

# 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:** Xofigo® (radium Ra 223 dichloride) IV A9606

**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 six months (6 injections only) and may **NOT** be renewed

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

- 178 billable units every 28 days
- 1 billable unit = 1 microcurie
- Xofigo (radium Ra 223 dichloride injection) is supplied in single-use vials containing 6 mL of solution at a concentration of 1,100 kBq/mL (30 microcurie/mL) with a total radioactivity of 6,600 kBq/vial (178 microcurie/vial)

**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
- ☐ Member has a diagnosis of prostate cancer that is castration-resistant
- ☐ Member has symptomatic bone metastases, and will be used in conjunction with denosumab or zoledronic acid

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- ☐ Member does **NOT** have any known visceral metastatic disease
- ☐ Medication will be used as a single agent
- ☐ Provider will follow the recommended dosage per weight and timeline indication detailed in the table below:

Indication	Dose
Prostate Cancer	<ul style="list-style-type: none"><li>55 kBq (1.49 microcurie) per kg body weight, given at 4 week intervals for 6 injections.</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.

***\*\*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