

Leqselvi (deuruxilitinib)
Effective 05/01/2026

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|------------------------------|--|---------------------|---|
| Plan | <input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange | Program Type | <input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy |
| Benefit | <input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit | | |
| Specialty Limitations | This medication has been designated specialty and must be filled at a contracted specialty pharmacy. | | |
| Contact Information | Medical Benefit | Phone: 833-895-2611 | Fax: 888-656-6671 |
| | Pharmacy Benefit | Phone: 800-711-4555 | Fax: 844-403-1029 |
| Exceptions | N/A | | |

Overview

Leqselvi (deuruxilitinib) is a Janus kinase (JAK) inhibitor indicated for the treatment of adults with severe alopecia areata.

Leqselvi is not recommended for use in combination with other JAK inhibitors, biologic immunomodulators, cyclosporine, or other potent immunosuppressants.

Coverage Guidelines

If member is new to the plan (as evidenced by coverage effective date of less than or equal to 90 days), submission of medical records documenting that the member is currently receiving treatment with the requested drug, excluding when the product is obtained as samples or via manufacturer's patient assistance programs

OR

Authorization may be granted when all the following criteria are met:

1. Member is 18 years of age or older
2. Diagnosis of severe alopecia areata
3. Member has at least 50% scalp hair loss (e.g., Severity of Alopecia Tool (SALT) score of 50 or higher)
4. Other forms of alopecia have been ruled out (e.g., androgenetic alopecia, trichotillomania, telogen effluvium, chemotherapy-induced hair loss, tinea capitis)
5. Trial and failure, contraindication, or intolerance to one of the following:
 - a. Litfulo (ritlecitinib)
 - b. Olumiant (baricitinib)

Continuation of Therapy

Requests for reauthorization will be approved when all of the following criteria are met:

1. Submission of medical records (e.g., chart notes) documenting improvement in signs and symptoms of alopecia areata from baseline (e.g., increased hair on scalp, eyebrows, eyelashes)

Limitations

1. Initial approvals will be granted for 36 weeks
2. Reauthorizations will be granted for 12 months
3. The following quantity limits apply:

| Drug Name and Dosage Form | Quantity Limit |
|---------------------------|-------------------|
| Leqselvi tablet | 2 tablets per day |

References

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2. Eisman S, Sinclair R. Ritlecitinib: an investigational drug for the treatment of moderate to severe alopecia areata. *Expert Opin Investig Drugs*. 2021;30(12):1169-1174.
3. Gilhar A, Etzioni A, Paus R. Alopecia areata. *N Engl J Med*. 2012;366(16):1515-1525.
4. Gupta AK, Wang T, Polla Ravi S, et al. Systematic review of newer agents for the management of alopecia areata in adults: Janus kinase inhibitors, biologics and phosphodiesterase-4 inhibitors. *J Eur Acad Dermatol Venereol*. 2023;37(4):666-679.
5. King B, Senna MM, Mesinkovska NA, et al. Efficacy and safety of deuruxolitinib, an oral selective Janus kinase inhibitor, in adults with alopecia areata: results from the phase 3 randomized, controlled trial (THRIVE-AA1). *J Am Acad Dermatol*. 2024 Jul 23:S0190-9622(24)02550-7.
6. Leqselvi [package insert], Cranbury, NJ: Sun Pharmaceutical Industries, Inc.; July 2024.
7. Liu M, Gao Y, Yuan Y, et al. Janus kinase inhibitors for alopecia areata: a systematic review and meta-analysis. *JAMA Netw Open*. 2023;6(6):e2320351.
8. Meah N, Wall D, York K, et al. The Alopecia Areata Consensus of Experts (ACE) study: Results of an international expert opinion on treatments for alopecia areata. *J Am Acad Dermatol*. 2020;83(1):123-130.
9. Mostaghimi A, Gao W, Ray M, et al. Trends in prevalence and incidence of alopecia areata, alopecia totalis, and alopecia universalis among adults and children in a US employer-sponsored insured population. *JAMA Dermatol*. 2023;159(4):411-418.
10. National Alopecia Areata Foundation. Related conditions. <https://www.naaf.org/alopecia-areata/related-conditions/>. 2024. Accessed October 7, 2024.
11. Ramírez-Marín HA, Tosti A. Evaluating the therapeutic potential of ritlecitinib for the treatment of alopecia areata. *Drug Des Devel Ther*. 2022;16:363-374.

Review History

09/10/2025 – Created and reviewed at September P&T. Effective 12/1/2025.

03/11/2026 – Reviewed and updated for March P&T. Administrative update - changing verbiage in reauthorization criteria from “documentation is submitted” to “submission of medical records (e.g., chart notes...” and updating language for members who are new to the Plan. Effective 05/01/2026.

