



Ilumya™ (tildrakizumab-asmn)

When requesting Ilumya (tildrakizumab-asmn), the individual requiring treatment must be diagnosed with an FDA-approved indication and meet the specific coverage guidelines and applicable safety criteria for the covered indication.

FDA-approved Indication

Ilumya is indicated for the treatment of adults with moderate-to-severe plaque psoriasis who are candidates for systemic therapy or phototherapy.

Coverage Guidelines

Plaque psoriasis

The individual must meet **all** of the following criteria for initial authorization:

- Had a 3-month trial of or intolerance to at least one biologic or traditional systemic agent for psoriasis (e.g., adalimumab, certolizumab, etanercept, infliximab, methotrexate, cyclosporine, acitretin, psoralen plus ultraviolet A light); OR
- Has a contraindication to methotrexate therapy; AND
- Is 18 years of age or older; AND
- Ilumya is prescribed by or in consultation with a dermatologist.

For reauthorization, the individual must show response to Ilumya therapy.

Approval duration (initial): 3 months

Approval duration (renewal): 12 months

Dosing Recommendation

The recommended dose is 100 mg administered by subcutaneous injection at weeks 0, 4, and not more frequently than once every 12 weeks thereafter.

References

1. Ilumya™ injection [prescribing information]. Whitehouse Station, NJ: Sun Pharmaceuticals; July 2020.
2. 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(4):1029-1072.
3. Nast A, Gisondi P, Ormerod AD, et al. European S3-Guidelines on the systemic treatment of psoriasis vulgaris – Update 2015 – Short version – EDF in cooperation with EADV and IPC. J Eur Acad Dermatol Venereol. 2015;29(12):2277-2294.
4. Reich K, Papp KA, Blauvelt A, et al. Tildrakizumab versus placebo or etanercept for chronic plaque psoriasis (reSURFACE 1 and reSURFACE 2): results from two randomised controlled, phase 3 trials. Lancet. 2017;390(10091):276-288.