

**Pharmacy Policy**

**Cosentyx (secukinumab)**

**Policy Number:** 9.117

**Revision Number:** R0

**Version Effective Date:** 1/1/2021

<p>Product Applicability <input type="checkbox"/> All Plan+ Products</p>	
<p><b>Well Sense Health Plan</b></p> <p><input checked="" type="checkbox"/> New Hampshire Medicaid</p>	<p><b>Boston Medical Center HealthNet Plan</b></p> <p><input type="checkbox"/> MassHealth - MCO</p> <p><input type="checkbox"/> MassHealth - ACO</p> <p><input type="checkbox"/> Qualified Health Plans/ConnectorCare/Employer Choice Direct</p> <p><input type="checkbox"/> Senior Care Options</p>

Note: Disclaimer and audit information is located at the end of this document.

**Prior Authorization Policy**

**Products Affected:**

- Cosentyx (secukinumab)

The Plan may authorize coverage of the above products for members meeting the following criteria:

<b>Covered Use</b>	All FDA approved indications not otherwise excluded
<b>Exclusion Criteria</b>	None
<b>Required Medical Information</b>	<ol style="list-style-type: none"> <li>1. A diagnosis of Ankylosing Spondylitis (AS); <b>AND</b> <ol style="list-style-type: none"> <li>a. An inadequate response or contraindication to at least one formulary NSAID</li> </ol> </li> <li>2. Diagnosis of moderate to severe plaque psoriasis (Ps); <b>AND</b>                      An inadequate response, or adverse reaction to two conventional therapies in any one of the following combinations (please note: these combinations DO NOT have to be used concurrently):                     <ol style="list-style-type: none"> <li>1. One topical agent plus one systemic agent; <b>OR</b></li> </ol> </li> </ol>

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	<ol style="list-style-type: none"> <li>2. One topical agent plus one phototherapy; <b>OR</b></li> <li>3. One systemic agent plus one phototherapy; <b>OR</b></li> <li>4. Two systemic agents; <b>OR</b></li> </ol> <ol style="list-style-type: none"> <li>b. A contraindication to all conventional therapies (topical agents, phototherapy, and systemic agents)</li> <li>c. .</li> </ol> <ol style="list-style-type: none"> <li>3. A diagnosis of Psoriatic Arthritis (PsA); <b>AND</b> <ol style="list-style-type: none"> <li>a. An inadequate response or intolerance to at least one formulary non-biologic DMARD for at least three months</li> <li>b.</li> </ol> </li> </ol>
<b>Age Restrictions</b>	18 years of age or older
<b>Prescriber Restriction</b>	AS: Prescribed by or in consultation with a rheumatologist PS: Prescribed by or in consultation with a dermatologist PsA: Prescribed by or in consultation with a dermatologist or rheumatologist
<b>Coverage Duration</b>	12 months
<b>Quantity Limit</b>	AS: 5 pens/syringes/vials for the first 28 days, then 2 pens /syringes/vials per 28 days thereafter Ps: 10 pens/syringes/vials for the first 28 days, then 2 pens/syringes/vials per 28 days thereafter PSA:5 pens/syringes/vials for the first 28 days, then 2 pens/syringes/vials per 28 days thereafter
<b>Other criteria</b>	Reauthorization: <ol style="list-style-type: none"> <li>1. Clinical condition has improved or stabilized</li> </ol>

### Applicable Coding:

None

### Clinical Background Information and References

1. Baeten D, Baraliakos X, Braun J, et al. Anti-interleukin-17A monoclonal antibody secukinumab in treatment of ankylosing spondylitis: a randomised, double-blind, placebo-controlled trial. *Lancet*. 2013 Nov 23;382(9906):1705-13.
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8. Enbrel prescribing information. Thousand Oaks, CA: Amgen Inc. and Pfizer Inc.; 2015 March.
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17. Langley RG, Elewski BE, Lebwohl M et al. Secukinumab in plaque psoriasis--results of two phase 3 trials. *N Engl J Med*. 2014;371(4):326-38.
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19. Lowes MA, Suárez-Fariñas M, Krueger JG. Immunology of psoriasis. *Annu Rev Immunol*. 2014;32:227-55.
20. Mason J, Mason AR, Cork MJ. Topical preparations for the treatment of psoriasis: a systematic review. *Br J Dermatol*. 2002; 146(3):351-64.
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Original Approval Date	Original Effective Date	Policy Owner	Approved by
12/1/2020	1/1/2021	Pharmacy Services	Pharmacy & Therapeutics (P&T) Committee, NH DHHS

Policy Revisions History			
Review Date	Summary of Revisions	Revision Effective Date	Approved by
12/1/2020	9.180 Cosentyx Policy retired, new policy created. Removed compliance requirement, update quantity limits to match labeling change, removed trial/fail requirements for Enbrel/Humira since	1/1/2021	P&T Committee, NH DHHS

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## Policy Revisions History

	Cosentyx/Enbrel/Humira Preferred on NH PDL		
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## Next Review Date

2021

## Other Applicable Policies

## Reference to Applicable Laws and Regulations, If Any

### Disclaimer Information

Medical Policies are the Plan's guidelines for determining the medical necessity of certain services or supplies for purposes of determining coverage. These Policies may also describe when a service or supply is considered experimental or investigational, or cosmetic. In making coverage decisions, the Plan uses these guidelines and other Plan Policies, as well as the Member's benefit document, and when appropriate, coordinates with the Member's health care Providers to consider the individual Member's health care needs.

Plan Policies are developed in accordance with applicable state and federal laws and regulations, and accrediting organization standards (including NCQA). Medical Policies are also developed, as appropriate, with consideration of the medical necessity definitions in various Plan products, review of current literature, consultation with practicing Providers in the Plan's service area who are medical experts in the particular field, and adherence to FDA and other government agency policies. Applicable state or federal mandates, as well as the Member's benefit document, take precedence over these guidelines. Policies are reviewed and updated on an annual basis, or more frequently as needed. Treating providers are solely responsible for the medical advice and treatment of Members.

The use of this Policy is neither a guarantee of payment nor a final prediction of how a specific claim(s) will be adjudicated. Reimbursement is based on many factors, including member eligibility and benefits on the date of service; medical necessity; utilization management guidelines (when applicable); coordination of benefits; adherence with applicable Plan policies and procedures; clinical coding criteria; claim editing logic; and the applicable Plan – Provider agreement.

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