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Section:	Prescription Drugs	**Effective Date:**	July 1, 2026
Subsection:	Topical Products	**Original Policy Date:**	April 7, 2017
Subject:	Dupixent	**Page:**	1 of 25

Last Review Date: June 11, 2026

Dupixent

Description

Dupixent (dupilumab)

Background
Dupixent (dupilumab) is a human monoclonal IgG4 antibody that inhibits interleukin-4 (IL-4) and interleukin-13 (IL-13) signaling by specifically binding to the interleukin-4 receptor alpha (IL-4Rα) subunit shared by the IL-4 and IL-13 receptor complexes. This blocks the IL-4 and IL-13 cytokine-induced inflammatory responses, including the release of proinflammatory cytokines, chemokines, nitric oxide, and IgE; however, the mechanism of action for Dupixent has not been definitively established (1).

Regulatory Status
FDA-approved indications: Dupixent is an interleukin-4 receptor alpha antagonist indicated: (1)

1. Atopic Dermatitis
    a. For the treatment of adult and pediatric patients aged 6 months and older with moderate-to-severe atopic dermatitis (AD) whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. Dupixent can be used with or without topical corticosteroids.
2. Asthma
    a. As an add-on maintenance treatment of adult and pediatric patients aged 6 years and older with moderate-to-severe asthma characterized by an eosinophilic phenotype or with oral corticosteroid dependent asthma.
        i. Limitations of Use: Not for the relief of acute bronchospasm or status asthmaticus.
3. Chronic Rhinosinusitis with Nasal Polyps

   a. As an add-on maintenance treatment in adult and pediatric patients aged 12 years and older with inadequately controlled chronic rhinosinusitis with nasal polyps (CRSwNP).
4. Eosinophilic Esophagitis
   a. For the treatment of adult and pediatric patients aged 1 year and older, weighing at least 15 kg, with eosinophilic esophagitis (EoE).
5. Prurigo Nodularis
   a. For the treatment of adult patients with prurigo nodularis (PN).
6. Chronic Obstructive Pulmonary Disease (COPD)
   a. As an add-on maintenance treatment of adult patients with inadequately controlled chronic obstructive pulmonary disease (COPD) and an eosinophilic phenotype.
      i. Limitations of Use: Not for the relief of acute bronchospasm.
7. Chronic Spontaneous Urticaria (CSU)
   a. For the treatment of adult and pediatric patients aged 2 years and older with chronic spontaneous urticaria (CSU) who remain symptomatic despite H1 antihistamine treatment.
      i. Limitations of Use: Not indicated for other forms of urticaria.
8. Bullous Pemphigoid
   a. For the treatment of adult patients with bullous pemphigoid (BP).
9. Allergic Fungal Rhinosinusitis
   a. For the treatment of adult and pediatric patients aged 6 years and older with allergic fungal rhinosinusitis (AFRS) who have a history of sino-nasal surgery.

Dupixent has warnings for hypersensitivity reactions, conjunctivitis, keratitis, and blepharitis, eosinophilic conditions, psoriasis, arthralgia and psoriatic arthritis, and parasitic infections. Patients should be monitored and Dupixent treatment should be discontinued if appropriate (1).

Patients should not discontinue systemic, topical, or inhaled corticosteroids abruptly upon initiation of Dupixent therapy. Steroids should be reduced gradually, if appropriate (1).

FEP adherence is defined as ≥50% utilization within the last 180 days.

Dupixent is approved to treat both asthma and COPD. Patients with features of both these conditions should follow treatment for asthma, per the Global Initiative for Asthma (GINA) report (2).

The safety and effectiveness of Dupixent in pediatric patients less than 6 months of age with atopic dermatitis have not been established. The safety and effectiveness of Dupixent in

pediatric patients less than 2 years of age with CSU have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 6 years of age with asthma and AFRS have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 1 year of age with eosinophilic esophagitis have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 12 years of age with CRSwNP have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 18 years of age with PN, COPD, and BP have not been established (1).

