

Prior Authorization Request Form for
acalabrutinib (Calquence)



JOHNS HOPKINS
MEDICINE

JOHNS HOPKINS
HEALTHCARE

7231 Parkway Drive, Suite 100, Hanover, MD 21076

**FAX Completed Form and
Applicable Progress Notes to:
(410) 424-4037**

USFHP Pharmacy Prior Authorization Form

To be completed by Requesting provider	
Drug Name:	Strength:
Dosage/Frequency (SIG):	Duration of Therapy:

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step 1 Please complete patient and physician information (please print):

Patient Name: _____	Physician Name: _____
Address: _____	Address: _____
Sponsor ID #: _____	Phone #: _____
Date of Birth: _____	Secure Fax #: _____

Step 2 Please complete the clinical assessment:

1. Is the patient GREATER THAN or EQUAL to 18 years of age?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No STOP Coverage not approved
2. Is Calquence being prescribed by or in consultation with a hematologist or oncologist?	<input type="checkbox"/> Yes Proceed to question 3	<input type="checkbox"/> No STOP Coverage not approved
3. Does the patient have pathologically confirmed relapsed or refractory mantle cell lymphoma (MCL), with documentation of monoclonal B cells that have a chromosome translocation t(11;14)(q13;q32) and/or overexpress cyclin D1 that has a short response duration to prior therapy (less than median progression free survival [PFS])?	<input type="checkbox"/> Yes Proceed to question 11	<input type="checkbox"/> No Proceed to question 4
4. Does the patient have a diagnosis of relapsed refractory therapy for chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL)?	<input type="checkbox"/> Yes Proceed to question 5	<input type="checkbox"/> No Proceed to question 7
5. Does the patient have CLL/SLL with del(17p)/TP53 mutation?	<input type="checkbox"/> Yes Proceed to question 9	<input type="checkbox"/> No Proceed to question 6
6. Does the patient fit into any of the following categories? o Younger than 65 years of age o 65 years of age or older with significant comorbidities o Frail patient with significant comorbidities (not able to tolerate purine analogs)	<input type="checkbox"/> Yes Proceed to question 9	<input type="checkbox"/> No STOP Coverage not approved

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7. Please provide the diagnosis.	_____ Proceed to question 8	
8. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?	<input type="checkbox"/> Yes Proceed to question 11	<input type="checkbox"/> No STOP Coverage not approved
9. Does the patient have evidence of a BTK C481S mutation?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 10
10. Does the patient have prior ibrutinib-refractory disease?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 11
11. Does the patient have significant cardiovascular disease such as uncontrolled or symptomatic arrhythmias, congestive heart failure, or myocardial infarction within 6 months of screening, or any Class 3 or 4 cardiac disease as defined by the New York Heart Association Functional Classification, or corrected QT interval (QTc) GREATER THAN 480 msec?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 12
12. Will the patient be monitored for bleeding, infection, cardiac arrhythmias, and cytopenias?	<input type="checkbox"/> Yes Proceed to question 13	<input type="checkbox"/> No STOP Coverage not approved
13. What is the patient's age/gender?	<input type="checkbox"/> Male - Sign and date below <input type="checkbox"/> Female of childbearing potential - Proceed to question 14 <input type="checkbox"/> Female not of childbearing potential - Sign and date below	
14. Has the patient been advised of the risk of significant fetal harm?	<input type="checkbox"/> Yes Proceed to question 15	<input type="checkbox"/> No STOP Coverage not approved
15. Is the patient breastfeeding?	<input type="checkbox"/> Yes Proceed to question 16	<input type="checkbox"/> No Sign and date below
16. Has it been confirmed that patients will not breastfeed during treatment and for at least 2 weeks following cessation of treatment?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

**Step
3**

I certify the above is true to the best of my knowledge. Please sign and date:

Prescriber Signature

Date

[08 April 2020]

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For Internal Use Only	
<input type="checkbox"/> Approved:	Duration of Approval: ____month(s)
<input type="checkbox"/> Denied:	Authorized By:
<input type="checkbox"/> Incomplete/Other:	PA#:
Date Faxed to MD:	Date Decision Rendered: