PRIOR AUTHORIZATION CRITERIA

| DRUG CLASS | TOPICAL RETINOIDS |
|--|------------------------------|
| BRAND NAME (generic) | |
| (30000) | ALTRENO |
| | (tretinoin) |
| | ATRALIN |
| | (tretinoin) |
| | ΑΥΙΤΑ |
| | (tretinoin) |
| | RETIN-A |
| | (tretinoin) |
| | RETIN-A MICRO |
| | (tretinoin) |
| | TWYNEO |
| | (tretinoin/benzoyl peroxide) |
| | VELTIN |
| | (clindamycin/tretinoin) |
| | ZIANA |
| | (clindamycin/tretinoin) |
| | |
| Status: CVS Caremark [®] Criteria | |
| Type: Initial Prior Authorization | |

POLICY

FDA-APPROVED INDICATIONS

Atralin, Avita, Retin-A

Atralin, Avita, and Retin-A are indicated for topical treatment of acne vulgaris. The safety and efficacy of this product in the treatment of other disorders have not been established.

Altreno (tretinoin) lotion 0.05%, Twyneo

Altreno (tretinoin) lotion 0.05% and Twyneo are indicated for the topical treatment of acne vulgaris in patients 9 years of age and older.

Tretinoins (Topical) PA Policy 355-A, 237-A UDR 08-2023

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Retin-A Micro

Retin-A Micro is indicated for topical application in the treatment of acne vulgaris.

Veltin, Ziana

Veltin gel 1.2%/0.025% and Ziana gel are indicated for the topical treatment of acne vulgaris in patients 12 years and older.

Compendial Uses

Keratosis follicularis (Darier's disease, Darier-White disease) 12,15-17

COVERAGE CRITERIA

The requested drug will be covered with prior authorization when the following criteria are met:

- The patient has a diagnosis of acne vulgaris
 - AND
 - The request is NOT for continuation of therapy
 - OR
 - The request is for continuation of therapy
 - AND
 - The patient has achieved or maintained a positive clinical response as evidenced by improvement (e.g., reduction in number of lesions, etc.)

OR

- The patient has a diagnosis of keratosis follicularis (Darier's disease, Darier-White disease) AND
 - The request is NOT for continuation of therapy
 - OR
 - \circ The request is for continuation of therapy
 - AND
- The patient has achieved or maintained a positive clinical response as evidenced by improvement

Duration of Approval (DOA):

- 355-A: Initial therapy DOA: 4 months; Continuation of therapy DOA: 36 months
- 237-A: Initial therapy DOA: 4 months; Continuation of therapy DOA: 12 months

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