

Effective: November 12, 2024

Guideline Type	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Non-Formulary <input type="checkbox"/> Step-Therapy <input type="checkbox"/> Administrative
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Applies to:

Commercial Products

- Harvard Pilgrim Health Care Commercial products; Fax: 617-673-0988
- Tufts Health Plan Commercial products; Fax: 617-673-0988
CareLinkSM – Refer to CareLink Procedures, Services and Items Requiring Prior Authorization

Public Plans Products

- Tufts Health Direct – A Massachusetts Qualified Health Plan (QHP) (a commercial product); Fax: 617-673-0988

Note: While you may not be the provider responsible for obtaining prior authorization, as a condition of payment you will need to ensure that prior authorization has been obtained.

Overview

Food and Drug Administration – Approved Indications

Enbrel (etanercept) is a tumor necrosis factor blocker indicated for:

Disease State	
Ankylosing Spondylitis	X
Juvenile Idiopathic Arthritis	X
Plaque Psoriasis	X
Psoriatic Arthritis	X
Rheumatoid Arthritis	X

Clinical Guideline Coverage Criteria

The plan may authorization coverage of Enbrel for Members when **ALL** of the following criteria are met:

Ankylosing Spondylitis

1. Documented diagnosis of ankylosing spondylitis
- AND**
2. Patient is at least 18 years of age
- AND**
3. Prescribed by or in consultation with a rheumatologist
- AND**
4. Documentation of **one (1)** of the following:
 - a. Inadequate response or adverse reaction to a prescription strength non-steroidal anti-inflammatory drug (e.g., celecoxib, diclofenac, ibuprofen, naproxen, meloxicam)
 - b. Contraindication to non-steroidal anti-inflammatory drugs
 - c. Previous treatment with a biologic agent indicated for the requested use
 - d. The patient is new to the plan and stable on the requested agent prior to enrollment

Juvenile Idiopathic Arthritis

1. Documented diagnosis of juvenile idiopathic arthritis
AND
2. Patient is at least 2 years of age
AND
3. Prescribed by or in consultation with a rheumatologist
AND
4. Documentation of **one (1)** of the following:
 - a. Inadequate response or adverse reaction to one disease modifying antirheumatic drug (e.g., methotrexate, leflunomide, hydroxychloroquine, sulfasalazine)
 - b. Contraindication to all traditional disease modifying antirheumatic drugs
 - c. Previous treatment with a biologic agent indicated for the requested use
 - d. The patient is new to the plan and has been stable on the requested agent prior to enrollment

Plaque Psoriasis

1. Documented diagnosis of plaque psoriasis
AND
2. Patient is at least 4 years of age
AND
3. Prescribed by or in consultation with a dermatologist
AND
4. Documentation of **one (1)** of the following:
 - a. Inadequate response to one of the following topical therapies: corticosteroids, vitamin D analogs, tazarotene, calcineurin inhibitors, anthralin, coal tar
 - b. Contraindication to all of the following topical therapies: corticosteroids, vitamin D analogs, tazarotene, calcineurin inhibitors, anthralin, and coal tar
 - c. Previous treatment with a biologic agent indicated for the requested use
 - d. The patient is new to the plan and has been stable on the requested agent prior to enrollment

Psoriatic Arthritis

1. Documented diagnosis of psoriatic arthritis
AND
2. Patient is at least 2 years of age
AND
3. Prescribed by or consultation with a rheumatologist or dermatologist

Rheumatoid Arthritis

1. Documented diagnosis of rheumatoid arthritis
AND
2. Patient is at least 18 years of age
AND
3. Prescribed by or in consultation with a rheumatologist
AND
4. Documentation of **one (1)** of the following:
 - a. Inadequate response or adverse reaction to one disease modifying antirheumatic drug (e.g., methotrexate, leflunomide, hydroxychloroquine, sulfasalazine)
 - b. Contraindication to all traditional disease modifying antirheumatic drugs
 - c. Previous treatment with a biologic agent indicated for the requested use
 - d. The patient is new to the plan and has been stable on the requested agent prior to enrollment

Limitations

1. Samples, free goods, or similar offerings of the requested medication do not qualify for an established clinical response and will not be considered for prior authorization.
 2. Documentation of a Member being a social drinker does not qualify as a medically acceptable contraindication or clinical inappropriateness to methotrexate therapy.
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Codes

None

References

1. Enbrel (etanercept) [prescribing information]. Thousand Oaks, CA: Immunex Corporation; September 2024.
 2. Braun J, Davis J et al. First update of the international ASAS consensus statement for the use of anti-TNF agents in patients with Ankylosing Spondylitis. *Ann Rheum Dis*. 2006; 65(3):316-20.
 3. Smith CH, Anstey AV, et al. British association of dermatologists' guidelines for use of biological interventions in psoriasis 2005. *Br J Dermatol* 2005; 153:486-497.
 4. Menter A, Strober BE, Kaplan DH, et al. Joint AAD-NPF guidelines of care for the management and treatment of psoriasis with biologics. *J Am Acad Dermatol* 2019;80:1029-72. HPHC Pharmacy PA Policy: Page 7 of 7.
 5. Fraenkel L, Bathon JM, England BR, et al. 2021 American College of Rheumatology Guideline for the Treatment of Rheumatoid Arthritis. *Arthritis Care Res*. 2021;73(7):924-939.
 6. Singh JA, Guyatt G, Ogdie A, et al. 2018 American College of Rheumatology/National Psoriasis Foundation guideline for the treatment of psoriatic arthritis. *Arthritis Rheumatol*. 2019;71(1):5-32.
 7. Ward MM, Deodhar A, Gensler LS, et al. 2019 Update of the American College of Rheumatology/Spondylitis Association of America/spondyloarthritis research and treatment network recommendations for the treatment of ankylosing spondylitis and nonradiographic axial spondyloarthritis. *Arthritis Rheumatol*. 2019;71(10):1599-1613.
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Approval And Revision History

September 13, 2022: Reviewed by the Pharmacy & Therapeutics Committee.

Subsequent endorsement date(s) and changes made:

- September 2022 e-vote: Effective January 1, 2023, Enbrel added as a preferred agent.
 - December 12, 2023: Expanded age requirements for psoriatic arthritis to 2 years of age based on supplemental indication (eff 1/1/2024).
 - November 12, 2024: No changes (eff 11/12/24)
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Background, Product and Disclaimer Information

Pharmacy Medical Necessity Guidelines have been developed for determining coverage for plan benefits and are published to provide a better understanding of the basis upon which coverage decisions are made. The plan makes coverage decisions on a case-by-case basis considering the individual member's health care needs. Pharmacy Medical Necessity Guidelines are developed for selected therapeutic classes or drugs found to be safe, but proven to be effective in a limited, defined population of patients or clinical circumstances. They include concise clinical coverage criteria based on current literature review, consultation with practicing physicians in the service area who are medical experts in the particular field, FDA and other government agency policies, and standards adopted by national accreditation organizations. The plan revises and updates Pharmacy Medical Necessity Guidelines annually, or more frequently if new evidence becomes available that suggests needed revisions.

For self-insured plans, coverage may vary depending on the terms of the benefit document. If a discrepancy exists between a Pharmacy Medical Necessity Guideline and a self-insured Member's benefit document, the provisions of the benefit document will govern.

Treating providers are solely responsible for the medical advice and treatment of members. The use of this policy is not a guarantee of payment or a final prediction of how specific claim(s) will be adjudicated. Claims payment is subject to member eligibility and benefits on the date of service, coordination of benefits, referral/authorization and utilization management guidelines when applicable, and adherence to plan policies and procedures and claims editing logic.