

# Opdualag™ (nivolumab/relatlimab-rmbw) (Intravenous)

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# I. Length of Authorization <sup>1</sup>

Coverage will be provided for 6 months and may be renewed.

# **II.** Dosing Limits

- A. Quantity Limit (max daily dose) [NDC Unit]:
  - Opdualag 240 mg/80 mg in a 20 mL single-dose vial: 2 vials per 28 days
- B. Max Units (per dose and over time) [HCPCS Unit]:
  - 160 billable units (480 mg nivolumab/160 mg relatlimab) every 28 days

# III. Initial Approval Criteria <sup>1</sup>

Coverage is provided for the following conditions:

• Patient is at least 12 years of age; **AND** 

## Universal Criteria 1-3

- Patient weighs at least 40 kg; AND
- Patient has not received previous therapy with a programmed death (PD-1/PD-L1)-directed therapy (e.g., cemiplimab, avelumab, pembrolizumab, atezolizumab, durvalumab, dostarlimab, retifanlimab, nivolumab, toripalimab, etc.), unless otherwise specified 4; AND

## Cutaneous Melanoma † ‡ Ф 1-4

- Patient has unresectable or metastatic\* disease; AND
  - o Used as first-line therapy; **OR**
  - Used as subsequent therapy for disease progression if single-agent anti-PD-1 therapy or combination checkpoint blockade was not previously used; OR
  - Used as re-induction therapy in patients with prior combination anti-PD-1/LAG-3
    therapy that resulted in disease control (i.e., complete response, partial response, or
    stable disease) with no residual toxicity, but with disease progression or relapse >3
    months after treatment discontinuation



\*Metastatic disease includes stage III unresectable/borderline resectable disease with clinically positive node(s) or clinical satellite/in-transit metastases, as well as unresectable local satellite/in-transit recurrence, unresectable nodal recurrence, and widely disseminated distant metastatic disease.

† FDA Approved Indication(s); ‡ Compendia Recommended Indication(s); ♠ Orphan Drug

## IV. Renewal Criteria 1-4

Coverage may be renewed based upon the following criteria:

- Patient continues to meet the universal and other indication-specific relevant criteria such
  as concomitant therapy requirements (not including prerequisite therapy), performance
  status, etc. identified in section III; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: severe infusion-related reactions, complications of allogeneic hematopoietic stem cell transplantation (HSCT), severe immune-mediated adverse reactions (i.e., pneumonitis, colitis, hepatitis, endocrinopathies, nephritis/renal dysfunction, myocarditis, dermatologic adverse reactions/rash, myocarditis, etc.), etc.; AND
- Disease response with treatment as defined by stabilization of disease or decrease in size of tumor or tumor spread

# Δ Notes:

- Patients who complete adjuvant therapy and progress ≥ 6 months after discontinuation are eligible to re-initiate PD-directed therapy for metastatic disease.
- Patients whose tumors, upon re-biopsy, demonstrate a change in actionable mutation (e.g., MSS initial biopsy; MSI-H subsequent biopsy) may be eligible to re-initiate PD-directed therapy and will be evaluated on a case-by-case basis.

# V. Dosage/Administration Δ1

Indication	Dose
Cutaneous	Adult and pediatric patients ≥12 years of age who weigh at least 40 kg:
Melanoma  • Administer 480 mg nivolumab and 160 mg relatlimab (contents of 2 vials)	
	intravenously every 4 weeks until disease progression or unacceptable toxicity.

# VI. Billing Code/Availability Information

## **HCPCS Code**:

• J9298 – Injection, nivolumab and relatlimab-rmbw, 3 mg/1 mg; 1 billable unit = 3 mg nivolumab/1 mg relatlimab-rmbw

### NDC:

Opdualag 240 mg of nivolumab and 80 mg of relatlimab per 20 mL single-dose vial: 00003-7125-xx



### VII. References

- 1. Opdualag [package insert]. Princeton, NJ; Bristol-Myers Squibb Company; March 2022. Accessed January 2024.
- 2. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) nivolumab-relatlimab. National Comprehensive Cancer Network, 2024. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed January 2024.
- 3. Tawbi HA, Schadendorf D, Lipson EJ; RELATIVITY-047 Investigators, et al. Relatlimab and nivolumab versus nivolumab in untreated advanced melanoma. *N Engl J Med*. 2022;386:24-34.
- 4. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) Melanoma: Cutaneous. Version 3.2023. National Comprehensive Cancer Network, 2024. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed January 2024.

# **Appendix 1 – Covered Diagnosis Codes**

ICD-10	ICD-10 Description	
C43.0	Malignant melanoma of lip	
C43.111	Malignant melanoma of right upper eyelid, including canthus	
C43.112	Malignant melanoma of right lower eyelid, including canthus	
C43.121	Malignant melanoma of left upper eyelid, including canthus	
C43.122	Malignant melanoma of left lower eyelid, including canthus	
C43.20	Malignant melanoma of unspecified ear and external auricular canal	
C43.21	Malignant melanoma of right ear and external auricular canal	
C43.22	Malignant melanoma of left ear and external auricular canal	
C43.30	Malignant melanoma of unspecified part of face	
C43.31	Malignant melanoma of nose	
C43.39	Malignant melanoma of other parts of face	
C43.4	Malignant melanoma of scalp and neck	
C43.51	Malignant melanoma of anal skin	
C43.52	Malignant melanoma of skin of breast	
C43.59	Malignant melanoma of other part of trunk	



C43.60	Malignant melanoma of unspecified upper limb, including shoulder	
C43.61	Malignant melanoma of right upper limb, including shoulder	
C43.62	Malignant melanoma of left upper limb, including shoulder	
C43.70	Malignant melanoma of unspecified lower limb, including hip	
C43.71	Malignant melanoma of right lower limb, including hip	
C43.72	Malignant melanoma of left lower limb, including hip	
C43.8	Malignant melanoma of overlapping sites of skin	
C43.9	Malignant melanoma of skin, unspecified	
Z85.820	Personal history of malignant melanoma of skin	

# Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

The preceding information is intended for non-Medicare coverage determinations. Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determinations (NCDs) and/or Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. Local Coverage Articles (LCAs) may also exist for claims payment purposes or to clarify benefit eligibility under Part B for drugs which may be self-administered. The following link may be used to search for NCD, LCD, or LCA documents: <a href="https://www.cms.gov/medicare-coverage-database/search.aspx">https://www.cms.gov/medicare-coverage-database/search.aspx</a>. Additional indications, including any preceding information, may be applied at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCD/LCA): N/A

Medicare Part B Administrative Contractor (MAC) Jurisdictions				
Jurisdiction	Applicable State/US Territory	Contractor		
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC		
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC		
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)		
6	MN, WI, IL	National Government Services, Inc. (NGS)		
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.		
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)		
N (9)	FL, PR, VI	First Coast Service Options, Inc.		
J (10)	TN, GA, AL	Palmetto GBA, LLC		
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA, LLC		
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.		
K (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)		
15	KY, OH	CGS Administrators, LLC		

