MEDAVIE BLUE CROSS™

PRIOR AUTHORIZATION REQUEST

PROCEDURES FOR PRIOR AUTHORIZATION

Completed forms can be faxed in confidence to 1-514-286-8480 for residents of Quebec and 1-844-661-2640 for residents of all other provinces

Upon receipt, this request will be confidentially reviewed according to payment criteria developed by Medavie Blue Cross in consultation with independent health care consultants. In some cases, additional clinical or diagnostic information may be required to process your claim.

For Quebec residents, the criteria for prior authorization are adjusted to meet the requirements of the Régie de l'assurance maladie du Québec (RAMQ).

- Prior Authorization is a pre-approval process to determine if certain products will be reimbursed under a member's benefit plan.
- Please complete entire form. Incomplete forms cannot be processed.
- **For certain medications, approval for reimbursement may be conditional on confirmation of enrollment in the patient support program.**
- Prior Authorization may be limited to a specified period or quantity of medication.
  Some Medavie Blue Cross plans may require you to purchase a drug requiring prior authorization from a preferred pharmacy*. If your prior authorization request is approved, a case manager may contact you, your physician, or Patient Assistance Program to provide information about the program and to arrange to have your prescription transferred to the preferred pharmacy.
  **Not applicable in Quebec*.
- In cases where a request for Prior Authorization is declined, Medavie Blue Cross is denying payment for a product and is not challenging the medical opinion of the physician nor rendering a medical opinion.
- Any costs associated with the completion of this form or obtainment of additional medical information are the responsibility of the member.
- Renewal of the Prior Authorization will be considered by Medavie Blue Cross upon request from the patient/member. The renewal request should include information from the physician supporting continued use of the medication.
- Prior Authorization coverage is contingent on your continued status as a Medavie Blue Cross cardholder or beneficiary.
- If this is a request under the *Mesure du patient d'exception* for a Quebec resident, please include a completed *Patient d'exception* form that can be found here: www.medaviebc.ca/en/resources, in addition to this document.
- If you would like more information about our Patient First Network, including how your Patient Support Program can become integrated with our new enhanced Prior Authorization processes, please send an e-mail to: patientfirstnetwork@medavie.bluecross.ca.

Residents of All Other Provinces
PO BOX 220, MONCTON (NB) E1C 8L3
TEL.: 1-800-667-4511 FAX: 1-844-661-2640

Residents of Quebec
PO BOX 3300, STATION B, MONTREAL (QC) H3B 4Y5
TEL.: 1-888-588-1212 FAX: 1-514-286-8480

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1 PHARMACY INFORMATION

This section is to be completed by the Professional coordinating the request on behalf of the member (PSP, Cancer Care Navigator or Pharmacy)

Decision communication preference: ☐ Fax, Number: _____ ☐ Telephone, Number: _____

Name of Program/Pharmacy: _____

Contact Name: _____ Contact E-mail: _____

2 PATIENT INFORMATION

Part A

Patient Name: _____ Date of Birth: _____ (mm/dd/yyyy)

E-mail address of patient (or of legal guardian if patient is underage): _____

Address: _____ Suite: _____ City: _____

Province: _____ Postal Code: _____ Telephone Number: _____

Policy Number: _____ ID Number: _____

Do you have valid Medicare coverage in your current province of residence? ☐ Yes ☐ No

Have you already purchased this prescription? ☐ Yes ☐ No

Please attach your paid-in-full receipt with this request form. If you have already submitted your receipt to Medavie Blue Cross, please indicate the date of the oldest receipt. Date: _____ (mm/dd/yyyy)

Part B – Coordination of Benefits

Do you or any dependant have coverage for this drug under any other plan or program? ☐ Yes ☐ No **If Yes, complete the following:**

Policy Number: _____ Carrier: _____

(If applicable, please attach Explanation of Benefits from prior carrier with complete form)

If the patient is a dependent, provide the birth day and month of the cardholder for the other carrier: _____ (mm/dd)

