

# Blinicyto<sup>®</sup> (Blinatumomab) Prior Authorization Form

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

## Drug Information

Physician billing (HCPCS code: \_\_\_\_\_)  Pharmacy billing (NDC: \_\_\_\_\_)

Dose: \_\_\_\_\_ Regimen: \_\_\_\_\_ Start Date (or date of next dose): \_\_\_\_\_

## 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. Please indicate the diagnosis and information:

**Philadelphia Chromosome-Negative (Ph-) Acute Lymphoblastic Leukemia (ALL)**

A. How will blinatumomab be used?

- As consolidation therapy as a component of multiphase chemotherapy
- As consolidation in adolescents/young adults or adults younger than 65 years of age without substantial comorbidity with persistent or late clearance minimal residual disease positive (MRD+) following a complete response to induction.
- As maintenance therapy in combination with mercaptopurine, vincristine, methotrexate, and prednisone (POMP) as a component of maintenance.
- For management of relapsed/refractory Ph- ALL.
- Other: \_\_\_\_\_

**Philadelphia Chromosome-Positive (Ph+) Acute Lymphoblastic Leukemia (ALL)**

A. How will blinatumomab be used?

- In combination with with a tyrosine kinase inhibitor (TKI) as frontline consolidation if not a candidate for multiagent chemotherapy.
- With or without a TKI as consolidation in adolescents/young adults or adults younger than 65 years of age without substantial comorbidity with persistent or late clearance MRD+ following a complete response to induction.
- As maintenance therapy in combination with POMP as a component of maintenance therapy if refractory to TKIs.
- For management of relapsed/refractory Ph+ ALL after failure of 2 TKIs.
- Other: \_\_\_\_\_

**If answer is none of the above, please indicate diagnosis:** \_\_\_\_\_

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

## Criteria

### For Initial Authorization: (continued)

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: \_\_\_\_\_

DRAFT

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**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.*

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