

## Cibingo

Patient Infor	mation:		
Name:			
Member ID:			
Address:			
City, State, Z	ip:		
Date of Birth:	•		
Prescriber Ir	oformation:		
Name:	ilomiation.		
NPI:			
Phone Numb	er		
Fax Number	01.		
Address:			
City, State, Z	ip:		
	· F ·		
Requested M	<b>dedication</b>		
Rx Name:			
Rx Strength			
Rx Quantity:			
Rx Frequenc	y:		
Rx Route of			
Administratio	n:		
Diagnosis an	d ICD Code:		
prescribed a medication:	edication for your be provided. Plea of the completed A: Please no Pharmacy pri s that are no	efit requires that we review certain requests for coverage with the prescriber. It patient that requires Prior Authorization before benefit coverage or coverage of ase complete the following questions then fax this form to the toll-free number list of form, prescription benefit coverage will be determined based on the place that supporting clinical documentation is required for Alignor authorization reviews can be subject to trial with addition to the listed within the criteria. The policies are subject to change the MDH transmittals and updates to treatment quidelines.	f additional sted below. an's rules.  LL PA mal ge based
	•		<u> </u>
		sis or indication? If checked, go to 2)	
[] Ot	her (If checked, r	no further questions)	
with		concurrently using the requested medication with a biologic or Yes thetic Disease-Modifying Antirheumatic Drug (DMARD)? uestions.]	No

3	Will the patient be concurrently using the requested medication with an anti- interleukin monoclonal antibody (such as, Dupixent or Adbry)? [If yes, no further questions.]	Yes	No
4	Will the patient be concurrently using the requested medication with other janus kinase inhibitors (such as, Rinvoq, Xeljanz/XR, Olumiant)? [If yes, no further questions.]	Yes	No
5	Will the patient be concurrently using the requested medication with Xolair (omalizumab subcutaneous injection)? [If yes, no further questions.]	Yes	No
6	Will the patient be concurrently using the requested medication with other potent immunosuppressants (such as, azathioprine, cyclosporine, or mycophenolate)? [If yes, no further questions.]	Yes	No
7	Is the requested medication being prescribed by or in consultation with an allergist, immunologist, or dermatologist? [If no, no further questions.]	Yes	No
8	Is the patient currently receiving the requested medication? [If no, skip to question 15.]	Yes	No
9	Has the patient been receiving medication samples for the requested medication? [If yes, skip to question 15.]	Yes	No
10	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 14.]	Yes	No
11	Has the patient already received at least 90 days of therapy with the requested medication? [If no, skip to question 15.]	Yes	No
12	Has documentation been submitted to confirm that the patient has experienced a beneficial clinical response, defined as improvement from baseline (prior to initiating the requested medication) in at least one of the following: A) Estimated body surface area affected, B) Erythema, C) Induration/papulation/edema, D) Excoriations, E) Lichenification, F) A decreased requirement for other topical or systemic therapies for atopic dermatitis? ACTION REQUIRED: Submit supporting documentation.  [If no, no further questions.]	Yes	No
13	Has documentation been submitted to confirm that when compared with baseline (prior to initiating the requested medication), the patient has experienced an	Yes	No

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	improvement in at least one symptom, such as decreased itching or scratching? ACTION REQUIRED: Submit supporting documentation. [No further questions.]		
14	Has documentation been provided to confirm that the clinical response of the patient's condition has improved compared to baseline? ACTION REQUIRED: Submit supporting documentation.  [If no, no further questions.]	Yes	No
15	Is the patient greater than or equal to 12 years of age? [If no, no further questions.]	Yes	No
16	Has documentation been submitted to confirm that the patient has a diagnosis of refractory, moderate to severe atopic dermatitis involving GREATER THAN OR EQUAL TO 10 percent of the body surface area according to the prescriber? ACTION REQUIRED: Submit supporting documentation. [If yes, skip to question 18.]	Yes	No
17	Has documentation been submitted to confirm that the patient has affected areas that are of the face, eyes/eyelids, skin folds, and/or genitalia? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
18	Has documentation been submitted to confirm that the patient has had a trial of at least two medium, medium-high, high, or super high potency topical corticosteroids unless intolerant, contraindicated, or unless treating the face/eyes/eyelid areas? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
19	Has documentation been submitted to confirm that the patient has had an intolerance, contraindication to, or treatment failure after at least 3 months with Dupixent (dupilumab)? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
20	Has the patient tried a topical calcineurin inhibitor for at least 28 consecutive days and inadequate efficacy was demonstrated? [If no, no further questions.]	Yes	No
21	Does the requested dose exceed Food and Drug Administration (FDA) approved label dosing for the requested indication?	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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