

PRIOR AUTHORIZATION REQUEST

<u>Opzelura</u>

| Patient Informati | n: | | |
|--|--|--|---|
| Name: | | | |
| Member ID: | | | |
| Address: | | | |
| City, State, Zip: | | | |
| Date of Birth: | | | |
| Prescriber Inforn | ation: | | |
| Name: | | | |
| NPI: | | | |
| Phone Number: | | | |
| Fax Number | | | |
| Address: | | | |
| City, State, Zip: | | | |
| Requested Medic | ation | | |
| Rx Name: | | | |
| Rx Strength | | | |
| Rx Quantity: | | | |
| Rx Frequency: | | | |
| Rx Route of | | | |
| Administration: | | | |
| Diagnosis and ICE | Code: | | |
| prescribed a medicated quantities can be prosuped upon receipt of the SECTION A: Prequests. Phare medications that | otion benefit requires that we review certain requests for coverage with the present for your patient that requires Prior Authorization before benefit coverage or covered. Please complete the following questions then fax this form to the toll-free number completed form, prescription benefit coverage will be determined based on ease note that supporting clinical documentation is required acy prior authorization reviews can be subject to trial with action are not listed within the criteria. The policies are subject to trial with action is required to the contract of the policies are subject to trial with action is required to the policies are subject to the contract of the policies are subject to the policies are subject t | erage of a mber liste the plan for ALL dditiona change | dditional d below. 's rules. <u>PA</u> |
| 1 Is the rec | est an INITIAL or CONTINUATION of therapy? | | |
| [] Initial (| checked, go to 7) | | |
| [] Continu | ion (If checked, go to 2) | | |
| | ent currently receiving the requested medication? to question 7.] | Yes | No |

If you have any questions, call: 1-888-258-8250

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| 3 | Has the patient been receiving medication samples for the requested medication? [If yes, skip to question 7.] | Yes | No |
|----|---|-----|----|
| 4 | Does the patient have a previously approved prior authorization (PA) on file with the current plan? [Note: If the patient does NOT have a previously approved PA on file for the requested medication with the current plan, the renewal request will be considered under initial therapy.] [If no, skip to question 7.] | Yes | No |
| 5 | Has the patient been established on therapy for AT LEAST 3 months? [If no, skip to question 7.] | Yes | No |
| 6 | Has the patient experienced a clinically significant benefit from the medication, as documented by the prescriber? [Note: Examples of a response to the requested medication for atopic dermatitis are marked improvements in erythema, induration/papulation/edema, excoriations, and lichenification; reduced pruritus; decreased requirement for other topical or systemic therapies; reduced body surface area affected with atopic dermatitis; or other responses observed.] [No further questions.] | Yes | No |
| 7 | What is the indication or diagnosis? | | |
| | [] Atopic dermatitis (If checked, go to 8) | | |
| | [] Nonsegmental Vitiligo (If checked, no further questions) | | |
| | [] Other (If checked, no further questions) | | |
| 8 | Is the patient greater than or equal to 2 years of age? [If no, no further questions.] | Yes | No |
| 9 | Does the patient have a documented diagnosis of mild to moderate atopic dermatitis? [If no, no further questions.] | Yes | No |
| 10 | Is the requested medication being prescribed by or in consultation with an allergist, immunologist, or dermatologist? [If no, no further questions.] | Yes | No |
| 11 | Does the patient have atopic dermatitis involvement estimated to be 3% to 20% of the body surface area (BSA) according to the prescriber? [If no, no further questions.] | Yes | No |
| 12 | Has the patient tried at least TWO medium-, medium-high, high-, and/or super-high-potency prescription topical corticosteroids? [If no, no further questions.] | Yes | No |

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| 13 | Was inadequate efficacy demonstrated with the topical corticosteroid therapy, according to the prescriber? [If no, no further questions.] | Yes | No |
|----|---|-----|----|
| 14 | Has the patient tried a topical calcineurin inhibitor (pimecrolimus or tacrolimus) for at least 28 consecutive days? [If no, no further questions.] | Yes | No |
| 15 | Was inadequate efficacy demonstrated with the topical calcineurin inhibitor (pimecrolimus or tacrolimus), according to the prescriber? [If no, no further questions.] | Yes | No |
| 16 | Has the patient tried Zoryve for at least 4 weeks? [If no, no further questions.] | Yes | No |
| 17 | Was inadequate efficacy demonstrated with Zoryve, according to the prescriber? [If no, no further questions.] | Yes | No |
| 18 | Does the dose of the requested medication exceed the Food and Drug Administration (FDA) approved labeled dosing for the indication (Dosing: a thin layer applied topically twice daily to affected areas of up to 20% body surface area)? | Yes | No |

Please document the diagnoses, symptoms, and/or any other information important to this review:

SECTION B: Physician Signature

PHYSICIAN SIGNATURE

DATE

FAX COMPLETED FORM TO: 1-833-896-0656

Disclaimer: An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

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