

Member Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Member ID#: \_\_\_\_\_

**Drug Information**

Physician billing (HCPCS code: \_\_\_\_\_)  Pharmacy billing (NDC: \_\_\_\_\_)  
Start Date (or date of next dose): \_\_\_\_\_ Dose: \_\_\_\_\_ Regimen: \_\_\_\_\_

**Billing Provider Information**

Provider NPI: \_\_\_\_\_ Provider Name: \_\_\_\_\_

Provider Phone: \_\_\_\_\_ Provider Fax: \_\_\_\_\_

**Prescriber Information**

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_

Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

**Criteria**

**For Initial Authorization (Initial approval will be for the duration of 6 months):**

1. Will blinatumomab be used as a single-agent? Yes \_\_\_ No \_\_\_

2. Please indicate the diagnosis and information:

**Acute Lymphoblastic Leukemia (ALL)**

A. What is the Philadelphia chromosome status of the leukemia?

Philadelphia chromosome negative (Ph-) ALL

Philadelphia chromosome positive (Ph+) ALL

Unknown

B. Does the patient have relapsed or refractory disease? Yes \_\_\_ No \_\_\_

C. Has member previously failed two Tyrosine Kinase Inhibitors (TKIs)? Yes \_\_\_ No \_\_\_

i. If yes, please list previously failed TKIs: \_\_\_\_\_

D. Will blinatumomab be used as consolidation in patient without substantial comorbidity with persistent or late clearance minimal residual disease positive (MRD+) following a complete response to induction? Yes \_\_\_ No \_\_\_

**If answer is none of the 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 blinatumomab? Yes \_\_\_ No \_\_\_

3. Has the member experienced adverse drug reactions related to blinatumomab 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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