

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)

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.

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Fax completed prior authorization request form to **888-601-8461** or submit Electronic Prior Authorization through CoverMyMeds[®] or SureScripts. All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at **AetnaBetterHealth.com/Oklahoma.**