

**I. Requirements for Prior Authorization of Monoclonal Antibodies – Anti-IL, Anti-IgE, Anti-TSLP (MABs – Anti-IL, Anti-IgE, Anti-TSLP)**

**A. Prescriptions That Require Prior Authorization**

All prescriptions for MABs – Anti-IL, Anti-IgE, Anti-TSLP must be prior authorized.

**B. Review of Documentation for Medical Necessity**

In evaluating a request for prior authorization of a prescription for a MAB – Anti-IL, Anti-IgE, Anti-TSLP, the determination of whether the requested prescription is medically necessary will take into account whether the beneficiary:

1. For Dupixent (dupilumab), see the Dupixent (dupilumab) policy; **OR**
2. Is prescribed the MAB – Anti-IL, Anti-IgE, Anti-TSLP for the treatment of a diagnosis that is indicated in the U.S. Food and Drug Administration (FDA)-approved package labeling OR a medically accepted indication; **AND**
3. Is age-appropriate according to FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**
4. Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**
5. Is prescribed the MAB – Anti-IL, Anti-IgE, Anti-TSLP by or in consultation with an appropriate specialist (i.e., pulmonologist, allergist, immunologist, dermatologist, hematologist/oncologist, rheumatologist, etc.); **AND**
6. If currently using a different MAB – Anti-IL, Anti-IgE, Anti-TSLP than requested, will discontinue the other MAB – Anti-IL, Anti-IgE, Anti-TSLP prior to starting the requested agent; **AND**
7. For a non-preferred MAB – Anti-IL, Anti-IgE, Anti-TSLP, **one** of the following:
  - a. **Both** of the following:
    - i. Has a history of therapeutic failure of or a contraindication or an intolerance to the preferred MABs – Anti-IL, Anti-IgE, Anti-TSLP approved or medically accepted for the beneficiary's indication
    - ii. For a non-preferred MAB – Anti-IL, Anti-IgE, Anti-TSLP with a corresponding biosimilar/brand biologic/unbranded biologic that is preferred on the Preferred Drug List (PDL), has a history of therapeutic failure of or a contraindication or an intolerance to the preferred corresponding biosimilar/brand biologic/unbranded biologic that would not be expected to occur with the requested drug
  - b. Has a current history (within the past 90 days) of being prescribed the same non-preferred MAB – Anti-IL, Anti-IgE, Anti-TSLP (does not apply to non-preferred MABs – Anti-IL, Anti-IgE, Anti-TSLP when a corresponding biosimilar/brand biologic/unbranded

See the PDL for the list of preferred MABs – Anti-IL, Anti-IgE, Anti-TSLP at:  
<https://papdl.com/preferred-drug-list>;

**AND**

8. For a diagnosis of asthma, **both** of the following:
- a. Has an asthma severity that is consistent with the FDA-approved indication for the prescribed MAB – Anti-IL, Anti-IgE, Anti-TSLP despite maximal therapeutic doses of or contraindication or intolerance to standard asthma controller drugs based on current national treatment guidelines for the diagnosis and management of asthma
  - b. Will use the requested MAB – Anti-IL, Anti-IgE, Anti-TSLP in addition to standard asthma controller drugs as recommended by current national treatment guidelines for the diagnosis and management of asthma;

**AND**

9. For a diagnosis of chronic spontaneous urticaria, **both** of the following:
- a. Has a history of urticaria for a period of at least six weeks
  - b. **One** of the following:
    - i. Requires systemic steroids to control urticarial symptoms
    - ii. Has a history of therapeutic failure of or a contraindication or an intolerance to maximum tolerated doses of an H1 antihistamine taken for at least two weeks;

**AND**

10. For a diagnosis of eosinophilic granulomatosis with polyangiitis (EGPA), **all** of the following:
- a. Has a diagnosis of EGPA supported by **all** of the following:
    - i. A history of asthma,
    - ii. A history of absolute blood eosinophil count  $\geq 1000$  cells/microL or blood eosinophil level  $>10\%$  of leukocytes,
    - iii. A history of at least **one** of the following:
      - a) Histopathological evidence of **one** of the following:
        - (i) Eosinophilic vasculitis,
        - (ii) Perivascular eosinophilic infiltration,
        - (iii) Eosinophil-rich granulomatous inflammation,
      - b) Neuropathy, mono or poly (motor deficit or nerve conduction abnormality),
      - c) Pulmonary infiltrates, non-fixed,

- d) Sino-nasal abnormality,
  - e) Cardiomyopathy,
  - f) Glomerulonephritis,
  - g) Alveolar hemorrhage,
  - h) Palpable purpura,
  - i) Positive test for ANCA,
- b. **One** of the following:
- i. Requires systemic glucocorticoids to maintain remission
  - ii. Has a contraindication or an intolerance to systemic glucocorticoids,
- c. For a beneficiary with severe EGPA as defined by national treatment guidelines, has a history of therapeutic failure of or a contraindication or an intolerance to rituximab or cyclophosphamide;

