HEALTHY U MEDICAID

PRIOR AUTHORIZATION REQUEST FORM

Brand Name Atopic Dermatitis Agents

Adbry™, Cibinqo™, Dupixent®, Eucrisa®, Opzelura™, Rinvoq®

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for assistance: 385-425-5094

Disclaimer: Prior Authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:		ID#:		
DOB:	Gender:		Phys	sician:	
Office Phone:	Office Fax:		Offic	ce Contact:	
Height/Weight:					
Member must try formulary preferred dru preferred products has not been successfu reason for failure. Reasons for failure mu Product being requested: ☐ Adbry™, ☐ C Dosing/Frequency:	ul, you must submit which prefer ist meet the Health Plan medical	red prod necessi	lucts have ty criteria	e been tried, dates of treatment, and i.	
If the request is for reauthorization, proceed to reauthorization section.					
Questions		Yes	No	Comments/Notes	
EUCRISA					
 Does the member have a diagnosis dermatitis? 	of mild to moderate atopic			Please provide documentation	
2. Does documentation show that the member has had an adequate trial and failure of at least one topical corticosteroid?				Please provide documentation	
3. Does documentation show that the member has had an adequate trial and failure of topical tacrolimus or topical pimecrolimus?				Please provide documentation	
OPZELURA					
1. Is the request made by a provider specializing in dermatology, allergy, or immunology?					
2. Does documentation show a confirmed diagnosis of mild to moderate atopic dermatitis in a non-immune compromised individual who is not adequately controlled with topical prescription therapies or when these therapies are not advisable?				Please provide documentation	
3. Is the affected area less than 20% of	of body surface area?			Please provide documentation	
4. Does the quantity requested excee	ed one tube per 30 days?			Please provide documentation	
 5. Has the member had an adequate trial with the following: a topical calcineurin inhibitor, such as pimecrolimus or tacrolimus, 				Please provide documentation	

two medium to high potency corticosteroids (e.g.,						
triamcinolone acetonide 0.1%, mometasone furoate 0.1%,						
betamethasone dipropionate 0.05%, desoximetasone						
0.05%), and						
phototherapy? SYSTEMIC AGENTS						
Is the request made by a provider specializing in dermatology,	I					
allergy, or immunology?						
Has the member had an adequate trial with at least two			Please provide documentation			
moderate to very high potency prescription corticosteroids?			ricase provide documentation			
If unable to tolerate corticosteroids due to the treatment area			Please provide documentation			
(e.g. face, genitals, etc.), has the member had an adequate trial			r rease provide decamentation			
with a calcineurin inhibitor such as topical tacrolimus?						
4. Has the member tried phototherapy?			Please provide documentation			
5. Has the member had a trial of at least one of the following			Please provide documentation			
in the past 6 months:			P			
oral corticosteroid						
intramuscular steroid						
• cyclosporine						
azathioprine						
methotrexate						
 mycophenolate 						
CIBINQO						
1. Has the member had a 3-month trial and failure of Dupixent			Please provide documentation			
and Adbry, unless contraindicated?						
2. Does Clinical documentation show that Tb and Hepatitis			Please provide documentation			
Screening have been done?						
RINVOQ		I				
 Has the member had a 3-month trial and failure of Adbry, Cibinqo and Dupixent, unless contraindicated? 			Please provide documentation			
2. Does Clinical documentation show that Tb and Hepatitis			Please provide documentation			
Screening have been done?						
REAUTHORIZATION						
1. Is the request for reauthorization of atopic dermatitis therapy?						
2. Is there evidence of positive clinical response?			Please provide documentation			
What medications and/or treatment modalities have been tried in the past for this condition? Please document						
name of treatment, reason for failure, treatment dates, etc.						
Additional information:						
Additional morniduon.						

Physician	Signature:
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Policy PHARM-HU-135 Origination Date: 04/20/2022 Reviewed/Revised Date: 01/18/2023 Next Review Date: 01/18/2024 Current Effective Date: 02/01/2023

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