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Commercial PA Criteria Effective: January 1, 2022

Prior Authorization: Ilumya [™]

Products Affected: Ilumya (tildrakizumab-asmn) for subcutaneous injection

<u>Medication Description</u>: Ilumya, an interleukin (IL)-23 blocker, is indicated for the treatment of adults with moderate to severe plaque psoriasis who are candidates for systemic therapy or phototherapy. It is administered subcutaneously at Weeks 0 and 4 and then once every 12 weeks thereafter

Covered Uses: Adults with moderate to severe plaque psoriasis who are candidates for systemic therapy or phototherapy

Exclusion Criteria:

1. Concurrent Use with other Biologics or with Targeted Synthetic Disease-Modifying Antirheumatic Drugs (DMARDs).

Required Medical Information:

- 1. Diagnosis
- 2. Concurrent Medications
- 3. Other therapies tried

Prescriber Restriction: The medication is prescribed by or in consultation with a dermatologist.

Age Restriction: 18 years and older

Coverage Duration:

Initial: 3 Months Conitnuation: 1 year 12 Months

Other Criteria:

I. Initial Criteria

1. Plaque Psoriasis

Initial Therapy: Approve if the patient meets the following criteria

A. Patient has tried at least ONE traditional systemic agent for psoriasis for at least 3 months unless intolerant; OR Note: Examples include methotrexate, cyclosporine, acitretin, or psoralen plus ultraviolet A light (PUVA). An exception to the requirement for a trial of one traditional systemic agent for psoriasis can be made if the patient has already had a 3-month trial or previous intolerance to at least one biologic other than the requested medication. Women of childbearing age may be given special consideration for approval without systemic therapy when topical and phototherapy options have been tried and failed. A biosimilar

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of the requested biologic does not count. A patient who has already tried a biologic for psoriasis is not required to "step back" and try a traditional systemic agent for psoriasis.

- B. Patient has a contraindication to methotrexate, as determined by the prescriber; AND
- C. The medication is prescribed by or in consultation with a dermatologist AND
- D. Patient has a documented failure of, or intolerance to, **TWO** of the following medications *Note: A trial of multiple adalimumab products counts as ONE product.*

Plaque Psoriasis (TWO of the following)				
Enbrel				
Adalimumab Product				
Otezla				
Skyrizi				
Stelara SC				
Taltz				
Tremfya				

II. Continuation Criteria

1. Plaque Psoriasis

- A. Patient has experienced a clinical response as determined by the prescribing physician; AND
- B. Patient has not experienced unacceptable toxicity from the drug.

References:

1. Ilumya[™] injection [prescribing information]. Whitehouse Station, NJ: Sun Pharmaceuticals; October 2019.

Policy Revision history

Rev #	Type of Change	Summary of Change	Sections Affected	Date
1	New Policy	New Policy	All	12/8/2021
2	Update	Added Step through TWO preferred products (Enbrel, adalimumab, Otezla, Skyrizi SC, Stelara SC, Taltz or Tremfya) for Initial Criteria for Psoriasis	Other Criteria	05/11/2023
3	Update	Coverage duration updated to include initial 3 month therapy. Added Note: A trial of multiple adalimumab products counts as ONE product.	Coverage duration	12/20/23