Related policies
Adbry, Cibinqo, Cinqair, Doxepin cream 5%, Eohilia, Eucrisa, IL-5 Antagonists, Nemluvio, Rinvoq, Tezspire, Xolair

Policy

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

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

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

Prior-Approval Requirements

Age 6 months of age or older

Diagnosis

Patient must have the following:

Moderate-to-severe atopic dermatitis (AD) (eczema)

AND ALL of the following with provided documentation (e.g., medical records, laboratory reports):

1. Inadequate treatment response, intolerance, or contraindication to **ONE** medication in **EACH** of the following categories:
    a. 18 years of age or older:
        a. Topical calcineurin inhibitor (see Appendix 1)
        b. **High** potency topical corticosteroid (see Appendix 2)
    b. 2 to 17 years of age:

a. Topical calcineurin inhibitor (see Appendix 1)
b. Topical corticosteroid (see Appendix 2)

c. 6 months to less than 2 years of age:
a. Topical corticosteroid (see Appendix 2)

2. Baseline evaluation of the condition using **ONE** of the following scoring tools:
a. Investigator's Static Global Assessment (ISGA) with a score ≥ 3 (e.g., *https://www.eczemacouncil.org/assets/docs/Validated-Investigator-Global-Assessment-Scale_vIGA-AD_2017.pdf*)
b. Eczema Area and Severity Index (EASI) with a score ≥ 16 (e.g., *https://dermnetnz.org/topics/easi-score/)*
c. Patient-Oriented Eczema Measure (POEM) with a score ≥ 8 (e.g., *https://jamanetwork.com/data/Journals/DERM/11776/dea40003f1.png*)
d. Scoring Atopic Dermatitis (SCORAD) index with a score ≥ 15 (e.g., *https://dermnetnz.org/topics/scorad/)*
3. **NOT** used in combination with another non-topical Prior Authorization (PA) medication for atopic dermatitis (see Appendix 3)
4. **NOT** given concurrently with live vaccines
5. 12 years of age and older **ONLY**: Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 6 years of age or older

Diagnosis

Patient must have the following:

Moderate-to-severe asthma

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Patient has **ONE** of the following:

a. Asthma with eosinophilic phenotype with eosinophil count greater than or equal to 150 cells/mcL in the past 90 days **OR** 300 cells/mcL in the past 12 months
b. Oral corticosteroid dependent asthma with **ONE** of the following:
    i. 1 month of daily oral corticosteroid use within the last 3 months
    ii. Patient currently requires oral corticosteroids

2. Exacerbation history in the past year of **ONE** of the following:
    a. ≥ 2 moderate asthma exacerbations
    b. ≥ 1 severe asthma exacerbation leading to hospitalization
3. Inadequate control of asthma symptoms after a minimum of 3 months of compliant use with greater than or equal to 50% adherence with **ONE** of the following within the past 6 months:
    a. Inhaled corticosteroids & long acting beta$_2$ agonist
    b. Inhaled corticosteroids & long acting muscarinic antagonist
4. **NOT** used for the emergency relief of acute bronchospasm or status asthmaticus
5. **NO** dual therapy with another monoclonal antibody for the treatment of asthma or COPD (see Appendix 4)
6. **NOT** given concurrently with live vaccines
7. Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 1 year of age or older

Diagnosis

Patient must have the following:

Eosinophilic esophagitis (EoE)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Patient has ≥15 intraepithelial eosinophils per high-power field (eos/hpf)
2. Symptoms of dysphagia (e.g., pain while swallowing, drooling, sensation of food getting stuck in the throat or chest)
3. Inadequate treatment response, intolerance, or contraindication to a proton pump inhibitor (PPI)
4. Patient weight ≥ 15 kg
5. **NOT** given concurrently with live vaccines
6. 11 years of age and older **ONLY**: Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 12 years of age or older

Diagnosis

Patient must have the following:

Chronic rhinosinusitis with nasal polyps (CRSwNP)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Inadequate treatment response, intolerance, or contraindication to a 3-month trial of **EACH** of the following:
    a. **TWO** nasal corticosteroid sprays
    b. **ONE** oral corticosteroid
2. Prescribed by or recommended by an otolaryngologist (ENT)
3. **NO** dual therapy with another monoclonal antibody for the treatment of CRSwNP (see Appendix 5)
4. **NOT** given concurrently with live vaccines
5. 18 years of age and older **ONLY**: Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 18 years of age or older

Diagnosis

Patient must have the following:

