



Pharmacy Prior Approval Request for Monoclonal Antibodies: Dupixent for Prurigo Nodularis

Beneficiary Information

1. Beneficiary Last Name: _____ 2. First Name: _____
3. Beneficiary ID #: _____ 4. Beneficiary Date of Birth: _____ 5. Beneficiary Gender: _____

Prescriber Information

6. Prescribing Provider NPI #: _____
7. Requester Contact Information - Name: _____ Phone #: _____ Ext. _____

Drug Information

8. Drug Name: _____ 9. Strength: _____ 10. Quantity Per 30 Days: _____
11. Length of Therapy (in days): [] up to 30 Days [] 60 Days [] 90 Days [] 120 Days [] 180 Days [] 365 Days [] Other _____

Clinical Information

1. Is the beneficiary age 18 years of age or older? [] Yes [] No
2. Does the beneficiary have a diagnosis of Prurigo Nodularis? [] Yes [] No
3. Has the beneficiary tried and failed, or has contraindication, or intolerance to at least one preferred medium to very high potency topical steroid? [] Yes [] No
4. Is Dupixent being prescribed by or in consultation with a dermatologist, allergist, or immunologist? [] Yes [] No
For continuation of therapy, please answer questions 1-5
5. While on Dupixent, has the beneficiary had continued clinical benefit from baseline supported by medical records? [] Yes [] No
** Please provide medical records documenting the beneficiary's current Prurigo Nodularis status and response to Dupixent treatment**

Signature of Prescriber: _____ Date: _____

(Prescriber Signature Mandatory)

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.