

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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