

Medical prior authorization form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206

Commercial (Traditional) This form applies to: Medicaid This request is: Urgent (life threatening) Non-Urgent (standard review) Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function. **Opdivo**® (nivolumab) Member Last Name: ___ First Name: DOB: _____ Gender: ____ ID #: _____ Primary Care Physician: Phys. Phone: Phys. Fax: _____ Requesting Physician: Physician Address: Physician NPI: Contact Name: Provider Signature: _____ **Product and Billing Information** □ New Request □ Continuation Request Drug product: Opdivo 100 mg/10mL Start date: Opdivo 40 mg / 4 mL solution Date of last dose: Date of next dose: Dose:_____ Dose Frequency:____ Height: Weight: Place of administration: ☐ Physician's office ☐ Outpatient infusion Facility: ☐ Home infusion NPI: Fax: Facility: _____ Billing: ☐ Physician to buy and bill ☐ Facility to buy and bill ☐ Specialty Pharmacy ICD-10 Diagnosis code(s):

Drug cost information

The wholesale acquisition cost for each 10 mg dose of Opdivo is \$239.80. The cost of treatment with this drug will vary depending on the patient's circumstances. Each one-year treatment is likely to be more than \$149,600.



Precertification Requirements

Patient must have one of the following diagnoses and meet listed criteria:

- Unresectable or metastatic melanoma. For Opdivo + Yervoy requests, must have first tried Opdivo or Keytruda as individual agents;
- Metastatic non-small cell lung cancer (NSCLC) that has progressed with previous platinum-based chemotherapy and EGFR or ALK targeted therapies for EGFR or ALK mutated disease;
- 3. Adjuvant treatment of melanoma with lymph node involvement/metastatic disease after undergoing complete resection
- 4. Advanced renal cell carcinoma that has progressed after anti-angiogenic therapy;
- 5. Advanced renal cell carcinoma in poor to intermediate risk disease in combination with Yervoy;
- 6. Advanced renal cell carcinoma in favorable risk disease in combination with Yervoy: must have first tried Sutent (sunitinib) or Votrient (pazopanib);
- 7. Hodgkin lymphoma that progressed or relapsed after autologous stem cell transplantation and post-transplant brentuximab vedotin (Adcetris).
- 8. Metastatic squamous cell head and neck cancer (non-nasopharyngeal) with disease progression on or after platinum-containing chemotherapy, not amendable to surgery.
- 9. Locally advanced or metastatic urothelial carcinoma, in patients with disease progression on or following platinum-containing chemotherapy, or within 12 months of neoadjuvant or adjuvant platinum-containing chemotherapy.
- 10. Microsatellite instability-high (MSI-H) or mismatch repair deficient (dMMR) metastatic colorectal cancer, that has progressed after treatment with a fluoropyrimidine-, oxaplatin-, and/or irinotecan-based therapy.
- 11. Hepatocellular carcinoma previously treated with sorafenib (Nexavar®), (Child-Pugh Class A or B7 only)

Requests for any condition not listed as covered require evidence of current medical literature that substantiates the drug's efficacy or that recognized oncology organizations generally accept the treatment for the condition.

Priority Health Precertification Documentation		
What condition is this drug being requested for? Unresectable or metastatic melanoma Metastatic non-small cell lung cancer (NSCLC) Adjuvant treatment of melanoma with lymph node involvement/metastatic disease after undergoing complete resection Advanced renal cell carcinoma Classical Hodgkin lymphoma Metastatic squamous cell head and neck cancer (non-nasopharyngeal) Locally advanced or metastatic urothelial carcinoma Metastatic colorectal cancer (MSI-H or dMMR) Hepatocellular carcinoma Other – rationale for use:		
For Unresectable or metastatic melanoma: A. Does the patient have BRAF v600 mutation positive unresectable or metastatic melanoma? Yes No B. If BRAF v600 mutation positive, what prior therapies have been tried?		
Dates:		
C. If this is a request for Opdivo + Yervoy combination therapy, please indicate what prior therapies have been tried:		
For adjuvant treatment of melanoma with lymph node involvement/metastatic disease after undergoing complete		
resection: A. Has the patient undergo a complete resection? Yes No		



FO	r metastatic NSCLC:
A.	Please indicate which of the following best describe the patient's disease type:
	☐ EGFR mutation-positive metastatic NSCLC
	☐ ALK mutation-positive metastatic NSCLC
	Metastatic NSCLC without EGFR or ALK mutations
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B.	Please provide which therapies have been tried:
	Dates: Dates:
	Dates.
C.	Does the member have squamous NSCLC or non-squamous NSCLC?
0.	Squamous NSCLC
	☐ Non-squamous NSCLC
	Unknown; rationale:
	r metastatic renal cell carcinoma:
A.	Please provide which anti-angiogenic therapies have been tried:
	Dates:
	Dates:
	and and the Bullet for the second second
	r classical Hodgkin lymphoma:
	Has the patient relapsed or progressed after autologous hematopoietic stem cell transplant AND post-transplantation
	brentuximab vedotin (Adcetris)?
	☐ No
Fo	r metastatic squamous cell head and neck cancer:
	Does the patient have nasopharyngeal cancer?
Λ.	
R	Please provide which therapies have been tried:
	Dates:
	Dates:
C.	Is patient a candidate for surgery?
	☐ Yes
	□ No
Fo	r locally advanced or metastatic urothelial carcinoma:
A.	Please provide which therapies have been tried:
	Dates:
	Dates:
	and the factor of the same of
	r metastatic colorectal cancer (MSI-H/dMMR):
Α.	Please provide which therapies have been tried:
	Drug:Dates:
	Drug:Dates:
	n h amata a alliulan a angin ama
	r hepatocellular carcinoma:
Α.	Has the patient been previously treated with sorafenib (Nexavar®)?
	☐ Yes
	□ No
R	Please provide Child-Pugh Class
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