Calquence (acalabrutinib) **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.

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

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:	
REQUESTER (if different than prescriber):	OFFICE CONTACT PERSON:	

MEDICAL DIODENI	
VIEDICAL DISPENS	SING INFORMATION

MEDICATION NAME:			
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
NEW THERAPY	RENEWAL IF F	ENEWAL: DATE THERAPY	INITIATED:
DURATION OF THERAPY (SPECIFIC DATES):			
Continued on next name			

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MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:			
1. HAS THE PATIENT TRIED ANY YES (if yes, complete below)	OTHER MEDICATIONS FOR THIS	CONDITION?		
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:		
2. LIST DIAGNOSES:		ICD-10:		
 Mantle cell lymphoma Chronic lymphocytic leukemia Small lymphocytic lymphoma(
Other diagnosis:	ICD-10 Code(s):			
3. REQUIRED CLINICAL INFORMATO SUPPORT A PRIOR AUTHORIZ	ATION: PLEASE PROVIDE ALL REL ZATION.	EVANT CLINICAL INFORMATION		
Is patient going to be using drug	in combination with a clinical trial?	P 🗌 Yes 🔲 No		
For diagnosis of mantle cell lymphoma: Has the patient ever been treated for mantle cell lymphoma? Has the patient had at least one prior therapy for mantle cell lymphoma?* Yes No *Please submit documentation.				
Has the patient been previously treated with another Bruton tyrosine kinase (BTK) inhibitor such as Imbruvica (ibrutinib)? u Yes u No				
Is patient ineligible for an autologous hematopoietic stem cell transplant(HSCT)? Yes No*Please submit documentation. 				
Will patient use Calquence(acalabrutinib) in combination with rituximab and bendamustine? □ Yes □ No * <i>Please submit documentation.</i>				
For diagnosis of chronic lymphocytic leukemia(CLL) or small lymphocytic lymphoma(SLL): Has patient received any prior systemic therapies for CLL /SLL? □ Yes □ No				
Will patient be using Calquence in combination with obinutuzumab(Gazyva©)? \Box Yes \Box No				
Has patient received at least one prior systemic therapy for CLL/SLL? Yes No *Please submit documentation. 				
Will patient be using Calquence as Monotherapy? Yes No 				
Renewal Request:				
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Is patient continuing to demonstrate a positive clinical response? Yes	No *Please submit
documentation.	

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: Prime Therapeutics Management Prior Authorization Program Attn: CP-4201 P.O. Box 64811 St. Paul, MN 55164-0811 Phone: 877-228-7909

