

The purpose of this form is to obtain information required to assess your drug claim. For additional information regarding Prior Authorization and Health Case Management, please visit our Great-West Life website at 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 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
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 Opdivo? 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 Opdivo)

Tell us if you have coverage with any other benefits plan

Does the patient have drug coverage with 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 Opdivo 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 Opdivo? 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:

3mg/kg IV every 2 weeks

1mg/kg every 3 weeks x 4 doses, followed by 3mg/kg every 2 weeks (in combination with ipilimumab)

Other (please specify): _____

Provide rationale: _____

Patients weight: _____ kg (for weight-based dosing)

Date determined (MM/YYYY): _____

2. Health Canada Indication (include date of initial diagnosis) (MM/YYYY): _____

Unresectable or metastatic melanoma in previously untreated patients – monotherapy

Unresectable or metastatic melanoma in previously untreated patients – in combination with ipilimumab (Yervoy)

Unresectable or metastatic melanoma and disease progression following ipilimumab

Locally advanced or metastatic non-small cell lung cancer (NSCLC) with progression on or after platinum-based chemotherapy

Advanced or metastatic renal cell carcinoma (RCC) in patients who have received prior anti-angiogenic therapy - monotherapy

Intermediate/poor-risk advanced or metastatic RCC - in combination with ipilimumab (Yervoy)

Recurrent or metastatic squamous cell carcinoma of the head and neck (SCCHN)

Classical Hodgkin lymphoma that has relapsed or progressed after 1) autologous stem cell transplantation (ASCT) and brentuximab vedotin, or 2) 3 or more lines of systematic therapy including ASCT

Advanced or metastatic hepatocellular carcinoma who are intolerant to or have progressed on sorafenib therapy - monotherapy

Other (Approved by Health Canada): _____

Complete questions 1 - 6 and Other Condition (Health Canada approved)

Genetic test results are not required

Is this drug being prescribed in accordance with approved Health Canada indications'?

Yes, complete questions 1 - 6 and Physician's information

No, prescribed use is not approved by Health Canada: _____

Complete questions 1 - 6 and Off-label use

Physician's Information (continued) (please print)

***1*Approved Health Canada Indications and Clinical Use for Opdivo:**

- Treatment of unresectable or metastatic BRAF V600 wild-type melanoma in previously untreated adults.
- Treatment of unresectable or metastatic BRAF V600 mutation-positive melanoma in previously untreated adults.
- In combination with ipilimumab for the treatment of unresectable or metastatic melanoma in previously untreated adults.
- Treatment of patients with unresectable or metastatic melanoma and disease progression following ipilimumab and, if BRAF V600 mutation-positive, a BRAF inhibitor.
- Treatment of adult patients with locally advanced or metastatic non-small cell lung cancer (NSCLC) with progression on or after platinum-based chemotherapy. Patients with EGFR or ALK genomic tumour aberrations should have disease progression on a therapy for these aberrations prior to receiving Opdivo.
- Treatment of adult patients with advanced or metastatic renal cell carcinoma (RCC) who have received prior anti-angiogenic therapy.
- In combination with ipilimumab, is indicated for the treatment of adult patients with intermediate/poor-risk advanced or metastatic RCC.
- Treatment of recurrent or metastatic squamous cell carcinoma of the head and neck (SCCHN) in adults progressing on or after platinum-based therapy.
- Treatment of classical Hodgkin lymphoma that has relapsed or progressed after 1) autologous stem cell transplantation (ASCT) and brentuximab vedotin, or 2) 3 or more lines of systematic therapy including ASCT
- Monotherapy treatment of advanced (not amenable to curative therapy or localtherapeutic measures) or metastatic hepatocellular carcinoma (HCC) who are intolerant to or have progressed on sorafenib therapy.

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 this drug 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: _____ _____

Physician's Information (continued) (please print)

Unresectable or Metastatic Melanoma – Genetic test results are not required

Has patient received previous therapy for the treatment of their melanoma? Yes No

If yes, please provide details in therapy chart above, including details regarding disease progression.

Will Opdivo be used in combination with ipilimumab (Yervoy)? Yes No

If yes, a separate form for Yervoy does not need to be filled out.

Metastatic Non-Small Cell Lung Cancer (NSCLC) – Genetic test results are not required

Has patient received previous therapy for the treatment of their NSCLC? Yes No

If yes, please provide details in therapy chart above, including details regarding disease progression.

Metastatic Renal Cell Carcinoma (RCC) – Genetic test results are not required

Has patient received previous therapy for the treatment of their RCC? Yes No

If yes, please provide details in medication chart above, including details regarding disease progression.

Will Opdivo be used in combination with ipilimumab (Yervoy)? Yes No

If yes, a separate Yervoy form does not need to be filled out.

Recurrent or Metastatic Squamous Cell Carcinoma of the Head and Neck (SCCHN) – Genetic test results are not required

Has patient received previous therapy for the treatment of their SCCHN? Yes No

If yes, please provide details in medication chart above, including details regarding disease progression.

Classical Hodgkin Lymphoma (cHL) – Genetic test results are not required

Has patient received previous therapy, including stem cell transplantation for the treatment of their cHL? Yes No

If yes, please provide details in therapy chart on the previous page, including details regarding disease progression.

Will Opdivo be used as monotherapy? Yes No

Please indicate patient's current ECOG score: _____

Advanced or metastatic hepatocellular carcinoma – Genetic test results are not required

Has the patient been previously treated and had disease progression or intolerance with sorafenib therapy? Yes No

If yes, please provide details in therapy chart above, including details regarding disease progression.

Will Opdivo be used as monotherapy? Yes No

Please indicate the patient's current ECOG score: _____

Renewal Request – Genetic test results are not required

Has patient experienced any disease progression while on Opdivo? Yes No

Please include any relevant medical records to support response, such as a copy of any recent CT or MRI reports and/or clinic notes.

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 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 this drug has been prescribed off-label instead of an alternative 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 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.

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Winnipeg MB R3C 3A5**

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