Public-Funded Program – Have you applied for coverage through a public-funded program? ☐ Yes ☐ No

If No, please indicate why: _____

Part C – Authorization

I understand that the personal information I have provided herein is collected and used by Medavie Blue Cross to administer the terms of my policy or the group policy of which I am an eligible member, recommend suitable products and services that I am eligible for as a member of a policy, and other applicable purposes, as described in the Medavie Blue Cross Privacy Statement at **www.medaviebc.ca.**

Depending on the type of coverage I carry, limited personal information such as claim, health and/or financial related data may be collected from and/or released to following third parties as required for the purposes of administering and managing the benefits outlined in the policy of which I am an eligible member. These third parties may include healthcare providers, other insurance companies, regulatory authorities and investigative bodies, services providers, and/or the cardholder of any contract under which I am a participant.

Where allowed by law, my information may be shared with Medavie Blue Cross employees or service providers in jurisdictions other than where it was collected. If I am a resident of Quebec, this includes transferring or disclosing my personal information to Medavie Blue Cross employees or service providers outside of that province.

I understand that my consent is only valid for the time it is needed to achieve the purposes outlined herein, unless I withdraw it. I understand I may withdraw my consent at any time. However, in some instances doing so may prevent Medavie Blue Cross from providing me with certain products or services that may be useful to me and/or my dependents. This consent complies with federal and provincial privacy laws.

For more details about our information practices, including how your personal information is protected, how to access or correct personal information, or if you have concerns or questions, please see our Medavie Blue Cross Privacy Statement available at www.medaviebc.ca or call 1-800-667-4511.

Signature of Patient: **Date:** _____ (mm/dd/yyyy)

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3 SPECIALTY DRUG INFORMATION

Name of patient: _____ Date of Birth: _____

Policy Number: _____ ID Number: _____

E-mail address of patient or of legal guardian if patient is underage: _____

3A Patient Support Program (PSP) Enrollment

Is patient enrolled in the Patient Support Program? ☐ No ☐ Yes, specify Program ID #: _____

Indicate the name of the Patient Support Program: _____

PSP phone #: _____ PSP Fax #: _____

Product Name	Strength	Dosage	Diagnosis
LYNPARZA (OLAPARIB)			

Patient weight: _____ ☐ lbs ☐ kg Number of vials / syringes per dose: _____

Date treatment was initiated: _____ (mm/dd/yyyy) Expected duration of treatment: _____

Date of diagnosis: _____ (mm/dd/yyyy) Was treatment initiated in hospital? ☐ Yes ☐ No

Where is medication being administered? _____

Indicate the specialty of the physician who initiated or recommended the treatment: _____

Indicate if the disease or injury is work related: ☐ Yes ☐ No

For Initial Request, please complete sections 3B and 3D. For Renewals, please complete sections 3C and 3D.

3B Initial Request

Please list all the drugs that were previously tried, or could not be tried because of contraindications. The information for the following drug categories (relative to each diagnosis) must be provided, including details on the contraindication if applicable.

- **Breast cancer: radiotherapy, anthracycline- or taxane-based chemotherapy, definitive local treatment, endocrine therapy, any other treatment**
- **Epithelial ovarian, fallopian tube or primary peritoneal cancer: PARP inhibitor, platinum-based chemotherapy, Zejula (niraparib), Lynparza (olaparib)**
- **Prostate cancer: androgen synthesis inhibitor, second-generation androgen receptor inhibitor, PARP inhibitor**
- **Adenocarcinoma of the pancreas: platinum-based chemotherapy (FOLFIRINOX, Gemcitabine + cisplatin, FOLFOX, CAPOX)**

Category	Product Name	Dosage	Duration of Treatment	Response to Treatment or Contraindication

Is the drug being prescribed according to the Health Canada product monograph? ☐ Yes ☐ No

*NOTE: Do not provide genetic test results.

