



New Hampshire Medicaid Fee-for-Service (FFS) Program
Prior Authorization/Non-Preferred Drug Approval Form
Skin Disorders

PATIENT LAST NAME:

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PATIENT FIRST NAME:

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5. Has the patient been treated with a topical calcineurin inhibitor (e.g., pimecrolimus or tacrolimus) in the past? Yes No

If yes, provide drug name and duration of therapy:

6. Has the patient been treated with a topical phosphodiesterase-4 inhibitor (e.g., crisaborole) Yes No in the past?

If yes, provide drug name and duration of therapy:

7. **Systemic treatment only:** Will the patient also receive therapy with any other monoclonal antibody biologic (e.g., tezepelumab, omalizumab, mepolizumab, reslizumab, dupilumab)? Yes No

8. **Nemlilio only:** Will the patient receive topical corticosteroid and/or topical calcineurin inhibitor therapy until the disease is adequately controlled? Yes No

Other Indications (9–13)

9. Does the patient have a diagnosis of nonsegmental vitiligo? Yes No

10. Does the patient have a diagnosis of prurigo nodularis? Yes No

11. What is the patient's age? _____

12. Is the prescriber a dermatologist, immunologist, or allergist **or** has one been consulted? Yes No

13. Provide any additional information that would help in the decision-making process.

If additional space is needed, please use a separate sheet.



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SECTION IV: NON-PREFERRED DRUG APPROVAL CRITERIA

Chapter 188 of the Laws of 2004 requires that Medicaid only cover non-preferred drugs upon a finding of medical necessity by the prescribing physician. Chapter 188 requires that you base your determination of medical necessity on the following criteria.

Allergic reaction. Describe reaction:

Drug-to-drug interaction. Describe reaction:

Previous episode of an unacceptable side effect or therapeutic failure. Provide clinical information:

Clinical contraindication, co-morbidity, or unique patient circumstance as a contraindication to a preferred drug. Provide clinical information:

Age-specific indications. Provide patient age and explain:

Unique clinical indication supported by FDA approval or peer-reviewed literature. Explain and provide a reference:

Unacceptable clinical risk associated with therapeutic change. Please explain:

I certify that the information provided is accurate and complete to the best of my knowledge and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

PREScriber's SIGNATURE: _____ **DATE:** _____

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