

## Radium-223 Dichloride - Castration-Resistant Prostate Cancer

(This form must be completed before the first dose is dispensed.)

### 1. Patient Profile

- \* Surname: .....
- \* Given Name: .....
- \* OHIN: ..... \* Chart Number: .....
- \* Postal Code: .....
- \* Height (cm): ..... \* Weight (kg): .....
- \* BSA (m<sup>2</sup>): ..... \* Gender:  Male  Female  Other
- \* Date of Birth: .....  
Day    Month    Year
- \* Site: .....
- \* Attending Physician (MRP- Most Responsible Physician): .....
- Requested Prior Approval  Yes \* Patient on Clinical Trial  Yes  No
- Other (specify): .....
- Specify Arm:  
 Standard of care arm  Experimental arm  
 Blinded / Unknown

### Prior Approval Request

- \* Select the appropriate  
prior approval  
scenario:

- 1-Unknown primary (submit pathology report and clinic note)
- 2-Clinical document review (identify the patient history that needs to be reviewed against eligibility criteria in Additional Comments below)
- 3-Regimen modification - schedule (complete questions a and b)
- 4-Regimen modification - drug substitutions (complete questions a and c)
- 5-Withholding a drug in combination therapy from start of treatment (complete questions d, e and f)
- 6-Maintenance therapy delay (submit clinic note)
- 7-Prior systemic therapy clinical trials (complete question g)
- 8-Modification due to supply interruption/drug shortage
- Other (specify)

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**All relevant supporting documentation must be submitted at the time of prior approval. Documentation may include a pathology report, clinic note, and/or CT scans.**

a. Co-morbidities / toxicity / justification:

.....

b. Intended regimen schedule: .....

c. Intended regimen: .....

d. Drug(s) to be held: .....

e. Rationale for holding drug(s): .....

f. Intention to introduce drug at a later date?  Yes

g. Prior clinical trial identifier (e.g., NCT ID, trial name) and treatment description (e.g., arm, drug/regimen): .....

h. Anticipated date of first treatment: \_\_\_\_\_  
Day      Month      Year

i. Additional comments:

.....

## 2. Eligibility Criteria

The patient must meet the following criteria:

- Patient has castration-resistant prostate cancer (CRPC) with symptomatic bone metastases and no known visceral metastatic disease.  Yes

## 3. Funded Dose

Please select if radium is administered pre- or post-docetaxel for metastatic castration resistant prostate cancer:  Pre-Docetaxel\*  
 Post-Docetaxel

- **\*If used in the pre-docetaxel setting, no subsequent funding will be considered in the post-docetaxel setting.**
- **Effective as of April 18, 2016:** The funded dose regimen is 55 kBq (1.49 microcurie) per kg body weight, given at 4-week intervals for a total of 6 injections.

## 4. Notes

Please check the following to confirm and acknowledge that:

- a consult with a medical or radiation oncologist has been done before starting radium therapy  Yes
- this enrolment will not be combined with cabazitaxel or enzalutamide or abiraterone for mCRPC  Yes
- if radium is funded in the pre-docetaxel setting, no subsequent funding will be considered in the post-docetaxel setting  Yes

## 5. Supporting Documents

- None required for this policy.

**In the absence of collecting supporting documentation:**

- CCO reserves the right to perform an audit on the patient's eligibility to receive reimbursement for this policy.
- In the event of an audit, CCO may request the following:
  - Consultation note or letter with medical or radiation oncologist documenting the radium treatment decision

Signature of Attending Physician (MRP-Most Responsible Physician): .....

.....  
Day    Month    Year

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