**AND**

11. For a diagnosis of hypereosinophilic syndrome (HES), **all** of the following:
- a. Has FIP1L1-PDGFR $\alpha$ -negative HES with organ damage or dysfunction,
  - b. Has a blood eosinophil count  $\geq 1000$  cells/microL,
  - c. **One** of the following:
    - i. Requires or has required systemic glucocorticoids to maintain remission
    - ii. Has a contraindication or an intolerance to systemic glucocorticoids;

**AND**

12. For all other diagnoses, has a history of therapeutic failure of or a contraindication or an intolerance to first line therapy(ies) if applicable according to consensus treatment guidelines; **AND**
13. For Xolair (omalizumab) for a diagnosis of asthma, has a diagnosis of allergen-induced asthma (allergic asthma confirmed by either a positive skin test or radioallergosorbent test) to an unavoidable perennial aeroallergen (e.g., pollen, mold, dust mite, etc.); **AND**
14. For Cinqair (reslizumab) for a diagnosis of asthma with an eosinophilic phenotype, has an absolute blood eosinophil count  $\geq 400$  cells/microL; **AND**
15. For Nucala (mepolizumab) for a diagnosis of asthma, has asthma with an eosinophilic phenotype with absolute blood eosinophil count  $\geq 150$  cells/microL; **AND**
16. For Fasenra (benralizumab), has asthma with an eosinophilic phenotype with absolute blood eosinophil count  $\geq 150$  cells/microL;

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to

meet the medical needs of the beneficiary, the request for prior authorization will be approved.

FOR RENEWALS OF PRIOR AUTHORIZATION FOR MABs – ANTI-IL, ANTI-IgE, ANTI-TSLP: The determination of medical necessity of a request for renewal of a prior authorization for a MAB – Anti-IL, Anti-IgE, Anti-TSLP that was previously approved will take into account whether the beneficiary:

1. Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**
2. Is prescribed a MAB – Anti-IL, Anti-IgE, Anti-TSLP by or in consultation with an appropriate specialist (i.e., pulmonologist, allergist, immunologist, dermatologist, rheumatologist, etc.); **AND**
3. Is not using the requested MAB – Anti-IL, Anti-IgE, Anti-TSLP in combination with another MAB – Anti-IL, Anti-IgE, Anti-TSLP; **AND**
4. For a diagnosis of asthma, **both** of the following:
  - a. Has measurable evidence of improvement in the severity of the asthma condition
  - b. Continues to use the requested MAB – Anti-IL, Anti-IgE, Anti-TSLP in addition to standard asthma controller drugs as recommended by current national treatment guidelines for the diagnosis and management of asthma;

**AND**

5. For a diagnosis of chronic spontaneous urticaria, **both** of the following:
  - a. Experienced improvement of symptoms
  - b. Has a documented rationale for continued use;

**AND**

6. For a diagnosis of HES or EGPA, has **one** of the following:
  - a. Measurable evidence of improvement in disease activity
  - b. Reduction in use of systemic glucocorticoids for this indication;

**AND**

7. For a non-preferred MAB – Anti-IL, Anti-IgE, Anti-TSLP with a corresponding biosimilar/brand biologic/unbranded biologic that is preferred on the PDL, has a history of therapeutic failure of or a contraindication or an intolerance to the preferred corresponding biosimilar/brand biologic/unbranded biologic that would not be expected to occur with the requested drug

See the PDL for the list of preferred MABs – Anti-IL, Anti-IgE, Anti-TSLP at:

<https://papdl.com/preferred-drug-list>;

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

#### C. Clinical Review Process

Prior authorization personnel will review the request for prior authorization and apply the clinical guidelines in Section B. above to assess the medical necessity of a prescription for a MAB – Anti-IL, Anti-IgE, Anti-TSLP. If the guidelines in Section B. are met, the reviewer will prior authorize the prescription. If the guidelines are not met, the prior authorization request will be referred to a physician reviewer for a medical necessity determination. Such a request for prior authorization will be approved when, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary.

## MONOCLONAL ANTIBODIES (MABs) – ANTI-IL, ANTI-IgE, ANTI-TSLP

### PRIOR AUTHORIZATION FORM (form effective 1/9/2023)

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		Total # of pages: _____		Prescriber name:	
Name of office contact:				Specialty:	
Contact's phone number:				NPI:	State license #:
LTC facility contact/phone:				Street address:	
Beneficiary name:				City/state/zip:	
Beneficiary ID#:		DOB:	Phone:	Fax:	

### CLINICAL INFORMATION

Drug requested:	Strength:	Dosage form (pen, vial, etc):
Dose & directions:	Quantity:	Duration: _____ months
Diagnosis:	Dx code ( <i>required</i> ):	Weight: _____ lbs / kg
Has the beneficiary used the requested medication in the past 90 days? <i>Submit documentation.</i>		<input type="checkbox"/> Yes – date of last dose: _____ <input type="checkbox"/> No
Is the requested medication being prescribed by or in consultation with a specialist?		<input type="checkbox"/> Yes <i>Submit documentation of consultation, if applicable.</i> <input type="checkbox"/> No

Complete all sections that apply to the beneficiary and this request. **Check all that apply and submit documentation for each item.**