Approved indications from Health Canada:

Clinical information

ECOG score (date): _____ (_____) (mm/dd/yyyy)

Pharmacological treatment:

☐ First-line treatment ☐ Second-line treatment ☐ Third-line treatment or more

The cancer is:

☐ Metastatic ☐ Nonmetastatic ☐ Relapsed ☐ Locally advanced ☐ Other (specify): _____

3 SPECIALTY DRUG INFORMATION

Name of patient: _____ Date of Birth: _____

Policy Number: _____ ID Number: _____

E-mail address of patient or of legal guardian if patient is underage: _____

3B Initial Request (cont'd)

Clinical information (cont'd)

Administration of treatment:

☐ As monotherapy

☐ In combination with Zytiga (abiraterone) and prednisone or prednisolone

☐ In combination with hormone therapy. Specify: _____

☐ In combination with other medications. Specify: _____

BRCA gene:

☐ Wild-type

☐ Confirmed mutation → ☐ Deleterious ☐ Suspected deleterious

↳ ☐ Germline ☐ Somatic

↳ ☐ BRCA1 ☐ BRCA2 ☐ ATM

☐ Other. Specify: _____

1. **Breast cancer**

→ Estrogen receptor: ☐ Positive ☐ Negative → Progesterone receptor: ☐ Positive ☐ Negative

→ HER2 receptor: ☐ Positive ☐ Negative → In situ hybridization (ISH) result: ☐ Positive ☐ Negative

→ Immunohistochemistry (IHC) score: _____

Adjuvant treatment of breast cancer: ☐ Yes ☐ No Early stage: ☐ Yes ☐ No

Is the cancer considered to be resistant to PARP inhibitors? ☐ Yes ☐ No

Previous treatments:

Did the patient receive definitive local treatment? ☐ Yes ☐ No → If yes, specify: _____

Did the patient receive neoadjuvant or adjuvant chemotherapy? ☐ Yes ☐ No

↳ If yes, specify the number of cycles: _____

Indicate the date the above treatments were completed (if applicable): _____ (mm/dd/yyyy)

Have all options been exhausted for endocrine therapy? ☐ Yes ☐ No

Is endocrine therapy inappropriate? ☐ Yes ☐ No

↳ If yes, indicate the reason: _____

Has the cancer been completely resected? ☐ Yes ☐ No

Number of treatment(s) received in the metastatic setting: _____

Date of the last treatment (surgery, radiotherapy or chemotherapy): _____ (mm/dd/yyyy)

3 SPECIALTY DRUG INFORMATION

Name of patient: _____ Date of Birth: _____

Policy Number: _____ ID Number: _____

E-mail address of patient or of legal guardian if patient is underage: _____

3B Initial Request (cont'd)

1. **Breast cancer (cont'd)**

Pathologically confirmed positive lymph nodes and treatment with adjuvant chemotherapy: ☐ Yes ☐ No

↳ If yes, indicate the number of positive lymph nodes before chemotherapy: _____

Incomplete response to neoadjuvant chemotherapy: ☐ Yes ☐ No

↳ If yes, indicate the CPS+EG score: _____

2. **Epithelial ovarian, fallopian tube or primary peritoneal cancer**

Diagnosis: ☐ Epithelial ovarian cancer ☐ Fallopian tube cancer ☐ Primary peritoneal cancer

↳ ☐ Serous ovarian cancer ☐ Endometrioid ovarian cancer ☐ Other (specify): _____

Indicate the grade of the cancer: ☐ High ☐ Low FIGO stage: ☐ I ☐ II ☐ III ☐ IV

Cancer maintenance treatment: ☐ Yes ☐ No

Is the cancer considered to be resistant to PARP inhibitors? ☐ Yes ☐ No

Did the patient receive treatment with Lynparza (olaparib) before relapse? ☐ Yes ☐ No

