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<b>Section:</b>	Prescription Drugs	<b>Effective Date:</b>	January 1, 2025
<b>Subsection:</b>	Antineoplastic Agents	<b>Original Policy Date:</b>	April 15, 2022
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**Last Review Date:** December 13, 2024

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## Opdualag

### Description

#### Opdualag (nivolumab and relatlimab-rmbw)

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#### Background

Opdualag is a human IgG4 monoclonal antibody and is a combination of nivolumab, a programmed death receptor-1 (PD-1) blocking antibody, and relatlimab, a lymphocyte activation gene-3 (LAG-3) blocking antibody. Antagonism of the LAG-3 pathway promotes T-cell proliferation and cytokine secretion. Upregulation of PD-1 ligands occurs in some tumors which inhibits T-cell proliferation and cytokine production. PD-1 blocking by nivolumab decreases tumor growth. LAG-3 blockade potentiates the anti-tumor activity of PD-1 blockage, inhibiting tumor growth and promoting tumor regression (1).

#### Regulatory Status

FDA-approved indication: Opdualag is indicated for the treatment of adult and pediatric patients 12 years of age or older with unresectable or metastatic melanoma (1).

Opdualag carries warnings regarding severe and fatal immune-mediated adverse reactions, infusion-related reactions, and complication of allogenic hematopoietic stem cell transplantation (1).

Opdualag can also cause fetal harm when administered to a pregnant woman. Pregnant women should be advised of the potential risk to a fetus. Females of reproductive potential should be advised to use effective contraception during treatment with Opdualag and for at least 5 months after the last dose of Opdualag (1).

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The safety and effectiveness of Opdualag have not been established in pediatric patients less than 12 years of age or in pediatric patients 12 years or older who weigh less than 40 kg (1).

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## Related Policies

Keytruda, Opdivo

## Policy

*This policy statement applies to clinical review performed for pre-service (Prior Approval, Precertification, Advanced Benefit Determination, etc.) and/or post-service claims.*

Opdualag may be considered **medically necessary** if the conditions indicated below are met.

Opdualag may be considered **investigational** for all other indications.

## Prior-Approval Requirements

**Age** 12 years of age or older

### Diagnosis

Patient must have the following:

1. Unresectable or metastatic melanoma
  - a. Age 12-17 **only**: patient weight  $\geq$  40 kg

**AND ALL** of the following:

- a. Prescriber agrees to monitor liver enzymes, creatinine, and thyroid function at baseline and periodically during treatment
- b. Prescriber agrees to discontinue treatment for any immune mediated adverse reaction (encephalitis, nephritis, rash, decreased renal function, and endocrinopathies) or disease progression
- c. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Opdualag and for 5 months after the last dose

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## Prior – Approval *Renewal* Requirements

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**Age** 12 years of age or older

## Diagnosis

Patient must have the following:

1. Unresectable or metastatic melanoma
  - a. Age 12-17 **only**: patient weight  $\geq$  40 kg

**AND ALL** of the following:

- a. **NO** disease progression or unacceptable toxicity
- b. Prescriber agrees to monitor liver enzymes, creatinine, and thyroid function periodically during treatment
- c. Prescriber agrees to discontinue treatment for any immune mediated adverse reaction (encephalitis, nephritis, rash, decreased renal function, and endocrinopathies) or disease progression
- d. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Opdualag and for 5 months after the last dose

## Policy Guidelines

### Pre - PA Allowance

None

### Prior - Approval Limits

**Quantity** 6 single-dose vials every 84 days

**Duration** 12 months

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### Prior – Approval *Renewal* Limits

Same as above

## Rationale

### Summary

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Opdualag is a combination of nivolumab and relatlimab and is indicated for unresectable or metastatic melanoma. Opdualag has warnings including severe and fatal immune-mediated adverse reactions and infusion-related reactions. The safety and effectiveness of Opdualag have not been established in pediatric patients less than 12 years of age or in pediatric patients 12 years or older who weigh less than 40 kg (1).

Prior authorization is required to ensure the safe, clinically appropriate, and cost-effective use of Opdualag while maintaining optimal therapeutic outcomes.

### References

1. Opdualag [package insert]. Princeton, NJ: Bristol-Myers Squibb Company; March 2024.
2. NCCN Drugs & Biologics Compendium<sup>®</sup> Nivolumab and relatlimab-rmbw 2024. National Comprehensive Cancer Network, Inc. Accessed on October 10, 2024.

### Policy History

Date	Action
April 2022	Addition to PA
June 2022	Annual review and reference update
September 2022	Annual review and reference update
September 2023	Annual review and reference update
March 2024	Annual review and reference update
September 2024	Annual review and reference update
December 2024	Annual review and reference update

### Keywords

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**This policy was approved by the FEP<sup>®</sup> Pharmacy and Medical Policy Committee on December 13, 2024 and is effective on January 1, 2025.**