

# Liposomal Doxorubicin - Platinum - Resistant Ovarian, Fallopian Tube, or Primary Peritoneal 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:

a. Patient has previously been treated with platinum-containing chemotherapy:

- with Paclitaxel  
 without Paclitaxel

Please specify: \_\_\_\_\_

b. Please select one of the following:

- disease has relapsed less than 6 months following therapy  
 tumour has progressed during therapy or not responding to therapy

c. Patient has reasonable performance status with symptoms that are likely to be alleviated if response is achieved  Yes

## 4. Funded Dose

- Liposomal doxorubicin 50 mg/m<sup>2</sup> every 4 weeks
- Liposomal doxorubicin 40 mg/m<sup>2</sup> every 4 weeks (if used with bevacizumab 10 mg/kg every 2 weeks)

## 5. Notes

1. Patients with primary platinum refractory disease (i.e., disease that has progressed while on front-line platinum-based chemotherapy) are not eligible for bevacizumab in the platinum-resistant setting.
2. Liposomal doxorubicin is only funded once (i.e., as one line of therapy, either as a single agent or as part of a combination regimen) for the treatment of recurrent ovarian, fallopian tube, or primary peritoneal cancer.

## 6. 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