



Provider Help Desk
1 (844) 236-1464

Request for Prior Authorization
Tralokinumab-ldrm (Adbry)
(PLEASE PRINT - ACCURACY IS IMPORTANT)

FAX Completed Form To
1 (877) 733-3195

Form with fields for IA Medicaid Member ID #, Patient name, DOB, Patient address, Provider NPI, Prescriber name, Phone, Prescriber address, Fax, Pharmacy name, Address, Phone, Pharmacy NPI, Pharmacy fax, NDC.

Prior authorization (PA) is required for tralokinumab-ldrm (Adbry). Requests for non-preferred agents may be considered when documented evidence is provided that the use of preferred agent(s) would be medically contraindicated.

- 1. Request adheres to all FDA approved labeling for requested drug and indication, including age, dosing, contraindications, warning and precautions, drug interactions, and use in specific populations; and
2. Patient has a diagnosis of moderate to severe atopic dermatitis; and
3. Is prescribed by or in consultation with a dermatologist; and
4. Patient has failed to respond to good skin care and regular use of emollients; and
5. Patient has documentation of an adequate trial and therapy failure with at least one preferred medium to high potency topical corticosteroid for a minimum of 2 consecutive weeks; and
6. Patient has documentation of a previous trial and therapy failure with a topical immunomodulator for a minimum of 4 weeks; and
7. Patient has documentation of a previous trial and therapy failure with cyclosporine or azathioprine; and
8. Patient will continue with skin care regimen and regular use of emollients.

If criteria for coverage are met, initial authorization will be given for 16 weeks to assess the response to therapy. Request for continuation of therapy will require documentation of a positive response to therapy and documentation patient will continue with skin care regimen and regular use of emollients. The required trials may be overridden when documented evidence is provided that use of these agents would be medically contraindicated.

Non-Preferred

Adbry

Strength Usage Instructions Quantity Day's Supply

Diagnosis:

Prescriber Specialty: Dermatologist Other (specify):

If other, note consultation with dermatologist: Consultation date:

Physician name, specialty & phone:

Has patient failed to respond to good skin care and regular use of emollients? Yes No

Will patient continue with skin care regimen and regular use of emollients? Yes Emollient to be used: No

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Preferred Medium to High Potency Topical Corticosteroid Trial:

Drug name & dose: _____ Trial dates: _____

Failure reason: _____

Preferred Topical Immunomodulator Trial:

Drug name & dose: _____ Trial dates: _____

Failure reason: _____

Cyclosporine or Azathioprine Trial:

Drug name & dose: _____ Trial dates: _____

Failure reason: _____

Requests for continuation therapy:

Does patient have a documented positive response to therapy?

Yes (describe): _____

No

Will patient continue with skin care regimen and regular use of emollients?

Yes Emollient to be used: _____ No

Medical or contraindication reason to override trial requirements: _____

Attach lab results and other documentation as necessary.

Prescriber signature (Must match prescriber listed above.)	Date of submission
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IMPORTANT NOTE: In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member continues to be eligible for Medicaid.