cell lung cancer (NSCLC)**

A. Is diagnosis metastatic NSCLC? Yes \_\_\_\_\_ No \_\_\_\_\_

B. Anaplastic lymphoma kinase (ALK) positivity? Yes \_\_\_\_\_ No \_\_\_\_\_

**If diagnosis is not listed, please provide diagnosis:** \_\_\_\_\_

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

\_\_\_\_\_

\_\_\_\_\_

**For Continued Authorization:**

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

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

3. Has the member experienced adverse drug reactions related to brigatinib 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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