



Mektovi® (Binimetinib) Prior Authorization Form

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Pharmacy billing (NDC: _____)

Dose: _____ Regimen: _____ Start Date: _____

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:

1. Please indicate the diagnosis and information:

Unresectable or Metastatic Melanoma

- A. Does member have BRAF V600E or V600K mutation? Yes ___ No ___
- B. Will binimetinib be used in combination with encorafenib? Yes ___ No ___

Non-Small Cell Lung Cancer (NSCLC)

- A. Is diagnosis metastatic NSCLC? Yes ___ No ___
- B. Does member have BRAF V600E mutation? Yes ___ No ___
- C. Will binimetinib be used in combination with encorafenib? Yes ___ No ___

If answer is none of the above, please indicate diagnosis: _____

Additional Information: _____

For Continued Authorization:

1. Date of last dose: _____
 2. Does patient have any evidence of progressive disease while on binimetinib therapy? Yes ___ No ___
 3. Has the member experienced any adverse drug reactions related to binimetinib 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.

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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