Prurigo nodularis (PN)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Inadequate treatment response, intolerance, or contraindication to a **high** potency topical steroid (see Appendix 2)
2. Inadequate treatment response, intolerance, or contraindication to phototherapy **OR** a topical calcineurin inhibitor (see Appendix 1)
3. Baseline evaluation of the condition using the Investigator's Global Assessment (IGA) for prurigo nodularis with a score ≥ 3 (e.g., *https://www.medicaljournals.se/acta/html-editor/table-pdf/big/5947/5947_30402.png*)
3. **NOT** given concurrently with live vaccines

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 18 years of age or older

Diagnosis

Patient must have the following:

Chronic obstructive pulmonary disease (COPD)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Patient has **ONE** of the following
    a. Eosinophil count greater than or equal to 150 cells/mcL in the past 90 days **OR** 300 cells/mcL in the past 12 months
    b. Oral corticosteroid dependent COPD with **ONE** of the following:
        i. 1 month of daily oral corticosteroid use within the last 3 months
        ii. Patient currently requires oral corticosteroids
2. Exacerbation history in the past year of **ONE** of the following:
    a. ≥ 2 moderate COPD exacerbations
    b. ≥ 1 severe COPD exacerbation leading to hospitalization
3. Inadequate control of COPD symptoms after a minimum of 3 months of compliant use with greater than or equal to 50% adherence with **ONE** of the following within the past 6 months:
    a. Long acting beta$_2$ agonist & long acting muscarinic antagonist & inhaled corticosteroid
    b. Long acting beta$_2$ agonist & long acting muscarinic antagonist if inhaled corticosteroids are contraindicated
4. **NOT** used for the emergency relief of acute bronchospasm
5. **NO** dual therapy with another monoclonal antibody for the treatment of asthma or COPD (see Appendix 4)
6. **NOT** given concurrently with live vaccines
7. Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 2 years of age or older

Diagnosis

Patient must have the following:

Chronic spontaneous urticaria (CSU)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Symptomatic after at least **TWO** previous trials of H1-antihistamines
2. Baseline urticaria activity score (UAS)
   (e.g., *https://www.mdcalc.com/urticaria-activity-score-uas*)
3. **NO** dual therapy with another monoclonal antibody for the treatment of CSU (see Appendix 6)
4. **NOT** given concurrently with live vaccines
5. 12 years of age and older **ONLY:** Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 18 years of age or older

Diagnosis

Patient must have the following:

Bullous pemphigoid (BP)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Inadequate treatment response, intolerance, or contraindication to a 3-month trial of **TWO** of the following:
   a. **High** potency topical corticosteroid (see Appendix 2)
   b. Oral corticosteroid
   c. Tetracycline antibiotic or dapsone
2. **NOT** given concurrently with live vaccines

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 6 years of age or older

Diagnosis

Patient must have the following:

Allergic Fungal Rhinosinusitis (AFRS)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Confirmed diagnosis of AFRS by CT scan with characteristic findings of AFRS or positive fungal stain of nasal contents removed during surgery
2. Patient has history of sino-nasal surgery
3. Inadequate treatment response, intolerance, or contraindication to a 3-month trial of **EACH** of the following:
    a. **TWO** nasal corticosteroid sprays
    b. **ONE** oral corticosteroid
4. Prescribed by or recommended by an otolaryngologist (ENT)
5. **NOT** given concurrently with live vaccines

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Prior – Approval *Renewal* Requirements

Age 6 months of age or older

Diagnosis

Patient must have the following:

Atopic dermatitis (AD) (eczema)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Documented improvement of the condition using **ONE** of the following scoring tools:
    a. ISGA – decrease from baseline by at least 2 points (e.g., *https://www.eczemacouncil.org/assets/docs/Validated-Investigator-Global-Assessment-Scale_vIGA-AD_2017.pdf*)
    b. EASI – decrease from baseline by at least 75% (e.g., *https://dermnetnz.org/topics/easi-score/*)
    c. POEM – decrease from baseline by at least 3 points (e.g., *https://jamanetwork.com/data/Journals/DERM/11776/dea40003f1.png*)
    d. SCORAD – decrease from baseline by at least 50% (e.g., *https://dermnetnz.org/topics/scorad/)*
2. Patient has been adherent to Dupixent therapy
3. **NOT** used in combination with another non-topical Prior Authorization (PA) medication for atopic dermatitis (see Appendix 3)
4. **NOT** given concurrently with live vaccines
5. 12 years of age and older **ONLY**: Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 6 years of age or older

Diagnosis

Patient must have the following:

Asthma

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Decreased exacerbations **OR** improvement in symptoms
2. Patient has been adherent to Dupixent therapy

