

**Topical Immunomodulators
Eucrisa (crisaborole)
Effective 08/01/2022**

Plan	<input type="checkbox"/> MassHealth <input checked="" type="checkbox"/> MassHealth (PUF) <input type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization <input checked="" type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit (NLX)		
Specialty Limitations	N/A		
Contact Information	Specialty Medications		
	All Plans	Phone: 866-814-5506	Fax: 866-249-6155
	Non-Specialty Medications		
	MassHealth	Phone: 877-433-7643	Fax: 866-255-7569
	Commercial	Phone: 800-294-5979	Fax: 888-836-0730
	Exchange	Phone: 855-582-2022	Fax: 855-245-2134
	Medical Specialty Medications (NLX)		
	All Plans	Phone: 844-345-2803	Fax: 844-851-0882
Exceptions	N/A		

Overview

No PA	Drugs that require PA
Elidel [®] (pimecrolimus) §	Eucrisa [®] (crisaborole) ^{PD}
Protopic [®] # (tacrolimus topical)	

^{PD} Preferred Drug: A trial of the preferred drug or clinical rationale for prescribing a non-preferred drug within a therapeutic class. Please note, for Immune Suppressants – Topical agents, a trial with a preferred agent is not required prior to approval of a non-preferred agent.

§ Brand Preferred over generic equivalents: A trial of the preferred drug or clinical rationale for prescribing the non-preferred drug generic equivalent.

This is a brand-name drug with FDA "A"-rated generic equivalents. PA is required for the brand, unless a particular form of that drug (for example, tablet, capsule or liquid) does not have an FDA "A"-rated generic equivalent.

Coverage Guidelines

Authorization may be reviewed on a case by case basis for members new to AllWays Health Partners who are currently receiving treatment with requested medication excluding when the product is obtained as samples or via manufacturer’s patient assistance programs

OR

Authorization may be granted if the member meets **ALL** following criteria and documentation has been submitted:

1. The member has a diagnosis of atopic dermatitis (eczema)
2. The member is ≥ 3 months of age



3. The member meets **ONE** of the following:
 - a. Paid claims or physician documented of inadequate response or adverse reaction to **ONE** topical corticosteroid **OR** topical calcineurin inhibitor (e.g. pimecrolimus or tacrolimus)
 - b. Contraindication to **BOTH** topical corticosteroids and topical calcineurin inhibitors
4. **ONE** of the following:
 - a. Request is for 60 gram/month. If the request does not document quantity, may be approved for the 60 gram/month if member meets criteria.
 - b. Medical necessity for exceeding the quantity limits (i.e. large surface area of lesion)

Continuation of Therapy

Reauthorizations by prescriber will infer a positive response to therapy.

Limitations

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

Eucrisa	60 grams per 30 days
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Brand preferred over generic equivalent

In addition to any prior authorization requirements, generic medications listed below require a trial of the preferred drug or clinical rationale for prescribing the non-preferred drug generic equivalent.

- Pimecrolimus

References

1. Eucrisa Ointment 2% (crisaborole) [prescribing information]. New York, NY: Pfizer Labs; April 2020.
2. Protopic (tacrolimus) [prescribing information]. Madison, NJ: LEO Pharma Inc.; February 2019.
3. Elidel (pimecrolimus) [prescribing information]. Bridgewater, NJ: Valeant Pharmaceuticals; December 2017.

Review History

10/01/2020 – Updated MH partial unified formulary; retired ST criteria and switched to PA; added Eucrisa PA criteria, added QL, Effective 1/1/2021

11/17/2021 – Updated and reviewed for Nov P&T; matched MH UPPL for 1/1/2022. No clinical changes.

06/22/2022 - Reviewed and updated for June P&T; matched MH UPPL. Guideline updated to remove Protopic from the brand preferred over generic list. Effective 08/01/2022.

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