

<u>Otezla</u>

Patient Information:

Name: Member ID:

Address:				
City, State, Zip:				
Date of Birth:				
	4.			
Prescriber Inforn	nation:			
Name:				
NPI:				
Phone Number:				
Fax Number				
Address:				
City, State, Zip:				
Requested Medic	cation			
Rx Name:				
Rx Strength				
Rx Quantity:				
Rx Frequency:				
Rx Route of				
Administration:				
Diagnosis and ICE) Code:			
quantities can be pro Upon receipt of the SECTION A: P requests. Phare medications tha	vided. Plea completed lease no macy pri at are no	repatient that requires Prior Authorization before benefit coverage or consecutive the following questions then fax this form to the toll-free reports of form, prescription benefit coverage will be determined based of the that supporting clinical documentation is required or authorization reviews can be subject to trial with the tritle of the trial with the criteria. The policies are subject to the trial with the criteria. The policies are subject to the trial with the criteria. The policies are subject to the trial with the criteria.	number listed by the plan's in the plant	elow. rules.
targeted [Note: Ex (for exan etanerce	synthetic of camples of nple, Huminpt SC proo	medication be used in combination with other biologics or with a disease-modifying antirheumatic drugs (DMARDs)? biologics include but not limited to adalimumab SC products ra, biosimilars), Actemra (IV or SC), Cimzia, Cosentyx, and duct (for example, Enbrel, biosimilars), Ilumya, Skyrizi, Kevzara,	Yes No	
biosimila Stelara (rs), a rituxi IV or SC),	/ or SC), an infliximab IV products (for example, Remicade, imab IV products (for example, Rituxan, biosimilars), Siliq, Taltz, Tremfya, Entyvio, or Simponi (Aria or SC). Examples of disease-modifying antirheumatic drugs (DMARDs) include but If you have any		
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	not limited to Cibinqo, Olumiant, Rinvoq, Xeljanz, Xeljanz XR.] [If yes, no further questions.]		
2	Is the patient currently receiving the requested medication? [If no, skip to question 8.]	Yes	No
3	Has the patient been receiving medication samples for the requested medication? [If yes, skip to question 8.]	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 8.]	Yes	No
5	Has the patient been established on therapy for AT LEAST 3 months? [If no, skip to question 8.]	Yes	No
6	What is the indication or diagnosis? [] Behcet's disease (If checked, go to 7)		
	[] Plaque psoriasis (If checked, go to 7)		
	[] Psoriatic arthritis (If checked, go to 7)		
	[] Ankylosing spondylitis (If checked, no further questions)		
	[] Rheumatoid arthritis (If checked, no further questions)		
	[] Other (If checked, no further questions)		
7	Has documentation been provided to confirm that the patient has had a clinically significant response to therapy, as determined by the prescriber? ACTION REQUIRED: Submit supporting documentation. [No further questions.]	Yes	No
8	What is the indication or diagnosis? [] Behcet's disease (If checked, go to 9)		
	[] Plaque psoriasis (If checked, go to 13)		
	[] Psoriatic arthritis (If checked, go to 18)		
	[] Ankylosing spondylitis (If checked, no further questions)		
	[] Rheumatoid arthritis (If checked, no further questions)		

	[] Other (If checked, no further questions)		
9	Is the patient greater than or equal to 18 years of age? [If no, no further questions.]	Yes	No
10	Is the requested medication being prescribed by or in consultation with a rheumatologist or dermatologist? [If no, no further questions.]	Yes	No
11	Does the patient have oral ulcers or other mucocutaneous involvement? [If no, no further questions.]	Yes	No
12	Has the patient tried AT LEAST TWO other systemic therapy for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance to AT LEAST TWO systemic agents for Behcet's disease? ACTION REQUIRED: Submit supporting documentation. [Note: Examples of systemic therapies include colchicine, azathioprine, thalidomide, interferon alpha, tumor necrosis factor inhibitors (for example, an adalimumab product {Humira, biosimilars}, an etanercept product {Enbrel, biosimilars}, Cimzia {certolizumab pegol subcutaneous injection}, Simponi {golimumab subcutaneous injection}, Simponi Aria {golimumab intravenous infusion}, or an infliximab product {Remicade, biosimilars})]. [If yes, skip to question 24.] [If no, no further questions.]	Yes	No
13	Is the patient greater than or equal to 6 years of age? [If no, no further questions.]	Yes	No
14	Is the requested medication being prescribed by or in consultation with a dermatologist? [If no, no further questions.]	Yes	No
15	Has the patient tried AT LEAST TWO traditional systemic agents for psoriasis for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance to AT LEAST TWO traditional systemic agents for psoriasis? ACTION REQUIRED: Submit supporting documentation. [Note: Examples of traditional systemic agents for psoriasis include methotrexate, cyclosporine, acitretin tablets.] [If no, no further questions.]	Yes	No
16	Has the patient tried BOTH preferred TNF Inhibitors, an adalimumab product (Simlandi, Hadlima, Yusimry, adalimumab- adbm) AND Enbrel (etanercept) for psoriasis for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance or has a contraindication? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions]	Yes	No
17	Has the patient tried a preferred ustekinumab product (Yesintek, Pyzchiva,	Yes	No

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	Steqeyma) for psoriasis AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance or has a contraindication? ACTION REQUIRED: Submit supporting documentation. [If yes, skip to question 24.] [If no, no further questions.]		
18	Is the patient greater than or equal to 18 years of age? [If no, no further questions.]	Yes	No
19	Has the patient tried AT LEAST TWO conventional synthetic disease-modifying antirheumatic drugs (DMARDs) for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance to AT LEAST TWO conventional DMARDs? [Note: Examples of conventional synthetic DMARDs include methotrexate (oral or injectable), leflunomide, hydroxychloroquine, sulfasalazine.] [If no, no further questions.]	Yes	No
20	Has the patient tried a preferred adalimumab product (Simlandi, Hadlima, Yusimry, adalimumab- adbm) for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance or has a contraindication? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
21	Has documentation been provided to confirm that the patient has had a treatment failure with a preferred ustekinumab product (Yesintek, Pyzchiva, Steqeyma) for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance or has a contraindication? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
22	Has documentation been provided to confirm that the patient has had a treatment failure with Xeljanz (tofacitinib) for AT LEAST 3 months OR has documentation been provided to confirm that the patient had an intolerance or has a contraindication? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
23	Is the requested medication being prescribed by or in consultation with a rheumatologist or dermatologist? [If no, no further questions.]	Yes	No
24	Does the requested dose exceed FDA approved label dosing for the requested indication? [Note: FDA approved dosing: 10 mg to 30 mg twice daily.]	Yes	No

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



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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