

Copiktra (duvelisib)
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: [PRIMETHERAPEUTICS.COM/NOPP](https://www.primetherapeutics.com/nopp)

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.

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1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION? <input type="checkbox"/> YES (if yes, complete below) <input type="checkbox"/> NO		
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE): 	DURATION OF THERAPY (SPECIFY DATES): 	RESPONSE/REASON FOR FAILURE/ALLERGY:
2. LIST DIAGNOSES:		ICD-10:
<input type="checkbox"/> Relapsed or refractory chronic lymphocytic leukemia(CLL) <input type="checkbox"/> Small lymphocytic leukemia (SLL) <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____		
3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.		
Clinical Information: <u>Diagnosis of relapsed or refractory chronic lymphocytic leukemia(CLL) or small lymphocytic leukemia(SLL):</u> Has patient had at least two prior therapies? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>Please provide documentation with dates of service.</i> Has the patient had an autologous transplant within 6 months of starting Copiktra? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>Please provide documentation with dates of service.</i> Has the patient had and allogeneic transplant? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>Please provide documentation with dates of service.</i> Has the patient been previously treated with another P13K inhibitor such as Zydelig(idelalisib)? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient been previously treated with a Bruton's inhibitor such as Imbruvica (ibrutinib)? <input type="checkbox"/> Yes <input type="checkbox"/> No Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review? <hr/> <hr/>		
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		



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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: Prime Therapeutics Management Prior Authorization Program
Attn: CP - 4201
P.O. Box 64811
St. Paul, MN 55164-0811