3. **NOT** used for the emergency relief of acute bronchospasm or status asthmaticus
4. **NO** dual therapy with another monoclonal antibody for the treatment of asthma or COPD (see Appendix 4)
5. **NOT** given concurrently with live vaccines
6. Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 1 year of age or older

Diagnosis

Patient must have the following:

Eosinophilic esophagitis (EoE)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. 11 years of age and older **ONLY**: Decrease in intraepithelial eosinophils per high-power field (eos/hpf) from baseline
2. Improvement in symptoms of dysphagia
3. Patient weight ≥ 15 kg
4. Patient has been adherent to Dupixent therapy
5. **NOT** given concurrently with live vaccines
6. 11 years of age and older **ONLY:** Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 12 years of age or older

Diagnosis

Patient must have the following:

Chronic rhinosinusitis with nasal polyps (CRSwNP)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Improvement in sino-nasal symptoms
2. Decreased utilization of oral corticosteroids
3. Patient has been adherent to Dupixent therapy
4. **NO** dual therapy with another monoclonal antibody for the treatment of CRSwNP (see Appendix 5)
5. **NOT** given concurrently with live vaccines
6. 18 years of age and older **ONLY:** Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 18 years of age or older

Diagnosis

Patient must have the following:

Prurigo nodularis (PN)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Documented improvement of the condition using IGA for prurigo nodularis with a decrease from baseline by at least 2 points (e.g., *https://www.medicaljournals.se/acta/html-editor/table-pdf/big/5947/5947_30402.png*)
2. Patient has been adherent to Dupixent therapy
3. **NOT** given concurrently with live vaccines

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 18 years of age or older

Diagnosis

Patient must have the following:

Chronic obstructive pulmonary disease (COPD)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Decreased exacerbations **OR** improvement in symptoms
2. Patient has been adherent to Dupixent therapy
3. **NOT** used for the emergency relief of acute bronchospasm
4. **NO** dual therapy with another monoclonal antibody for the treatment of asthma or COPD (see Appendix 4)
5. **NOT** given concurrently with live vaccines
6. Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 2 years of age or older

Diagnosis

Patient must have the following:

Chronic spontaneous urticaria (CSU)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Decrease in urticaria activity score (UAS), such as improvement in pruritic wheals, hives, and itching
   (e.g., *https://www.mdcalc.com/urticaria-activity-score-uas*)
2. Patient has been adherent to Dupixent therapy
3. **NO** dual therapy with another monoclonal antibody for the treatment of CSU (see Appendix 6)
4. **NOT** given concurrently with live vaccines
5. 12 years of age and older **ONLY:** Patient **MUST** have tried the preferred product(s) (see Appendix 7) if adjudicated through the pharmacy benefit unless the patient has a valid medical exception (e.g., inadequate treatment response, intolerance, contraindication)

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 18 years of age or older

Diagnosis

Patient must have the following:

Bullous pemphigoid (BP)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Improvement in symptoms
2. Patient has been adherent to Dupixent therapy
3. **NOT** given concurrently with live vaccines

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Age 6 years of age or older

Diagnosis

Patient must have the following:

Allergic Fungal Rhinosinusitis (AFRS)

AND ALL of the following with provided documentation (e.g., medical records, laboratory values):

1. Improvement in sino-nasal symptoms
2. Patient has been adherent to Dupixent therapy
3. **NOT** given concurrently with live vaccines

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Policy Guidelines

Pre - PA Allowance

None

Prior - Approval Limits

Quantity

Strength	Diagnosis	Quantity
100 mg	Asthma	12 injections per 168 days **OR**
	Atopic dermatitis	N/A
	Chronic rhinosinusitis with nasal polyps	N/A
	Eosinophilic esophagitis	N/A

