

Enhanced Specialty Guideline Management Treatment of Atopic Dermatitis Cibinqo

Products Referenced by this Document

Drugs that are listed in the following table include both brand and generic and all dosage forms and strengths unless otherwise stated. Over-the-counter (OTC) products are not included unless otherwise stated.

Brand Name	Generic Name
Cibinqo	abrocitinib

Indications

This program applies to Cibinqo for the treatment of atopic dermatitis. For indications other than atopic dermatitis, refer to the Specialty Guideline Management program for Cibinqo. Coverage will be provided if all the approval criteria are met and the member has no exclusions to the prescribed therapy.

Documentation

Submission of the following information is necessary to initiate the prior authorization review:

Initial requests:

• Member's chart notes or medical records showing affected area(s) and affected body surface area (where applicable).

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 Member's chart notes, medical record documentation and claims history of prerequisite therapies (including topical calcineurin inhibitors,topical corticosteroids, or biologics/targeted synthetic drugs) including dosage, duration, and response to therapy. If prerequisite therapy is not advisable, documentation of why topical corticosteroid, topical calcineurin inhibitor, and/or systemic drug product is/are not advisable for the member.

Continuation requests:

• Documentation (e.g., chart notes) that the member has experienced a positive clinical response to therapy as evidenced by low disease activity or improvement in signs or symptoms of atopic dermatitis.

Prescriber Specialties

This medication must be prescribed by or in consultation with a dermatologist or allergist/immunologist.

Coverage Criteria

Authorization of 4 months may be granted for members 12 years of age or older for treatment of moderate-to-severe atopic dermatitis when the member has had an inadequate response or intolerance to at least one biologic (e.g., Dupixent, Adbry) or a targeted synthetic drug (e.g., Rinvoq) in the past 180 days.

Authorization of 4 months may be granted for treatment of moderate-to-severe atopic dermatitis in members 12 years of age or older when all of the following criteria are met:¹⁻⁶

- Affected body surface is greater than or equal to 10% body surface area OR crucial body areas (e.g., hands, feet, face, neck, scalp, genitals/groin, intertriginous areas) are affected.
- Member meets either of the following:
 - Member has had an inadequate treatment response with either of the following in the past 180 days:
 - A high potency or super-high potency topical corticosteroid (see Appendix)
 - o A topical calcineurin inhibitor
 - The use of high potency or super-high potency topical corticosteroid and topical calcineurin inhibitor are not advisable for the member (e.g., due to contraindications, prior intolerances, potency not appropriate for member's age).
- Member has had an inadequate response to treatment with a systemic drug product (e.g., oral cyclosporine, azathioprine, methotrexate, mycophenolate mofetil) indicated for the treatment of atopic dermatitis, or use of these therapies are not advisable for the member.

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Continuation of Therapy

Authorization of 12 months may be granted for members 12 years of age or older (including new members) who are using the requested medication for moderate-to-severe atopic dermatitis and who achieve or maintain a positive clinical response as evidenced by low disease activity (i.e., clear or almost clear skin), or improvement in signs and symptoms of atopic dermatitis (e.g., redness, itching, oozing/crusting).

Other^{1,7}

Member has had a documented negative tuberculosis (TB) test (which can include a tuberculosis skin test [TST] or an interferon-release assay [IGRA]) within 6 months of initiating therapy for persons who are naïve to biologic drugs or targeted synthetic drugs associated with an increased risk of TB.

If the screening testing for TB is positive, there must be further testing to confirm there is no active disease (e.g., chest x-ray). Do not administer the requested medication to members with active TB infection. If there is latent disease, TB treatment must be started before initiation of the requested medication.

Member cannot use the requested medication concomitantly with any other biologic drug, targeted synthetic drug, or potent immunosuppressant such as azathioprine or cyclosporine.

Dosage and Administration

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Appendix

Potency	Drug	Dosage form	Strength
I. Super-high	Augmented betamethasone	Ointment, Lotion, Gel	0.05%
potency (group 1)	dipropionate		
I. Super-high	Clobetasol propionate	Cream, Gel, Ointment, Solution,	0.05%
potency (group 1)		Cream (emollient), Lotion,	
-		Shampoo, Foam, Spray	
I. Super-high	Fluocinonide	Cream	0.1%
potency (group 1)			

