

Brukinsa® (Zanubrutinib) Prior Authorization Form

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Pharmacy billing (NDC: _____) Start Date (or date of next dose): _____

Dose: _____ Dosing Regimen: _____

Billing Provider 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. Has member received at least 1 prior therapy? Yes ___ No ___

Marginal Zone Lymphoma (MZL)

A. Has member received at least 1 prior anti-CD20 monoclonal antibody-based therapy?
Yes ___ No ___

Waldenström's Macroglobulinemia

A. Will Brukinsa® be used as primary therapy? Yes ___ No ___
B. Will Brukinsa® be used as subsequent treatment? Yes ___ No ___

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

Follicular Lymphoma (FL)

A. Will Brukinsa® be used as third line or subsequent therapy for no response, relapsed, or progressive disease? Yes ___ No ___
B. Will Brukinsa® be used in combination with obinutuzumab? Yes ___ No ___

If diagnosis is not listed above, please indicate diagnosis: _____

Additional Information: _____

For Continued Authorization:

1. Date of last dose: _____

2. Does member have any evidence of progressive disease while on zanubrutinib? Yes ___ No ___

3. Has the member experienced any adverse drug reactions related to zanubrutinib therapy? Yes ___ No ___

If yes, please specify adverse reactions: _____

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. 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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