BRAND NAME (generic)

EUCRISA (crisaborole)

Status: CVS Caremark[®] Criteria Type: Initial Prior Authorization with Quantity Limit

POLICY

FDA-APPROVED INDICATIONS

Eucrisa is indicated for topical treatment of mild to moderate atopic dermatitis in adult and pediatric patients 3 months of age and older.

COVERAGE CRITERIA

Atopic Dermatitis

Authorization may be granted when the requested drug is being prescribed for mild to moderate atopic dermatitis when ALL of the following criteria are met:

- The patient is 3 months of age or older
- The patient meets ONE of the following criteria:
 - The patient is less than 2 years of age
 - The requested drug will be used on sensitive skin areas (e.g., face, genitals, or skin folds) and the following criteria is met:
 - The patient experienced an inadequate treatment response, intolerance, or contraindication to a topical calcineurin inhibitor
 - The patient experienced an inadequate treatment response, intolerance, or contraindication to a topical calcineurin inhibitor AND a medium or higher potency topical corticosteroid
- If additional quantities are being requested, then 5 percent or greater body surface area is affected

CONTINUATION OF THERAPY

Atopic Dermatitis

Authorization may be granted when the requested drug is being prescribed for mild to moderate atopic dermatitis when ALL of the following criteria are met:

- The patient is 3 months of age or older
- The patient has achieved or maintained a positive clinical response as evidenced by improvement [(e.g., improvement in or resolution of any of the following signs and symptoms: erythema (redness), edema (swelling), xerosis (dry skin), erosions, excoriations (evidence of scratching), oozing and crusting, lichenification (epidermal thickening), OR pruritus (itching)]
- If additional quantities are being requested, then 5 percent or greater body surface area is affected

QUANTITY LIMITS APPLY

60 grams per 25 days* or 180 grams per 75 days* Greater than 5% BSA: 120 grams per 25 days* or 360 grams per 75 days* *The duration of 25 days is used for a 30-day fill period and 75 days is used for a 90-day fill period to allow time for refill processing.

DURATION OF APPROVAL (DOA)

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• 1565-C: Initial therapy DOA: 3 months; Continuation of therapy DOA: 12 months

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