

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

1. Diagnosis of unresectable or metastatic melanoma? Yes \_\_\_ No \_\_\_
  - A. If answer is 'yes' to question 1, please check all of the following that apply:
    - BRAF V600E or V600K mutation detected by an FDA-approved test
    - Wild-type BRAF melanoma
    - Used as first-line therapy in combination with vemurafenib
    - Used as second-line therapy or subsequent therapy with vemurafenib
      - i. If cobimetinib is being used as second-line therapy or subsequent therapy, please provide member's ECOG performance status (0-5): \_\_\_\_\_
2. If answer is 'no' to question 1, please provide diagnosis: \_\_\_\_\_

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

**For Continued Authorization:**

1. Does member have any evidence of progressive disease while on cobimetinib? Yes \_\_\_ No \_\_\_
2. Has member experienced any adverse drug reactions related to cobimetinib 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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