Eligibility Form

Cabazitaxel - Metastatic Castration Resistant Prostate Cancer

(This form must be completed <u>before</u> the first dose is dispensed.)

| 1. Patient Profile | | | | | | | |
|---|---|--|--|--|--|--|--|
| * Surname: | | | | | | | |
| * Given Name: | | | | | | | |
| * OHIN: | * Chart Number: | | | | | | |
| * Postal Code: | | | | | | | |
| * Height (cm): | * Weight (kg): | | | | | | |
| * BSA (m ²): | * Gender: O Male O Female O Other | | | | | | |
| * Date of Birth: | Day Month Year | | | | | | |
| * Site: | | | | | | | |
| * Attending Physician | (MRP- Most Responsible Physician): | | | | | | |
| Requested Prior Ap | proval 🗌 Yes \star Patient on Clinical Trial 🔘 Yes 💮 No | | | | | | |
| Other (specify): | | | | | | | |
| Specify Arm: Standard of care Blinded / Unkno | • | | | | | | |
| Prior Approval R | Request | | | | | | |
| * Select the appropriate prior approval scenario: | 1-Unknown primary (submit pathology report ○ 2-Clinical document review (identify the patient and clinic note) history that needs to be reviewed against eligibility criteria in Additional Comments below) 3-Regimen modification - schedule (complete ○ 4-Regimen modification - drug substitutions questions a and b) (complete questions a and c) 5-Withholding a drug in combination therapy ○ 6-Maintenance therapy delay (submit clinic note) from start of treatment (complete questions d, e and f) 7-Prior systemic therapy clinical trials (comple ○ 8-Modification due to supply interruption/drug question g) 9-COVID-19 pandemic: use of chemotherapy ○ Other (specify) after sequential androgen receptor axis-targeted agents (ARATs) | | | | | | |

| pa | thology report, | clinic no | te, and/o | r CT scans |
|--|--|-------------|------------|------------|
| a. Co-m | norbidities / toxid | city / just | ification: | |
| | | | | |
| b. Inten | ided regimen dule: | | | |
| c. Inten | ided regimen: | | | |
| d. Drug | (s) to be held: | | | |
| | onale for ng drug(s): | | | |
| | ntion to duce drug at a date? | ☐ Yes | | |
| ident NCT name treati desc arm, | clinical trial ifier (e.g., ID, trial e) and ment ription (e.g., | | | |
| h. Anticipated date of first treatment: | | | | |
| | | Day | Month | Year |
| i. Addit | tional comments | S: | | |
| | | | | |
| | | | | |
| | | | | |

All relevant supporting documentation must be submitted at the time of prior approval. Documentation may include a

The patient must meet the following criteria:

2. Eligibility Criteria

| Cabazitaxel will be used in combination with prednisone for the treatment of metastatic castration-resistant | | | | | | | | | | | |
|--|------------|-------------|---------------------------|-------|--|--|--|--|--|--|--|
| 3. Funded Dose | | | | | | | | | | | |
| Cabazitaxel 20 mg/m ² or 25 mg/m ² IV every 3 weeks (with 10 mg oral | prednis | one daily | v) until disease progress | sion. | | | | | | | |
| 4. Notes | | | | | | | | | | | |
| 1. Cabazitaxel is not funded if used in combination with abiraterone, or e | nzalutan | nide, or ra | adium-223 for mCRPC | | | | | | | | |
| Cabazitaxel is funded in the mCRPC setting in patients who have prog chemotherapy and an androgen-receptor-axis-targeted agent (ARAT), setting(s). | | | | • | | | | | | | |
| 5. Supporting Documents | | | | | | | | | | | |
| None required at time of enrolment. | | | | | | | | | | | |
| In the event of an audit, the following should be available to document • Clinic note documenting treatment history. | eligibilit | y: | | _ | | | | | | | |
| Signature of Attending Physician (MRP- Most Responsible Physician): | <u></u> | | | | | | | | | | |
| | Day | Month | Year | | | | | | | | |
| Form 992 | | | | | | | | | | | |