	Prurigo nodularis	N/A
	Chronic obstructive pulmonary disease	N/A
	Chronic spontaneous urticaria	N/A
	Bullous pemphigoid	N/A
	Allergic fungal rhinosinusitis	N/A
200 mg	Asthma	14 injections per 168 days **OR**
	Atopic dermatitis	
	Chronic rhinosinusitis with nasal polyps	N/A
	Eosinophilic esophagitis	12 injections per 168 days **OR**
	Prurigo nodularis	N/A
	Chronic obstructive pulmonary disease	N/A
	Chronic spontaneous urticaria	14 injections per 168 days **OR**
	Bullous pemphigoid	N/A
	Allergic fungal rhinosinusitis	12 injections per 168 days **OR**
300 mg	Asthma	14 injections per 168 days **OR**
	Atopic dermatitis	
	Chronic rhinosinusitis with nasal polyps	12 injections per 168 days **OR**
	Eosinophilic esophagitis	24 injections per 168 days **OR**
	Prurigo nodularis	14 injections per 168 days **OR**
	Chronic obstructive pulmonary disease	12 injections per 168 days **OR**
	Chronic spontaneous urticaria	14 injections per 168 days **OR**
	Bullous pemphigoid	14 injections per 168 days **OR**
	Allergic fungal rhinosinusitis	12 injections per 168 days

Duration 24 weeks

Prior – Approval *Renewal* Limits

Quantity

Strength	Diagnosis	Quantity
100 mg	Asthma	6 injections per 84 days **OR**
	Atopic dermatitis	N/A

	Chronic rhinosinusitis with nasal polyps	N/A
	Eosinophilic esophagitis	N/A
	Prurigo nodularis	N/A
	Chronic obstructive pulmonary disease	N/A
	Chronic spontaneous urticaria	N/A
	Bullous pemphigoid	N/A
	Allergic fungal rhinosinusitis	N/A
200 mg	Asthma	6 injections per 84 days **OR**
	Atopic dermatitis	
	Chronic rhinosinusitis with nasal polyps	N/A
	Eosinophilic esophagitis	6 injections per 84 days **OR**
	Prurigo nodularis	N/A
	Chronic obstructive pulmonary disease	N/A
	Chronic spontaneous urticaria	6 injections per 84 days **OR**
	Bullous pemphigoid	N/A
	Allergic fungal rhinosinusitis	6 injections per 84 days **OR**
300 mg	Asthma	6 injections per 84 days **OR**
	Atopic dermatitis	
	Chronic rhinosinusitis with nasal polyps	6 injections per 84 days **OR**
	Eosinophilic esophagitis	12 injections per 84 days **OR**
	Prurigo nodularis	6 injections per 84 days **OR**
	Chronic obstructive pulmonary disease	6 injections per 84 days **OR**
	Chronic spontaneous urticaria	6 injections per 84 days **OR**
	Bullous pemphigoid	6 injections per 84 days **OR**
	Allergic fungal rhinosinusitis	6 injections per 84 days

Duration 12 months

Rationale

Section:	Prescription Drugs	**Effective Date:**	July 1, 2026
Subsection:	Topical Products	**Original Policy Date:**	April 7, 2017
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Summary
Dupixent (dupilumab) is an interleukin-4 receptor alpha antagonist indicated for the treatment of atopic dermatitis (AD), asthma, eosinophilic esophagitis (EoE), chronic rhinosinusitis with nasal polyps (CRSwNP), prurigo nodularis (PN), chronic obstructive pulmonary disease (COPD), chronic spontaneous urticaria (CSU), bullous pemphigoid (PG), and allergic fungal rhinosinusitis. Dupixent has warnings for hypersensitivity reactions, conjunctivitis and keratitis, and parasitic infections. Patients should be monitored and Dupixent treatment should be discontinued if appropriate. The safety and effectiveness of Dupixent in pediatric patients less than 6 months of age with atopic dermatitis have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 2 years of age with CSU have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 6 years of age with asthma and AFRS have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 1 year of age with eosinophilic esophagitis have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 12 years of age with CRSwNP have not been established. The safety and effectiveness of Dupixent in pediatric patients less than 18 years of age with PN, COPD, and BP have not been established (1).

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

References
1. Dupixent [package insert]. Tarrytown, NY: Regeneron Pharmaceuticals Inc.; April 2026.
2. Global Initiative for Asthma. Global Strategy for Asthma Management and Prevention, 2024. Available from www.ginasthma.org.

