

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 _____

C. Will brigatinib be used as a single agent? 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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