Table. Relative Potency of Select Topical Corticosteroid Products⁸

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Potency	Drug	Dosage form	Strength
I. Super-high	Flurandrenolide	Tape	4 mcg/cm ²
potency (group 1)		- 1	- 3 -
I. Super-high	Halobetasol propionate	Cream, Lotion, Ointment, Foam	0.05%
potency (group 1)			
II. High potency	Amcinonide	Ointment	0.1%
(group 2)			
II. High potency	Augmented betamethasone	Cream	0.05%
(group 2)	dipropionate		
II. High potency	Betamethasone dipropionate	Ointment	0.05%
(group 2)			
II. High potency	Clobetasol propionate	Cream	0.025%
(group 2)			
II. High potency	Desoximetasone	Cream, Ointment, Spray	0.25%
(group 2)			
II. High potency	Desoximetasone	Gel	0.05%
(group 2)			
II. High potency	Diflorasone diacetate	Ointment, Cream (emollient)	0.05%
(group 2)			
II. High potency	Fluocinonide	Cream, Ointment, Gel, Solution	0.05%
(group 2)			
II. High potency	Halcinonide	Cream, Ointment	0.1%
(group 2)			
II. High potency	Halobetasol propionate	Lotion	0.01%
(group 2)			
III. High potency	Amcinonide	Cream, Lotion	0.1%
(group 3)			
III. High potency	Betamethasone dipropionate	Cream, hydrophilic emollient	0.05%
(group 3)			
III. High potency	Betamethasone valerate	Ointment	0.1%
(group 3)			
III. High potency	Betamethasone valerate	Foam	0.12%
(group 3)			
III. High potency	Desoximetasone	Cream, Ointment	0.05%
(group 3)			
III. High potency	Diflorasone diacetate	Cream	0.05%
(group 3)			
III. High potency	Fluocinonide	Cream, aqueous emollient	0.05%
(group 3)			
III. High potency	Fluticasone propionate	Ointment	0.005%
(group 3)			

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Potency	Drug	Dosage form	Strength
III. High potency	Mometasone furoate	Ointment	0.1%
(group 3)			
III. High potency	Triamcinolone acetonide	Cream, Ointment	0.5%
(group 3)			
IV. Medium	Betamethasone dipropionate	Spray	0.05%
potency (group 4)			
IV. Medium	Clocortolone pivalate	Cream	0.1%
potency (group 4)			
IV. Medium	Fluocinolone acetonide	Ointment	0.025%
potency (group 4)			
IV. Medium	Flurandrenolide	Ointment	0.05%
potency (group 4)		Ontinent	0.05%
IV. Medium	Hydrocortisone valerate	Ointment	0.2%
potency (group 4)			
IV. Medium	Mometasone furoate	Cream, Lotion, Solution	0.1%
potency (group 4)			
IV. Medium	Triamcinolone acetonide	Cream	0.1%
potency (group 4)			
IV. Medium	Triamcinolone acetonide	Ointment	0.05% and
potency (group 4)			0.1%
IV. Medium	Triamcinolone acetonide	Aerosol Spray	0.2 mg per
potency (group 4)			2-second
			spray
V. Lower-mid	Betamethasone dipropionate	Lotion	0.05%
potency (group 5)			
V. Lower-mid	Betamethasone valerate	Cream	0.1%
potency (group 5)			
V. Lower-mid	Desonide	Ointment, Gel	0.05%
potency (group 5)			
V. Lower-mid	Fluocinolone acetonide	Cream	0.025%
potency (group 5)			
V. Lower-mid	Flurandrenolide	Cream, Lotion	0.05%
potency (group 5)			
V. Lower-mid	Fluticasone propionate	Cream, Lotion	0.05%
potency (group 5)			
V. Lower-mid	Hydrocortisone butyrate	Cream, Lotion, Ointment,	0.1%
potency (group 5)		Solution	
V. Lower-mid	Hydrocortisone probutate	Cream	0.1%
potency (group 5)			

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Potency	Drug	Dosage form	Strength
V. Lower-mid	Hydrocortisone valerate	Cream	0.2%
potency (group 5) V. Lower-mid	Prednicarbate	Cream (emollient), Ointment	0.1%
potency (group 5)	Fredriical Date	Cream (enfoliency, Ontiment	0.176
V. Lower-mid	Triamcinolone acetonide	Lotion	0.1%
potency (group 5)			•
V. Lower-mid	Triamcinolone acetonide	Ointment	0.025%
potency (group 5)			
VI. Low potency (group 6)	Alclometasone dipropionate	Cream, Ointment	0.05%
VI. Low potency (group 6)	Betamethasone valerate	Lotion	0.1%
VI. Low potency (group 6)	Desonide	Cream, Lotion, Foam	0.05%
VI. Low potency (group 6)	Fluocinolone acetonide	Cream, Solution, Shampoo, Oil	0.01%
VI. Low potency (group 6)	Triamcinolone acetonide	Cream, lotion	0.025%
VII. Least potent (group 7)	Hydrocortisone (base, greater than or equal to 2%)	Cream, Ointment, Solution	2.5%
VII. Least potent (group 7)	Hydrocortisone (base, greater than or equal to 2%)	Lotion	2%
VII. Least potent (group 7)	Hydrocortisone (base, less than 2%)	Cream, Ointment, Gel, Lotion, Spray, Solution	1%
VII. Least potent (group 7)	Hydrocortisone (base, less than 2%)	Cream, Ointment	0.5%
VII. Least potent (group 7)	Hydrocortisone acetate	Cream	2.5%
VII. Least potent (group 7)	Hydrocortisone acetate	Lotion	2%
VII. Least potent (group 7)	Hydrocortisone acetate	Cream	1%

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