

The purpose of this form is to obtain information required to assess your drug claim. Approval for coverage of this drug may be reassessed at any time at Great-West Life's discretion. For additional information regarding Prior Authorization and Health Case Management, please visit our Great-West Life website at [www.greatwestlife.com](http://www.greatwestlife.com).

**IMPORTANT:** Please answer all questions. Your claim assessment will be delayed if this form is incomplete or contains errors.

**Any costs incurred for the completion of this form are the responsibility of the plan member/patient.**

Great-West Life recognizes and respects the importance of privacy. Personal information collected is used for the purposes of assessing eligibility for this drug and for administering the group benefits plan. For a copy of our Privacy Guidelines, or if you have questions about Great-West Life's personal information policies and practices (including with respect to service providers), refer to [www.greatwestlife.com](http://www.greatwestlife.com) or write to Great-West Life's Chief Compliance Officer.

I authorize Great-West Life, any healthcare provider, my plan administrator, any insurance or reinsurance company, administrators of government benefits or patient assistance programs or other benefits programs, other organizations, or service providers working with Great-West Life or any of the above, located inside or outside Canada, to exchange personal information when relevant and necessary for these purposes. I understand that personal information may be subject to disclosure to those authorized under applicable law within or outside Canada.

I acknowledge that the personal information is needed to assess eligibility for this drug and to administer the group benefits plan. I acknowledge that providing consent will help Great-West Life to assess my claim and that refusing to consent may result in delay or denial of my claim. Great-West Life reserves the right to audit the information provided on this form at any time and this consent extends to any audit of my claim. This consent may be revoked by me at any time by sending written instruction to that effect.

If the patient is a person other than myself, I confirm that the patient has given their consent to provide their personal information and for Great-West Life to use and disclose it as set out above.

I certify that the information given below is true, correct, and complete to the best of my knowledge. Failure to provide true, correct and complete information on this form could result in revocation of any approval decision, a requirement to repay paid claims or other appropriate action.

Plan Member's signature: \_\_\_\_\_ Date: \_\_\_\_\_

### **Form Completion Instructions:**

- 1. Complete "Patient Information" sections.**
- 2. Have the prescribing physician complete the "Physician Information" sections.**
- 3. Send all pages of the completed form to us by mail, fax or email as noted below.**

**Note:** As email is not a secure medium, any person with concerns about their prior authorization form/medical information being intercepted by an unauthorized party is encouraged to submit their form by other means.

**Mail to: The Great-West Life Assurance Company  
Drug Services  
PO Box 6000  
Winnipeg MB R3C 3A5**

**Fax to: The Great-West Life Assurance Company  
Fax 1-204-946-7664  
Attention: Drug Services**

**Email to: [gwldrug.services@gwl.ca](mailto:gwldrug.services@gwl.ca)  
Attention: Drug Services**

*(Continued on next page)*

**Plan Member Information – Complete all sections of this page (please print)**

Plan Member:		Patient Name:	
Plan Name:	Plan Number:	Plan Member ID Number:	
Patient Date of Birth (DD/MM/YYYY):	Address (number, street, city, province, postal code):		

Please indicate preferred contact number and if there are any times when telephone contact with you about your claim would be most convenient.

May we contact you by email? (Note that some correspondence may still need to be sent by regular mail).

Yes  No If yes, please provide email address: \_\_\_\_\_

**Tell us if you have been on this drug before**

Is the patient currently on, or previously been on Lynparza?  Yes  No

If Yes, a) indicate start date (DD/MM/YYYY): \_\_\_\_\_

b) coverage provided by: \_\_\_\_\_

(if coverage is not provided by Great-West Life please provide pharmacy print-out showing purchase of Lynparza)

**Tell us if you have coverage with any other benefits plan**

Does the patient have drug coverage under any other group benefits plan?  Yes  No

If Yes, name of other insurance company: \_\_\_\_\_

If other plan is with Great-West Life, tell us the plan and ID number: \_\_\_\_\_