Previous platinum-based chemotherapy treatments:	
First-line chemotherapy Protocol: _____ Number of cycles: _____ Date of last dose: _____ (mm/dd/yyyy)	Response: ☐ Complete ☐ Partial ☐ Other. Specify: _____ Objective tumour response: ☐ Yes ☐ No
Second-last chemotherapy Protocol: _____ Number of cycles: _____ Date of last dose: _____ (mm/dd/yyyy)	Has the disease progressed more than 6 months after the end? ☐ Yes ☐ No Response: ☐ Complete ☐ Partial ☐ Other. Specify: _____ Response duration: _____ months
Last chemotherapy Protocol: _____ Number of cycles: _____ Date of last dose: _____ (mm/dd/yyyy)	Objective tumour response: ☐ Yes ☐ No

3 SPECIALTY DRUG INFORMATION

Name of patient: _____ Date of Birth: _____
Policy Number: _____ ID Number: _____
E-mail address of patient or of legal guardian if patient is underage: _____

3B Initial Request (cont'd)

3. **Prostate cancer**

The cancer is: ☐ Castration resistant ☐ Hormone-sensitive (castration-sensitive)

Stage of cancer: ☐ I ☐ II ☐ III ☐ IV ☐ Other (specify): _____

Response to previous treatments:

Failure of androgen deprivation therapy: ☐ Yes ☐ No

↳ If yes, is the patient asymptomatic or mildly symptomatic? ☐ Yes ☐ No

Did patient have disease progression during treatment with a new hormonal agent/androgen receptor axis-targeted therapy (ARAT) (abiraterone, enzalutamide, apalutamide, and darolutamide) for prostate cancer? ☐ Yes ☐ No

↳ If yes, please specify the description of the cancer at the time of that treatment:

☐ Metastatic ☐ Nonmetastatic ☐ Hormone-sensitive ☐ Castration-resistant

Did the patient have disease progression during treatment with a PARP inhibitor (olaparib, talazoparib, niraparib) for prostate cancer? ☐ Yes ☐ No

↳ If yes, please specify the description of the cancer at the time of that treatment:

☐ Metastatic ☐ Nonmetastatic ☐ Hormone-sensitive ☐ Castration-resistant

Is this the first treatment the patient is receiving for their metastatic castration-resistant prostate cancer? ☐ Yes ☐ No

4. **Adenocarcinoma of the pancreas**

Maintenance treatment following first-line platinum-based chemotherapy: ☐ Yes ☐ No

Previous platinum-based chemotherapy treatment:	
First-line chemotherapy Protocol: _____ Duration of treatment: _____ weeks	Response: ☐ Complete ☐ Partial ☐ Stabilization of the disease ☐ Other. Specify: _____ Objective tumour response: ☐ Yes ☐ No

3 SPECIALTY DRUG INFORMATION

Name of patient: _____ Date of Birth: _____

Policy Number: _____ ID Number: _____

E-mail address of patient or of legal guardian if patient is underage: _____

3C Renewal Request

Please provide information on the evolution of the disease to evaluate the response to treatment.

Date of initial evaluation (pretreatment): _____ (mm/dd/yyyy)

Date of most recent evaluation: _____ (mm/dd/yyyy)

Information necessary to the assessment of this request

Response to treatment:

☐ Partial ☐ Complete ☐ Stabilization of the disease ☐ Progression of the disease

Response to treatment confirmed by radiographic evidence: ☐ Yes ☐ No

3D Additional Information

Please indicate any additional information pertaining to this request.

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HEALTH PROFESSIONAL STATEMENT

I certify that I have reviewed all pages of this request and that all information provided is true, correct and complete.

First Name: _____ Last Name: _____ Permit Number: _____

Specialty: _____

Clinic Name: _____

Address: _____ Suite: _____

City: _____ Province: _____ Postal Code: _____

E-mail: _____ Telephone: _____ Fax: _____

Signature: Date: _____
(mm/dd/yyyy)

It is important to provide the requested information in detail to help avoid delay in assessing claims for the above drug. This form may be subject to audit.

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