

## Calquence® (acalabrutinib) 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 the diagnosis and information:

**Mantle Cell Lymphoma (MCL)**

A. Will acalabrutinib be used as a single agent? Yes \_\_\_ No \_\_\_

**Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL)**

A. Will acalabrutinib be used as a single agent? Yes \_\_\_ No \_\_\_

B. Will acalabrutinib be used in combination with obinutuzumab? Yes \_\_\_ No \_\_\_

**Other** \_\_\_\_\_

Additional Information: \_\_\_\_\_

#### For Continued Authorization:

1. Date of last dose: \_\_\_\_\_

2. Does member have any evidence of progressive disease while on acalabrutinib? Yes \_\_\_ No \_\_\_

3. Has the member experienced any adverse drug reactions related to acalabrutinib 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 do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.*

#### 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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