

Nemluvio[®] (nemolizumab-ilto) Prior Authorization Form

Member Name: _____ Date of Birth: _____ Member ID#: _____

Drug Information

Pharmacy billing (NDC: _____) Start Date (or date of next dose): _____

Dose: _____ Regimen: _____

Pharmacy Information

Pharmacy NPI: _____ Pharmacy Name: _____

Pharmacy Phone: _____ Pharmacy Fax: _____

Prescriber Information

Prescriber NPI: _____ Prescriber Name: _____

Prescriber Phone: _____ Prescriber Fax: _____ Specialty: _____

Criteria**For Initial Authorization:****1. Please indicate the diagnosis and information:** Moderate-to-Severe Atopic Dermatitis Prurigo Nodularis (PN) Other: _____

A. Has the member failed 1 medium potency to very-high potency Tier-1 topical corticosteroid?

Yes No

i. If yes, please provide the medication and duration of treatment:

a. Drug: _____ Date of trial: _____

b. Was the trial at least 2 weeks in duration? Yes No ii. If no, is there a contraindication or documented intolerance to medium potency to very-high potency Tier-1 topical corticosteroids? Yes No

a. If yes, please describe: _____

B. Has the member failed 1 topical calcineurin inhibitor [e.g., Elidel[®] (pimecrolimus), Protopic[®] (tacrolimus)]?Yes No

i. If yes, please provide the medication and duration of treatment:

a. Drug: _____ Date of trial: _____

b. Was the trial at least 2 weeks in duration? Yes No

ii. If no, is there a contraindication or documented intolerance to topical calcineurin inhibitors?

Yes No

a. If yes, please describe: _____

C. Will Nemluvio[®] be used concurrently with other biologic medications? Yes No

i. If yes, please provide patient-specific information to support the concurrent use: _____

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Fax completed prior authorization request form to 888-601-8461 or submit Electronic Prior Authorization through CoverMyMeds[®] or SureScripts. All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at AetnaBetterHealth.com/Oklahoma.

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Criteria**For Initial Authorization: (continued)**

2. If diagnosis is **Moderate-to-Severe Atopic Dermatitis**, please provide the following:
- A. Is diagnosis adequately controlled with topical prescription therapies? Yes No
 - B. Does member agree to continue using a topical corticosteroid and/or a topical calcineurin inhibitor in combination with Nemluvio[®] until the disease has sufficiently improved? Yes No
 - C. Member's body surface area (BSA) of atopic dermatitis involvement: _____ Date taken: _____
 - D. Is Nemluvio[®] prescribed by a dermatologist, allergist, or immunologist? Yes No
 - i. If no, has the member been evaluated by a dermatologist, allergist, or immunologist within the last 12 months (or an advanced care practitioner with a supervising physician who is a dermatologist, allergist, or immunologist)? Yes No
 - E. Please provide a patient-specific, clinically significant reason why the member cannot use Adbry[®], Dupixent[®], and Ebgllyss[®]: _____
3. If diagnosis is **Prurigo Nodularis (PN)**, please provide the following:
- A. Member's weight: _____ Date Taken: _____
 - B. Member's Peak Pruritus Numeric Rating Scale (PP-NRS) score: _____
 - C. Does the member have ≥ 20 PN lesions? Yes No
 - D. Have all other causes of pruritus been ruled out? Yes No
 - E. Is Nemluvio[®] prescribed by a dermatologist, allergist, or immunologist? Yes No
 - i. If no, has the member been evaluated by a dermatologist, allergist, or immunologist for PN within the last 12 months (or an advanced care practitioner with a supervising physician who is a dermatologist, allergist, or immunologist)? Yes No
 - F. Please provide a patient-specific, clinically significant reason why the member cannot use Dupixent[®]: _____

For Continued Authorization:

- 1. Date of last dose: _____
- 2. Is the member responding well to treatment with Nemluvio[®]? Yes No
- 3. Has the member experienced any adverse drug reactions related to Nemluvio[®] therapy? Yes No

If yes, please specify adverse reactions: _____

Additional Information: _____

(Page 2 of 2)**Prescriber Signature: _____ Date: _____****I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Failure to complete this form in full will result in processing delays.**

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