Name of plan member: \_\_\_\_\_

Relationship to patient: \_\_\_\_\_

**Provide details and attach documentation of acceptance or decline:**

\_\_\_\_\_  
\_\_\_\_\_

**Tell us about any Provincial or other coverage you may have**

Does the patient have coverage under a provincial program or from any other source?  Yes  No

If Yes, name of program or other source: \_\_\_\_\_

Provide details and attach documentation of acceptance or decline: \_\_\_\_\_

Is the patient currently receiving disability benefits for the condition for which Lynparza has been prescribed?  Yes  No

**Tell us about any Patient Assistance Program you might be enrolled in**

Has the patient enrolled in the patient assistance program for Lynparza?  Yes  No

If Yes, please provide the following information:

1. Patient assistance program patient ID Number: \_\_\_\_\_

2. Patient assistance program contact person name and phone number:

Contact Name: \_\_\_\_\_ Phone Number: \_\_\_\_\_

(Continued on next page)

**Note to Physician:** In order to assess a patient's claim for this drug, we require detailed information on the patient's prescription drug history as requested below.

**Attach extra information if necessary. GENETIC TEST RESULTS ARE NOT REQUIRED**

**Physician's Information (please print)**

Name of prescribing physician: \_\_\_\_\_

Specialty: \_\_\_\_\_

Address (number, street, city, province, postal code): \_\_\_\_\_

Telephone Number (including area code): \_\_\_\_\_

Fax Number (including area code): \_\_\_\_\_

1. Prescribed dosage and regimen:

- 400mg twice daily
  - 300mg twice daily
  - Other (please specify): \_\_\_\_\_
- Provide rationale: \_\_\_\_\_

2. Health Canada indication (include date of initial diagnosis) (MM/YYYY): \_\_\_\_\_

- Ovarian, Fallopian Tube, or Peritoneal Cancer
- Metastatic Breast Cancer
- Other (approved by Health Canada): \_\_\_\_\_

Complete questions 1 – 6 and Other condition (Health Canada approved)

Is this drug being prescribed in accordance with approved Health Canada indications<sup>1</sup>?

- Yes, complete questions 1-6 and Physician's information
- No, condition not approved by Health Canada: \_\_\_\_\_

Complete questions 1-6 and Off-Label use

**<sup>1</sup>Approved Health Canada Indications and Clinical Use for Lynparza**

- Monotherapy for the maintenance treatment of adult patients with platinum-sensitive relapsed (PSR) BRCA wild type high-grade epithelial ovarian, fallopian tube or primary peritoneal cancer who are in response (complete response or partial response) to platinum-based chemotherapy, has been issued marketing authorization with conditions, pending the results of trials to verify its clinical benefit.
- Monotherapy for the maintenance treatment of adult patients with advanced BRCA-mutated high-grade epithelial ovarian, fallopian tube or primary peritoneal cancer who are in response (complete response or partial response) to first-line platinum-based chemotherapy. Patients must have confirmation of BRCA mutation (identified by either germline or tumour testing) before Lynparza treatment is initiated.
- Monotherapy for the maintenance treatment of adult patients with platinum-sensitive relapsed (PSR) BRCA-mutated high-grade epithelial ovarian, fallopian tube or primary peritoneal cancer who are in response (complete response or partial response) to platinum-based chemotherapy.
- Monotherapy for the treatment of adult patients with deleterious or suspected deleterious germline BRCA-mutated (gBRCAm), human epidermal growth factor receptor 2 (HER2)-negative metastatic breast cancer who have previously been treated with chemotherapy in the neoadjuvant, adjuvant or metastatic setting. Patients with hormone receptor (HR)- positive breast cancer should have progressed on or be considered inappropriate for endocrine therapy. Germline BRCA mutation must be confirmed before LYNPARZA treatment is initiated.

**Physician's Information (continued) (please print)**

3. What is the anticipated duration of treatment with this drug? \_\_\_\_\_

4. Where will treatment be administered?  Home  Physician's Office  Private clinic  Hospital in-patient  Hospital out-patient

5. Please provide medical rationale why Lynparza has been prescribed instead of an alternate drug in the same therapeutic class.

