

Ebglyss

Patient Info	ormation:			
Name:				
Member ID	:			
Address:				
City, State,	Zip:			
Date of Birt				
	1			
Prescriber	Information:			
Name:				
NPI:				
Phone Num	nber:			
Fax Numbe	r			
Address:				
City, State,	Zip:			
Requested	Medication			
Rx Name:				
Rx Strength	1			
Rx Quantity	/ :			
Rx Frequer	icy:			
Rx Route of	f			
Administrat	ion:			
Diagnosis a	and ICD Code:			
prescribed a requantities can Upon receipt SECTION requests.	hedication for your be provided. Plea of the completed A: Please no Pharmacy prints that are no	fit requires that we review certain requests for coverage with the p patient that requires Prior Authorization before benefit coverage or complete the following questions then fax this form to the toll-free reference, prescription benefit coverage will be determined based on the that supporting clinical documentation is required or authorization reviews can be subject to trial with a tlisted within the criteria. The policies are subject to trial with the criteria and updates to treatment quickless.	overage of number list on the plant of the p	of additional sted below. lan's rules. LL PA onal ge based
1 Is	the request an INIT	IAL or CONTINUATION of therapy?		
[] [nitial (If checked, g	o to 7)		
[] (Continuation (If che	cked, go to 2)		
		receiving the requested medication?	Yes	No
[If	no, skip to question	7.]		

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3	Has the patient been receiving medication samples of Ebglyss?	Yes	No
	[If yes, skip to question 7.]		
4	Does the patient have a previously approved prior authorization (PA) on file with the current plan?	Yes	No
	[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.]		
5	Will the patient be concurrently receiving the requested medication in combination with any monoclonal antibody, anti-IL4, anti-IL5, or TSLP inhibitor such as Adbry, Cinqair, Dupixent, Fasenra, Nemluvio, Nucala, Tezspire or Xolair?	Yes	No
	[If yes, no further questions.]		
6	Has the patient been taking the requested medication for at least 4 months and has experienced a clinically significant benefit from the medication, as documented by the prescriber? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[Note: Atopic dermatitis – Examples of a response to the requested medication 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.]		
7	What is the indication or diagnosis?		
	[] Atopic dermatitis (If checked, go to 8)		
	[] Other (If checked, no further questions)		
8	Is the patient greater than or equal to 12 years of age?	Yes	No
9	What is the weight of the patient?		
	[] Less than 40 kg (If checked, no further questions)		
	[] More than or equal to 40 kg (If checked, go to 10)		
10	Will the patient be concurrently receiving the requested medication in combination with	Yes	No

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	any monoclonal antibody, anti-IL4, anti-IL5, or TSLP inhibitor such Adbry, Cinqair, Dupixent, Fasenra, Nemluvio, Nucala, Tezspire or Xolair?		
	[If yes, no further questions.]		
11	Has the patient undergone treatment with any immunosuppressive medications within the last 4 weeks (e.g., systemic corticosteroids, cyclosporine, mycophenolate-mofetil, IFN-γ, Janus kinase inhibitors, azathioprine, methotrexate, etc.)?	Yes	No
	[If yes, no further questions.]		
12	Does the patient have a documented diagnosis of moderate to serve atopic dermatitis?	Yes	No
	[If no, no further questions.]		
13	Does the prescribed dosing exceed FDA approved indication?	Yes	No
	[If yes, no further questions.]		
14	Has documentation been provided to confirm that the patient has an IGA score of greater than or equal to 3 and an EASI score of greater than or equal to 16? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
15	Does the patient have atopic dermatitis involvement estimated to be greater than or equal to 10% of the body surface area (BSA) according to the prescriber? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If yes, go to question 17.]		
16	Does the patient have atopic dermatitis affecting the following areas: hands, face, feet, eyes/eyelids, neck, scalp, skin folds, and/or genitalia?	Yes	No
	[If no, no further questions.]		
17	Has the patient tried at least TWO medium-, medium-high, high-, and/or super-high-potency prescription topical corticosteroids? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
18	Has the patient tried tacrolimus ointment? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
19	Were the topical corticosteroids and tacrolimus ointment applied daily for at least 28 consecutive days?	Yes	No
	[If no, no further questions.]		

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20	Was inadequate efficacy demonstrated with the topical therapies, according to the prescriber? [If no, no further questions.]	Yes	No
21	Has the patient tried Zoryve? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
22	Was Zoryve applied daily for at least 56 consecutive days? [If no, no further questions.]	Yes	No
23	Was inadequate efficacy demonstrated with Zoryve, according to the prescriber? [If no, no further questions.]	Yes	No
24	Is the requested medication being prescribed by or in consultation with an allergist, immunologist, or dermatologist?	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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