Policy History

Date	Action
April 2017	Addition to PA Addition of EASI, POEM and SCORAD scoring tools to criteria for evaluation
June 2017	Annual review Addition of Dupixent into the Managed PA program Adjustment of the Baseline POEM and SCORAD values
May 2018	Addition of url links for scoring tools
June 2018	Annual editorial review
November 2018	Annual editorial review and reference update. Addition of asthma indication
March 2019	Decreased age requirement for atopic dermatitis from 18 and older to 12 and older and added 200 mg syringes for atopic dermatitis. Added no live vaccines requirement to asthma indication

June 2019	Annual review. Addition of the 50% adherence requirement to the asthma diagnosis
July 2019	Addition of indication: chronic rhinosinusitis with nasal polyposis (CRSwNP)
September 2019	Annual review
June 2020	Decreased age requirement for atopic dermatitis from 12 and older to 6 and older. Revised t/f steroid requirement for pediatric patients. Scoring tool links updated
September 2020	Annual review and reference update
March 2021	Annual editorial review. Investigator's Static Global Assessment link updated
May 2021	Revised the asthma eosinophil count to include ≥150 cells/mcL in the past 90 days. Updated Appendix 1 and 2
June 2021	Annual review
November 2021	Changed age requirement for asthma to 6 years and older per newest package insert. Added Dupixent 100mg to dosing chart. Revised initiation days supply and duration to accommodate new strength
December 2021	Annual review
January 2022	Changed requirement to t/f of TWO nasal corticosteroids sprays and ONE oral corticosteroid per FEP
March 2022	Annual review and reference update. Per SME: Changed asthma renewal requirement to "decreased exacerbations and/or improvement in symptoms"; Added asthma initiation requirement that patients with eosinophilic asthma must have prior acute exacerbation(s); Added asthma initiation option that patients with corticosteroid dependent asthma may be currently requiring oral corticosteroids.
April 2022	Addition of requirement for atopic dermatitis: "not used in combination with another non-topical PA medication for atopic dermatitis" and added Appendix 3. Added "emergency" to the requirement "not used for the emergency relief of acute bronchospasm or status asthmaticus"
June 2022	Annual review. Addition of indication per PI update: eosinophilic esophagitis. Per PI update, reduced atopic dermatitis age requirement to 6 months and older
September 2022	Annual review
October 2022	Per PI update, addition of indication prurigo nodularis (PN)
November 2022	Per FEP: addition of initiation requirement for CRSwNP, must be prescribed by or recommended by an ENT
December 2022	Annual review
January 2023	Changed Appendix 2 and moved fluradrenolide tape to very high potency
March 2023	Annual review and reference update
February 2024	Per PI update, decreased age requirement for EoE to 1 year and older and weight at least 15 kg. Added 200 mg to quantity chart for EoE
March 2024	Annual review

April 2024	Per FEP, revised CRSwNP initiation requirement to a 3-month trial of two nasal steroids and one oral steroid. Also added t/f of a PPI to EoE diagnosis initiation criteria
June 2024	Annual review and reference update
September 2024	Annual review
October 2024	Per PI update, changed indication to chronic rhinosinusitis with nasal polyps and decreased age to 12 and older. Also added COPD indication. Adjusted dual therapy for Asthma indication to prohibit use with another monoclonal antibody for asthma/COPD. Added Appendix 4 noting monoclonal antibodies for the treatment of Asthma or COPD. Per FEP, added requirement to t/f phototherapy or conventional treatment for prurigo nodularis. Added renewal requirement for patient to be adherent to Dupixent therapy to each indication in renewal criteria.
December 2024	Annual editorial review. Added Appendix 5 and no dual therapy requirement for CRSwNP
March 2025	Annual review
May 2025	Per PI update, added indication of CSU. Added Appendix 6
August 2025	Per PI update, added indication of BP
September 2025	Annual review
December 2025	Annual review. Moved Dupixent to non-preferred. Added documentation requirement for all indications. Added Appendix 7
March 2026	Annual review. Per FEP, for EoE continuation, added that only patients 11 and older need a decrease in intraepithelial eosinophils from baseline and per FEP, increased initiation duration to 24 weeks. Per SME, changed PN requirement from t/f conventional systemic therapy to t/f a topical calcineurin and removed BP requirement to t/f conventional systemic therapy
April 2026	Per PI update, added indication of AFRS
May 2026	Per PI update, decreased age requirement for CSU to 2 years and older
June 2026	Annual review

Keywords

This policy was approved by the FEP® Pharmacy and Medical Policy Committee on June 11, 2026 and is effective on July 1, 2026.

