

# OPTIMA HEALTH PLAN

## 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 1-844-668-1550**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If information provided is not complete, correct, or legible, authorization can be delayed.**

**For Medicare Members:** Medicare Coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD) and Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. They can be found at: <https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx>. Additional indications may be covered at the discretion of the health plan.

**Drug Requested:** **Zevalin<sup>®</sup>** (ibritumomab tiuxetan, Yttrium Y-90) **IV (A9543)**

**DRUG INFORMATION:** Authorization may be delayed if incomplete.

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

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

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

**Weight:** \_\_\_\_\_

- Standard Review. In checking this box, the timeframe does not jeopardize the life or health of the member or the member's ability to regain maximum function and would not subject the member to severe pain.

### **Quantity Limits:**

#### **A. Length of Authorization**

- Coverage will be provided for one administration and may **NOT** be renewed

#### **B. Max Units (per dose and over time) [HCPCS Unit]:**

- 1 billable unit [up to 40 millicuries; 1 billable unit = 40 mCi]

**CLINICAL CRITERIA:** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

### **Approval Criteria – Coverage cannot be renewed**

- Member is at least 18 years of age
- Requesting provider is an oncologist
- For female members of reproductive potential**, a negative pregnancy test has been confirmed
- Pre-treatment measurement of member's platelet level has been provided, and is **NOT** < 100,000 cells/mm<sup>3</sup>
- Member has adequate marrow cellularity of > 15%
- Member's medical condition has < 25% involvement of lymphoma in bone marrow
- Requested medication will be used as a single agent following two doses of rituximab

(Continued on next page)

- Member must meet **ONE** of the following:
  - Member has a diagnosis of relapsed or refractory, low-grade or follicular B-cell non-Hodgkin lymphoma (NHL), **AND** has not been previously treated with ibritumomab
  - Member has a diagnosis follicular NHL, previously untreated, **AND** has achieved a partial or complete response to first-line chemotherapy
- Provider will follow the recommended dosage per weight and timeline indication detailed in the table below:

Indication	Dose
All Indications	<ul style="list-style-type: none"> <li>• Administer rituximab 250 mg/m<sup>2</sup> Day 1; repeat dose on Day 7, 8, or 9</li> <li>• Within 4 hours of the second dose of rituximab, administer ibritumomab intravenously as follows:               <ul style="list-style-type: none"> <li>○ Normal platelet count: 0.4 mCi/kg (14.8 MBq/kg)</li> <li>○ Relapsed/refractory &amp; platelets 100,000 – 149,000/mcL: 0.3 mCi/kg (11.1 MBq/kg)</li> </ul> </li> <li>• Do not exceed the maximum dose of 32.0 mCi (11.84 MBq)</li> </ul>

**Reauthorization Criteria – Coverage cannot be renewed**

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

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

**OR**

- Specialty Pharmacy - PropriumRx**

For urgent reviews: Practitioner should call Optima Pre-Authorization Department if they believe a standard review would subject the member to adverse health consequences. Optima’s definition of urgent is a lack of treatment that could seriously jeopardize the life or health of the member or the member’s ability to regain maximum function.

(Continued on next page; signature page must be attached to this request form)

(Please ensure signature page is attached to form.)

***\*\*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.\****

Member Name: \_\_\_\_\_

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

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

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

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

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

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

\*Approved by Pharmacy and Therapeutics Committee: 7/21/2022  
REVISED/UPDATED: 8/10/2022.