



**Vanflyta (quizartinib)  
Prior Authorization Request Form**



Caterpillar Prescription Drug Benefit  
Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME: \_\_\_\_\_ MEMBER'S FIRST NAME: \_\_\_\_\_

**Instructions:** Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

**URGENT**

MEMBER INFORMATION		
LAST NAME:	FIRST NAME:	
PHONE NUMBER:	DATE OF BIRTH:	
STREET ADDRESS:		
CITY:	STATE:	ZIP CODE:
PATIENT INSURANCE ID NUMBER:		

MALE  FEMALE HEIGHT (IN/CM): \_\_\_\_\_ WEIGHT (LB/KG): \_\_\_\_\_ ALLERGIES: \_\_\_\_\_

IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: [HTTPS://MAGELLANRX.COM/MEMBER/EXTERNAL/COMMERCIAL/COMMON/DOC/EN-US/PHI DISCLOSURE AUTHORIZATION.PDF](https://MAGELLANRX.COM/MEMBER/EXTERNAL/COMMERCIAL/COMMON/DOC/EN-US/PHI_DISCLOSURE_AUTHORIZATION.PDF)

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE): \_\_\_\_\_  
AUTHORIZED REPRESENTATIVE'S PHONE NUMBER: \_\_\_\_\_

PRESCRIBER INFORMATION		
LAST NAME:	FIRST NAME:	
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:	
NPI NUMBER:	DEA NUMBER:	
PHONE NUMBER:	FAX NUMBER:	
STREET ADDRESS:		
CITY:	STATE:	ZIP CODE:
REQUESTOR (if different than prescriber):	OFFICE CONTACT PERSON:	

MEDICATION OR MEDICAL DISPENSING INFORMATION			
MEDICATION NAME:			
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
<input type="checkbox"/> NEW THERAPY	<input type="checkbox"/> RENEWAL	IF RENEWAL: DATE THERAPY INITIATED:	
DURATION OF THERAPY (SPECIFIC DATES):			

*Continued on next page*





**Vanflyta (quizartinib)**  
**Prior Authorization Request Form**  
 Caterpillar Prescription Drug Benefit  
 Phone: 877-228-7909 Fax: 800-424-7640



**MEMBER'S LAST NAME:** \_\_\_\_\_ **MEMBER'S FIRST NAME:** \_\_\_\_\_

<b>1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION?</b> <input type="checkbox"/> YES (if yes, complete below) <input type="checkbox"/> NO		
<b>MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):</b>	<b>DURATION OF THERAPY (SPECIFY DATES):</b>	<b>RESPONSE/REASON FOR FAILURE/ALLERGY:</b>
<b>2. LIST DIAGNOSES:</b>		<b>ICD-10:</b>
<input type="checkbox"/> Acute Myeloid Leukemia(AML) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____		
<b>3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.</b>		
<p><b>Is patient going to be using drug in a clinical trial?</b> <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Is the prescriber is an oncologist/hematologist? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Is the patient newly diagnosed acute myeloid leukemia (AML)? <input type="checkbox"/> Yes <input type="checkbox"/> No <b>Please provide documentation.</b></p> <p>Is the patient FLT3 internal tandem duplication (ITD)-positive? <input type="checkbox"/> Yes <input type="checkbox"/> No <b>Please provide documentation.</b></p> <p>Is the patient receiving standard "7+3" induction chemotherapy regimen of cytarabine 100 mg/m<sup>2</sup> per day (or 200 mg/m<sup>2</sup> per day) by continuous intravenous infusion and anthracycline (daunorubicin 60 mg/m<sup>2</sup> per day or idarubicin 12 mg/m<sup>2</sup> per day) by intravenous infusion? <input type="checkbox"/> Yes <input type="checkbox"/> No <b>Please provide documentation.</b></p> <p>Does patient have an Eastern Cooperative Oncology Group (ECOG) performance status 0-2? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Will VANFLYTA(quizartinib) be used as maintenance monotherapy following allogeneic hematopoietic stem cell transplantation (HSCT)? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Does patient have a diagnosis of AML secondary to prior chemotherapy or radiotherapy for other neoplasms? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Has patient been previously treated for AML? <input type="checkbox"/> Yes <input type="checkbox"/> No <b>Please provide documentation.</b></p> <p>Has patient been previously treated for AML with one of the following? <input type="checkbox"/> Yes <input type="checkbox"/> No <b>Please provide documentation.</b></p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Leukapheresis;</li> <li><input type="checkbox"/> Treatment for hyperleukocytosis with hydroxyurea</li> <li><input type="checkbox"/> Cranial radiotherapy for central nervous system (CNS) leukostasis</li> <li><input type="checkbox"/> Prophylactic intrathecal chemotherapy</li> <li><input type="checkbox"/>Growth factor/cytokine support</li> </ul> <p>Has patient had prior treatment with Vanflyta(quizartinib) or other FLT3-ITD inhibitors? <input type="checkbox"/> Yes <input type="checkbox"/> No <b>Please provide documentation.</b></p>		





**Vanflyta (quizartinib)**  
**Prior Authorization Request Form**  
 Caterpillar Prescription Drug Benefit  
 Phone: 877-228-7909 Fax: 800-424-7640



**MEMBER'S LAST NAME:** \_\_\_\_\_ **MEMBER'S FIRST NAME:** \_\_\_\_\_

**Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?**

**Please note:** Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.

**ATTESTATION:** I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

**Prescriber Signature or Electronic I.D. Verification:** \_\_\_\_\_ **Date:** \_\_\_\_\_

**CONFIDENTIALITY NOTICE:** The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return FAX) and arrange for the return or destruction of these documents.

**FAX THIS FORM TO: 800-424-7640**

**MAIL REQUESTS TO: Magellan Rx Management Prior Authorization Program Attn:CP-4201**

P.O.Box 64811

St. Paul, MN 55164-0811

Phone: 877-228-7909