#### INITIAL requests

**For a non-preferred drug in this class:** Does the beneficiary have a history of trial and failure of or contraindication or an intolerance to the preferred agents in this class that are approved or medically accepted for treatment of the beneficiary's condition? *Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred agents in this class.*

☐ Yes  
☐ No *Submit documentation.*

#### 1. For treatment of ASTHMA:

- ☐ Is currently receiving optimally titrated doses of or has a contraindication or an intolerance to the following (*check all that apply*):
- ☐ inhaled glucocorticoid      ☐ long-acting beta-agonist (LABA)
  - ☐ leukotriene modifier      ☐ other (eg, tiotropium, theophylline): \_\_\_\_\_

#### ☐ For an anti-IgE MAB (eg, XOLAIR):

- ☐ Has moderate-to-severe persistent asthma induced by an unavoidable perennial allergen (pollen, mold, dust mites, etc)
- ☐ Diagnosis confirmed by positive skin test or radioallergosorbent test (RAST)
- ☐ Has a serum total IgE measurement between 30 international units (IU)/mL and 1300 IU/mL

#### ☐ For an anti-IL MAB (eg, CINQAIR, FASENRA, NUCALA):

- ☐ Has asthma of an eosinophilic phenotype – Absolute blood eosinophil count: \_\_\_\_\_/mL Date obtained: \_\_\_\_\_
- ☐ Has severe asthma

#### ☐ For an anti-TSLP (eg, TEZSPIRE):

- ☐ Has severe asthma

#### 2. For treatment of CHRONIC SPONTANEOUS (IDIOPATHIC) URTICARIA:

- ☐ Has a history of urticaria for a period of ≥6 weeks
- ☐ Requires use of systemic steroids to control urticarial symptoms
- ☐ Tried and failed the maximally tolerated dose of an H1 antihistamine (eg, cetirizine/levocetirizine, fexofenadine, loratadine/desloratadine) taken for at least 2 weeks or has a contraindication or an intolerance to H1 antihistamines

### 3. For treatment of EGPA:

- ☐ Has a history of asthma
- ☐ Has an absolute blood eosinophil count  $\geq 1000$ /microliter
- ☐ Has a blood eosinophil level  $> 10\%$  of leukocytes
- ☐ Has evidence of the following (*check all that apply*):
  - ☐ histopathological evidence of:
    - ☐ eosinophilic vasculitis
    - ☐ perivascular eosinophilic infiltration
    - ☐ eosinophil-rich granulomatous inflammation
  - ☐ neuropathy (nerve deficit or conduction abnormality)
  - ☐ pulmonary infiltrates, non-fixed
  - ☐ sino-nasal abnormality
  - ☐ cardiomyopathy
  - ☐ glomerulonephritis
  - ☐ alveolar hemorrhage
  - ☐ palpable purpura
  - ☐ positive test for ANCA
- ☐ Requires systemic glucocorticoids to maintain remission
- ☐ Has a contraindication or an intolerance to systemic glucocorticoids
- ☐ Has severe EGPA as defined by national treatment guidelines
  - ☐ Tried and failed or has a contraindication or an intolerance to rituximab or cyclophosphamide

### 4. For treatment of HYPEREOSINOPHILIC SYNDROME (HES):

- ☐ Has documented FIP1L1-PDGFR $\alpha$ -negative HES
- ☐ Has organ damage or dysfunction
- ☐ Has a blood eosinophil count  $\geq 1000$ /microliter
- ☐ Requires or has required systemic glucocorticoids to maintain remission
  - ☐ Has a contraindication or an intolerance to systemic glucocorticoids

### 5. For treatment of NASAL POLYPS:

- ☐ Has a history of trial and failure of or contraindication or intolerance to nasal corticosteroids
- ☐ For an anti-IgE MAB (eg, XOLAIR):
  - ☐ Has a serum total IgE measurement between 30 international units (IU)/mL and 1500 IU/mL

## RENEWAL requests

### 1. For treatment of ASTHMA:

- ☐ Experienced measurable evidence of improvement in the severity of the asthma condition
- ☐ Will continue to use optimally titrated doses of or has a contraindication or an intolerance to the following (*check all that apply*):
  - ☐ inhaled glucocorticoid
  - ☐ long-acting beta-agonist (LABA)
  - ☐ leukotriene modifier
  - ☐ other (eg, tiotropium, theophylline): \_\_\_\_\_

### 2. For treatment of CHRONIC SPONTANEOUS (IDIOPATHIC) URTICARIA:

- ☐ Experienced an improvement in symptoms
- ☐ Document rationale for continued use: \_\_\_\_\_

### 3. For treatment of EGPA:

- ☐ Experienced measurable evidence of improvement in disease activity
- ☐ Reduction in use of systemic glucocorticoids for the treatment of EGPA

### 4. For treatment of HYPEREOSINOPHILIC SYNDROME (HES):

- ☐ Experienced measurable improvement in disease activity
- ☐ Reduction in use of systemic glucocorticoids for the treatment of HES

## PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION

Prescriber Signature:

Date:

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