

## Nab-PACLitaxel - Metastatic Breast 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 criteria a, b OR c, and d:

- a. The patient has metastatic breast cancer.  Yes
- b. Has had acute infusion reactions with paclitaxel or docetaxel considered by treating physicians to be due to the vehicle of the taxanes (Cremophor and polysorbate 80)  Yes
- c. Has experienced severe toxicity from previous administration of other taxanes (Severe toxicity could be due to pre-medications for the administration of the taxane or due to the taxane itself) <sup>a</sup>  Yes
- d. Please specify previous taxane:     Docetaxel                       Paclitaxel

## 4. Notes

<sup>a</sup>. excludes glycemic effects of steroids

- Nab-paclitaxel may be used in place of paclitaxel or docetaxel provided that the patient meets nab-paclitaxel eligibility criteria. The cost of paclitaxel or docetaxel is funded through the Systemic Treatment Quality-Based Procedure (ST-QBP) and is included in the band level pricing.

## 5. Supporting Documents

To ensure reimbursement of your claim, both the completed enrolment form and a copy of the required documentation (where applicable) must be submitted through CCO e-Claims.

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

.....  
Day    Month    Year