

PHARMACY POLICY STATEMENT

Indiana Medicaid

DRUG NAME	Nemluvio (nemolizumab-ilto)
BENEFIT TYPE	Pharmacy
STATUS	Prior Authorization Required

Nemluvio, approved in 2024, is an interleukin-31 receptor antagonist indicated for the treatment of adults with prurigo nodularis and the treatment of adults and pediatric patients 12 years of age and older with moderate-to-severe atopic dermatitis in combination with topical corticosteroids and/or calcineurin inhibitors when the disease is not adequately controlled with topical prescription therapies. It inhibits IL-31 signaling by binding selectively to IL-31 RA. IL-31 is a naturally occurring cytokine that is involved in pruritus, inflammation, epidermal dysregulation, and fibrosis.

Nemluvio (nemolizumab-ilto) will be considered for coverage when the following criteria are met:

Prurigo Nodularis (PN)

For **initial** authorization:

- 1. Member is at least 18 years of age; AND
- 2. Medication must be prescribed by or in consultation with a dermatologist; AND
- 3. Member has a documented diagnosis of prurigo nodularis with **ALL** of the following:
 - a) Chronic pruritis lasting more than 6 weeks;
 - b) Severe itch with repeated scratching;
 - c) At least 20 total lesions present; AND
- 4. Member has had a trial and failure of a medium to super high potency topical corticosteroid for at least 2 weeks. NOTE: If topical corticosteroids are contraindicated, must alternatively try and fail at least ONE of the following: topical emollient, capsaicin, calcipotriol, calcineurin inhibitor, phototherapy, or oral antihistamine; AND
- 5. Member has tried and failed Dupixent.
- 6. **Dosage allowed/Quantity limit:** quantity limit 2 pens per 4 weeks.
 - a) Less than 90 kg: administer 60 mg (two 30 mg injections) subcutaneously, followed by 30 mg given every 4 weeks.
 - b) 90 kg or more: administer 60 mg (two 30 mg injections) subcutaneously, followed by 60 mg given every 4 weeks.

If all the above requirements are met, the medication will be approved for 4 months.

For **reauthorization**:

1. Chart notes have been provided documenting clinically significant reduction of itch intensity and/or nodule clearance compared to baseline.

If all the above requirements are met, the medication will be approved for an additional 12 months.

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Atopic Dermatitis (AD)

For **initial** authorization:

- 1. Member is at least 12 years of age; AND
- Medication must be prescribed by or in consultation with a dermatologist, allergist, or immunologist;
- 3. Member has a diagnosis of moderate to severe AD; AND
- 4. Provider attests Nemluvio will be used in combination with topical corticosteroids and/or topical calcineurin inhibitors: AND
- 5. Member's atopic dermatitis involves 10% or more of the body surface area (BSA) OR involves highly visible or functional areas (e.g., neck, face, genitals, palms) and is significantly impairing quality of life;
- 6. Member has a documented trial and failure of **ONE** of the following:
 - a) Medium to very high potency topical corticosteroids for 2 weeks;
 - b) Topical calcineurin inhibitor (e.g., tacrolimus, pimecrolimus) for 6 weeks. *Note:* Eucrisa for 4 weeks is also acceptable.
- 7. **Dosage allowed/Quantity limit:** inject 60 mg (two 30 mg injections) subcutaneously, followed by 30 mg given every 4 weeks. After 16 weeks of treatment, for patients who achieve clear or almost clear skin, a dosage of 30 mg every 8 weeks is recommended. Quantity limit: 1 pen per 28 days.

If all the above requirements are met, the medication will be approved for 4 months.

For reauthorization:

1. Chart notes have been provided showing an improvement in signs and symptoms of disease such as fewer flares, less itching/erythema, improved quality of life, etc.

If all the above requirements are met, the medication will be approved for an additional 12 months.

CareSource considers Nemluvio (nemolizumab-ilto) not medically necessary for the treatment of conditions that are not listed in this document. For any other indication, please refer to the Off-Label policy.

DATE	ACTION/DESCRIPTION
09/10/2024	New policy for Nemluvio created.
02/17/2025	Added AD indication.

References:

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- 4. Satoh T, Yokozeki H, Murota H, et al. 2020 guidelines for the diagnosis and treatment of prurigo. *J Dermatol.* 2021;48(9):e414-e431. doi:10.1111/1346-8138.16067.
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- 9. Eichenfield LF, Tom WL, Berger TG, et al. Guidelines of care for the management of atopic dermatitis: section 2. Management and treatment of atopic dermatitis with topical therapies. *J Am Acad Dermatol.* 2014;71(1):116-132.

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- 11. Davis DMR, Drucker AM, Alikhan A, et al. Guidelines of care for the management of atopic dermatitis in adults with phototherapy and systemic therapies. *J Am Acad Dermatol.* 2024;90(2):e43-e56. doi:10.1016/j.jaad.2023.08.102
- 12. AAAAI/ACAAI JTF Atopic Dermatitis Guideline Panel, Chu DK, Schneider L, et al. Atopic dermatitis (eczema) guidelines: 2023 American Academy of Allergy, Asthma and Immunology/American College of Allergy, Asthma and Immunology Joint Task Force on Practice Parameters GRADE- and Institute of Medicine-based recommendations. *Ann Allergy Asthma Immunol.* 2024;132(3):274-312. doi:10.1016/j.anai.2023.11.00

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