

# OPTIMA HEALTH COMMUNITY CARE

## (MEDICAID)

### PHARMACY/MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

**Directions:** The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-844-348-3720**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **Incomplete form will delay authorization process.**

**Drug Requested:** **Besponsa<sup>®</sup>** (inotuzumab ozogamicin) **IV (J9999/C9028) (Medical)**

**DRUG INFORMATION:** Complete all information below or authorization process will be delayed.

**Drug Form/Strength/Month:** \_\_\_\_\_

**Dosing Schedule:** \_\_\_\_\_ **Length of Therapy:** \_\_\_\_\_

**Diagnosis:** \_\_\_\_\_ **ICD Code:** \_\_\_\_\_

**CLINICAL CRITERIA:** Check **ALL** items below for appropriate diagnosis. Authorization process will be delayed if boxes for diagnosis are **NOT** checked.

- Patient is age 18 years or older

**AND**

- Patient has a diagnosis of B-cell precursor acute lymphoblastic leukemia (ALL)

**AND**

• **Select one of the conditions that corresponds to the patient:**

- Patient shown to be Philadelphia Chromosome-positive, and is either relapsed OR refractory CD22 as defined in either condition below

- a. Patient has undergone treatment with at least one tyrosine kinase inhibitor {Imatinib (Gleevec<sup>®</sup>), Dasatinib (Sprycel<sup>®</sup>), Nilotinib (Tasigna<sup>®</sup>), Bosutinib (Bosulif<sup>®</sup>), Ponatinib (Iclusig<sup>®</sup>)}
- b. Patient has undergone 1 or 2 induction chemotherapy regimens for ALL

**OR**

- Patient shown to be Philadelphia Chromosome-negative and:
- a. Patient has undergone 1 or 2 induction chemotherapy regimens for ALL

**AND**

**Select below the therapy regimen/cycle phase for approval:**

- Cycle 1: 21 DAYS**

**DAY 1 - 0.8 mg/m<sup>2</sup>**

**DAY 8 - 0.5 mg/m<sup>2</sup>**

**DAY 15 - 0.5 mg/m<sup>2</sup>**

**total dose/cycle 1: 1.8 mg/m<sup>2</sup>**

- ❖ treatment cycle may be extended to 4 weeks if complete remission (CR) is achieved, or CR with incomplete hematologic recovery (CRi) and/or to allow for recovery from toxicity.

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**Subsequent cycles:**

- Patients who achieve CR or CRi: 28 DAYS

<b>DAY 1 - 0.5 mg/m<sup>2</sup></b>	<b>DAY 8 - 0.5 mg/m<sup>2</sup></b>	<b>DAY 15 - 0.5 mg/m<sup>2</sup></b>
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**total dose/cycle: 1.5 mg/m<sup>2</sup>**

- Patients who do NOT achieve CR or CRi: 28 DAYS

<b>DAY 1 - 0.8 mg/m<sup>2</sup></b>	<b>DAY 8 - 0.5 mg/m<sup>2</sup></b>	<b>DAY 15 - 0.5 mg/m<sup>2</sup></b>
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**total dose/cycle 1: 1.8 mg/m<sup>2</sup>**

- ❖ if CR or CRi is not achieved within 3 cycles, discontinue treatment.

**Medication being provided by (check box below that applies):**

- Location/site of drug administration: \_\_\_\_\_  
NPI or DEA # of administering location: \_\_\_\_\_

**OR**

- Specialty Pharmacy - PropriumRx

*\*Use of samples to initiate therapy **does not** meet step-edit/preauthorization criteria.\**

*\***Previous therapies will be verified through pharmacy paid claims or submitted chart notes.**\**

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

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

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

\*Approved by the Pharmacy and Therapeutic Committee: 2/15/2018  
UPDATED/REVISED: 6/19/2018; 8/16/2018; 10/8/2018; (Reformatted) 2/4/2019