Appendix 1

Relative Potency of Topical Calcineurin Inhibitors		
Drug	**Dosage Form**	**Strength**
Medium Potency		
Tacrolimus	Ointment	0.1%
Low Potency		
Tacrolimus	Ointment	0.03%
Pimecrolimus	Cream	1%

Appendix 2

Relative Potency of Selected Topical Corticosteroids		
Drug	**Dosage Form**	**Strength**
Super-High Potency		
Augmented betamethasone dipropionate	Ointment, Gel, Lotion	0.05%
Clobetasol propionate	Cream, Ointment, Lotion, Solution, Shampoo, Foam, Spray, Gel	0.05%
Fluocinonide	Cream	0.1%
Flurandrenolide	Tape	4 mcg/cm^2
Halobetasol propionate	Lotion, Cream, Ointment, Foam	0.05%
High Potency		
Amcinonide	Cream, Ointment	0.1%
Betamethasone dipropionate	Cream, Ointment	0.05%
Betamethasone valerate	Ointment	0.1%
	Foam	0.12%
Desoximetasone	Cream, Ointment, Spray	0.25%
	Cream, Ointment	0.05%
	Gel	0.05%
Diflorasone diacetate	Cream, Ointment	0.05%
Fluocinonide	Cream, Ointment, Gel, Solution	0.05%
Fluticasone propionate	Ointment	0.1%
Halcinonide	Cream, Ointment, Solution	0.1%
Halobetasol propionate	Lotion	0.01%

Mometasone furoate	Ointment	0.1%
Triamcinolone acetonide	Cream, Ointment	0.5%
Medium Potency		
Betamethasone dipropionate	Spray, Lotion	0.05%
Betamethasone valerate	Cream	0.1%
Clocortolone pivalate	Cream	0.1%
Desonide	Ointment, Gel	0.05%
Fluocinolone acetonide	Cream, Ointment	0.025%
Flurandrenolide	Cream, Ointment, Lotion	0.05%
Fluticasone propionate	Cream	0.05%
Hydrocortisone butyrate	Cream, Lotion, Ointment, Solution	0.1%
Hydrocortisone probutate	Cream	0.1%
Hydrocortisone valerate	Cream, Ointment	0.2%
Mometasone furoate	Cream, Lotion, Solution	0.1%
Prednicarbate	Cream, Ointment	0.1%
Triamcinolone acetonide	Cream, Ointment, Lotion	0.1%
	Ointment	0.05%
	Ointment	0.025%
	Aerosol spray	0.2 mg per 2 second spray
Low Potency		
Alclometasone dipropionate	Cream, Ointment	0.05%
Betamethasone valerate	Lotion	0.1%
Desonide	Cream, Lotion, Foam	0.05%
Fluocinolone acetonide	Cream, Solution, Shampoo, Oil	0.01%
Hydrocortisone	Cream, Ointment, Lotion, Solution	2.5%
	Cream, Gel, Lotion	2%
	Cream, Ointment, Liquid, Lotion, Solution	1%
	Cream, Ointment	0.5%
Hydrocortisone acetate	Cream	2.5%
	Lotion	2%

	Cream, Ointment	1%
Triamcinolone acetonide	Cream, Lotion	0.025%

Appendix 3 - List of Non-Topical PA Medications for Atopic Dermatitis

Generic Name	Brand Name
abrocitinib	Cibinqo
dupilumab	Dupixent
lebrikizumab-lbkz	Ebglyss
nemolizumab-ilto	Nemluvio
tralokinumab-ldrm	Adbry
upadactinib	Rinvoq

Appendix 4 - List of Monoclonal Antibodies for Asthma or COPD

Generic Name	Brand Name
benralizumab	Fasenra
dupilumab	Dupixent
mepolizumab	Nucala
omalizumab	Xolair
reslizumab	Cinqair
tezepelumab-ekko	Tezspire

Appendix 5 - List of Monoclonal Antibodies for CRSwNP

Generic Name	Brand Name
dupilumab	Dupixent
mepolizumab	Nucala
omalizumab	Xolair
tezepelumab-ekko	Tezspire

Appendix 6 - List of Monoclonal Antibodies for CSU

Generic Name	Brand Name
dupilumab	Dupixent
omalizumab	Xolair

Appendix 7 - List of Preferred Products

List of preferred products:
https://info.caremark.com/content/dam/enterprise/caremark/microsites/dig/pdfs/pa-fep/fep-misc/FEP_IndicationMedChx.pdf

Refer to formulary documents for confirmation of coverage:
https://www.fepblue.org/pharmacy/prescriptions#drug-lists