Iowa Department of Human Services

FAX Completed Form To
1 (800) 574-2515

Provider Help Desk
1 (877) 776-1567

Request for Prior Authorization
Crisaborole (Eucrisa)

(PLEASE PRINT – ACCURACY IS IMPORTANT)

IA Medicaid Member ID #	Patient name	DOB
Patient address		
Provider NPI	Prescriber name	Phone
Prescriber address		Fax
Pharmacy name	Address	Phone
Prescriber must complete all information above. It must be legible, correct, and complete or form will be returned.		
Pharmacy NPI	Pharmacy fax	NDC

Prior authorization (PA) is required for Eucrisa (crisaborole). Payment will be considered when patient has an FDA approved or compendia indication for the requested drug when the following criteria are met: 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 mild to moderate atopic dermatitis; and 3) Patient has failed to respond to good skin care and regular use of emollients; and 4) Patient has documentation of an adequate trial and therapy failure with one preferred medium to high potency topical corticosteroid for a minimum of 2 consecutive weeks; and 5) Patient has documentation of a previous trial and therapy failure with a topical immunomodulator for a minimum of 4 weeks; and 6) Patient will continue with skin care regimen and regular use of emollients. 7) Quantities will be limited to 60 grams for use on the face, neck, and groin and 100 grams for all other areas, per 30 days. The required trials may be overridden when documented evidence is provided that use of these agents would be medically contraindicated.

Non-Preferred

☐ Eucrisa

Strength	Usage Instructions	Quantity	Day's Supply

Diagnosis: _____

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

Document emollient use: Product name, dosing instructions & duration of use: _____

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

☐ Yes Emollient to be used: _____ ☐ No

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: _____

Affected area to be treated: _____

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

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.*