

### Vanflyta® (quizartinib) Prior Authorization Form

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

#### Drug Information

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

#### Pharmacy Information

Pharmacy NPI: \_\_\_\_\_ Pharmacy Name: \_\_\_\_\_  
Pharmacy Phone: \_\_\_\_\_ Pharmacy Fax: \_\_\_\_\_

#### Prescriber Information

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_  
Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

#### Criteria

##### For Initial Authorization:

1. Please indicate diagnosis and information:

**Acute Myeloid Leukemia (AML)**

A. Is AML newly diagnosed? Yes \_\_\_ No \_\_\_

B. Is disease positive for FLT3 internal tandem duplication (FLT3-ITD) as detected by an FDA-approved test? Yes \_\_\_ No \_\_\_

C. How will quizartinib be used? (*select one*)

In combination with standard anthracycline and cytarabine-based induction

In combination with standard cytarabine-based consolidation

As maintenance therapy following standard anthracycline and cytarabine-based induction and cytarabine-based consolidation

Other: \_\_\_\_\_

**If diagnosis is not listed above, please indicate diagnosis:** \_\_\_\_\_

Additional information: \_\_\_\_\_  
\_\_\_\_\_

##### For Continued Authorization:

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

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

3. Has member experienced adverse drug reactions related to quizartinib 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 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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