**Genetic test results are not required.**

\_\_\_\_\_

\_\_\_\_\_

**6. Drug and Treatment History – must be completed for every request.**

Drug(s) and Treatment(s) past and present	Dosing Regimen	Start Date (DD/MM/YYYY)	End Date (DD/MM/YYYY)	Clinical Results/Outcome
				<input type="checkbox"/> Failure <input type="checkbox"/> Intolerance <input type="checkbox"/> Other Clinical details: _____ _____
				<input type="checkbox"/> Failure <input type="checkbox"/> Intolerance <input type="checkbox"/> Other Clinical details: _____ _____
				<input type="checkbox"/> Failure <input type="checkbox"/> Intolerance <input type="checkbox"/> Other Clinical details: _____ _____

**Ovarian, Fallopian Tube or Peritoneal Cancer – Genetic test results are not required**

How many regimens of platinum-based chemotherapy has the patient completed? \_\_\_\_\_

Provide details in medication chart

Is Lynparza being prescribed as monotherapy?  Yes  No

Is the patient currently in response (complete response or partial response) to platinum-based chemotherapy?  Yes  No

If patient has received more than one regimen of platinum-based chemotherapy, has platinum sensitivity been confirmed? (defined as disease progressing at least 6 months after completion of the penultimate platinum chemotherapy)  Yes  No

**Breast Cancer – Genetic test results are not required**

Has the patient received prior treatment with anthracycline- and a taxane- containing regimen in adjuvant or metastatic setting?  Yes  No

Has the patient received prior endocrine therapy, if deemed appropriate for endocrine therapy?  Yes  No

Has the patient progressed with platinum-based therapy?  Yes  No

Has the patient received more than 2 prior lines of chemotherapy in the metastatic setting?  Yes  No

Has the patient had prior treatment with other PARP inhibitors?  Yes  No

**Other condition (Health Canada approved) – Genetic test results are not required**

Please provide any relevant information related to the disease and attach supporting documentation.

\_\_\_\_\_

\_\_\_\_\_

**Off-label use – Genetic test results are not required**

**Questions 1 - 6 must be completed.**

Date of initial diagnosis (DD/MM/YYYY): \_\_\_\_\_

Is there clinical evidence supporting the off-label use of this drug?  Yes  No

Provide clinical literature/studies to support the request for off-label use, such as:

- At least two Phase II or two Phase III clinical trials showing consistent results of efficacy; and
- Published recommendations in evidence-based guidelines supporting its use.

Provide medical rationale why Lynparza has been prescribed off-label instead of an alternate drug with an approved indication for this condition.

Provide any pertinent medical history or information to support this off-label request.

\_\_\_\_\_

\_\_\_\_\_

If this is a renewal request, provide documentation showing treatment efficacy since previous request.

\_\_\_\_\_

\_\_\_\_\_

**Note for Physician: To be eligible for reimbursement, Great-West Life may require your patient to purchase a drug requiring prior authorization from a pharmacy designated by Great-West Life. If applicable, a health case manager will contact you with further information.**

**I certify that the information provided is true, correct, and complete.**

Physician's Signature: \_\_\_\_\_ Date: \_\_\_\_\_

License Number: \_\_\_\_\_

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. The completed form can be returned to Great-West Life by mail, fax, or email.

**Note:** As email is not a secure medium, any person with concerns about their prior authorization form/medical information being intercepted by an unauthorized party is encouraged to submit their form by other means.

**Mail to: The Great-West Life Assurance Company  
Drug Services  
PO Box 6000  
Winnipeg MB R3C 3A5**

**Fax to: The Great-West Life Assurance Company  
Fax 1-204-946-7664  
Attention: Drug Services**

**Email to: [gwldrug.services@gwl.ca](mailto:gwldrug.services@gwl.ca)  
Attention: Drug Services**