

PRIOR AUTHORIZATION REQUEST FORM

EOC ID:

Opdivo (Nivolumab)

Phone: 800-728-7947 Fax back to: 866-880-4532

The Scott & White Health Plan Pharmacy Department manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above. Please note any information left blank or illegible may delay the review process.

Patient Name:	Prescriber Name: Supervising Physician:	
Member/Subscriber Number:	Fax:	Phone:
Date of Birth:	Office Contact:	
Group Number:	NPI:	State Lic ID:
Address:	Address:	
City, State ZIP:	City, State ZIP:	
Primary Phone:	Specialty/facility name (if applicable	e):
Drug Name and Strength:		
Directions / SIG:		
Please attach any pertinent medical history or information for this patient that may support approval. Please answer the following questions and sign.		
Q1. Please provide ICD code(s) for diagnosis.		
Q2. For which diagnosis is Opdivo (Nivolumab) being pres	cribed?	
Unresectable or metastatic melanoma		
☐ Metastatic, progressive non-small cell lung cancer (NSCLC)		
Advanced renal cell carcinoma		
☐ Classical Hodgkin lymphoma		
☐ Recurrent or metastatic squamous cell carcinoma of the head and neck		
☐ Locally advanced or metastatic urothelial carcinoma		
☐ Metastatic colorectal cancer		
Other (please specify)		
Q3. If you selected "other" in question 2, please provide documentation that use is consistent with a category 2A or higher recommendation per NCCN compendia or guidelines.		
Q4. Please indicate location of administration.		
☐ Home		



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Patient Name:	Supervising Physician:	
□ Long Term Care (LTC) facility □ Physician office (drug from office stock - buy and bill) □ Physician office (drug from pharmacy with a prescription)		
Q5. Is the prescriber an Oncologist or Hematologist? ☐ Yes ☐ No		
Q6. If using Opdivo for unresectable or metastatic melanoma, please select how Opdivo will be used from the options below. □ Opdivo will be used as a single agent for treatment of BRAF V600 wild-type or BRAF V600 mutation-positive disease □ Opdivo will be used in combination with ipilimumab (Yervoy) □ Other (Please Specify)		
Q7. If using for NSCLC and tumor has EGFR or ALK geno on approved EGFR or ALK directed therapy? Yes No N/A - Member does not have EGFR or ALK genomic to	mic tumor aberrations, has patient had disease progression umor aberrations	
Q8. If using Opdivo for NSCLC, squamous cell carcinoma have disease progression on or after platinum-containing of Yes	•	
Q9. If using Opdivo for advanced renal cell carcinoma, has	s patient received prior anti-angiogenic therapy?	
Q10. If using Opdivo for classical Hodgkin Lymphoma, has patient relapsed or progressed following one of the following? Autologous hematopoietic stem cell transplant (HSCT) and post-transplant brentuximab vedotin (Adcetris) or more lines of systemic therapy including autologous HSCT Other (please specify) None of the above		
Q11. If using Opdivo for metastatic colorectal cancer, does mismatch repair deficient (dMMR) with progression following irinotecan?		



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-	Prescriber Name:
Patient Name:	Supervising Physician:
Q12. Will the patient be using systemic corticosteroids and	/ or immunosuppressants while taking Opdivo?
☐ Yes ☐ No	
Q13. Does the patient have a history of severe immune-more requiring use of corticosteroids for 12 weeks for more?	ediated adverse reaction from treatment with ipilimumab,
☐ Yes ☐ No	
Q14. Additional Comments	
Prescriber Signature	Date
□ Expedited/Urgent - By checking this box and signing abov seriously jeopardize the life or health of the enrollee or the e	
	ssity denial. Requesting providers may speak to the SWHP medical ity to help impact the decision on a request before coverage has been

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