



## Otezla **Prior Authorization Request**

Send completed form to: Case Review Unit, CVS Caremark Prior Authorization Fax: 1-866-249-6155

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155. If you have questions regarding the prior authorization, please contact CVS Caremark at 1-866-814-5506. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

The recipient of this fax may make a request to opt-out of receiving telemarketing fax transmissions from CVS Caremark. There are numerous ways you may opt-out: The recipient may call the toll-free number at 877-265-2711, at any time, 24 hours a day/7 days a week. The recipient may also send an opt-out request via email to do not call@cvscaremark.com. An opt out request is only valid if it (1) identifies the number to which the request relates, and (2) if the person/entity making the request does not, subsequent to the request, provide express invitation or permission to CVS Caremark to send facsimile advertisements to such person/entity at that particular number. CVS Caremark is required by law to honor an opt-out request within thirty days of receipt.

Patient's Name:				
Pat	ient's ID:		Patient's Date of Birth:	
Phy	ysician's Name:			
Spe	ecialty:		NPI#:	
Phy	ysician Office Telephone:		Physician Office Fax:	
Rec	quest Initiated For:			
1.	Has the patient been diagnosed v ☐ Moderate to severe plaque pso ☐ Active psoriatic arthritis (PsA ☐ Other	oriasis )		
2.	What is the ICD-10 code?			
<u>Sec</u> 3.	Section A: Preferred Product - For Plaque Psoriasis  3. These are the primary preferred products for which coverage is provided for treatment of the following condition:  Plaque psoriasis: Humira (primary); Stelara/Taltz (secondary)*  *Note: Secondary preferred products for plaque psoriasis are Stelara and Taltz. These preferred product options only apply to members who have had a documented inadequate response or intolerable adverse event with Humira.			
to y	our office OR you may complete www.covermymeds.com/epa/car □ No	If Yes, please the PA electronically (elemark/ or call 1-866-452	e call 1-866-814-5506 to have the updated form faxed PA). You may sign up online via CoverMyMeds at:	
4.	Is this request for continuation of	f therapy with the reques	ted product?  \( \subseteq \text{ Yes} \) \( \subseteq \text{ No. } \textit{skip to #6} \)	
5.	Is the patient currently receiving the requested product through samples or a manufacturer's patient assistance program? If unknown, answer Yes.   Yes In No. If No., skip to Section B: All Requests			
6.	preferred products? Please indic <i>note(s)</i> . ☐ Humira:		intolerable adverse event with any of the following TION REQUIRED: If Yes, attach supporting chart  Intolerable adverse event Intolerable adverse event	
recip		on, distribution, or copying of this of	s solely for the use of individuals named above. If you are not the intended communication is prohibited. If you have received the fax in error, please PDPD SGM - 2/2018.	

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X_ Pre	criber or Authorized Signature Date (mm/dd/yy)			
info	t that this information is accurate and true, and that documentation supporting this nation is available for review if requested by CVS Caremark or the benefit plan sponsor.			
17.	oes the patient have a clinical reason to avoid pharmacologic treatment with methotrexate, cyclosporine or citretin?    Yes   No    If Yes, indicate the clinical reason:			
16.	as the patient experienced an inadequate response, or has an intolerance to phototherapy (e.g., UVB, PUV) pharmacologic treatment with methotrexate, cyclosporine or acitretin? <i>If Yes, no further questions</i> $\square$ Y No			
15.	less than 5% of BSA affected, are crucial body areas (e.g., hands, feet, face, neck, scalp, genitals/groin, tertriginous areas) affected? ☐ Yes ☐ No			
	n C: Plaque Psoriasis  That is the percentage of body surface area (BSA) affected? %			
Cor	lete the following section based on the patient's diagnosis, if applicable.			
13.	<ul> <li>3. Has the patient received any of the following medications?</li> <li>If Yes, please indicate the most recent medication and skip to diagnosis section.</li> <li>□ Cosentyx</li> <li>□ Enbrel</li> <li>□ Humira</li> <li>□ Inflectra</li> <li>□ Remicade</li> <li>□ Renflexis</li> <li>□ Stelara</li> <li>□ Taltz</li> <li>□ Tremfya</li> <li>□ No</li> </ul>			
12.	Has the patient achieved or maintained positive clinical response to treatment as evidenced by low disease activity or improvement in signs and symptoms? <i>If Yes, no further questions</i> $\square$ Yes $\square$ No			
11.	How long has the patient been receiving the requested medication? months If less than 4 months, no further questions.			
10.	Is the patient currently receiving Otezla through samples or a manufacturer's patient assistance program?  Yes In No Unknown For plaque psoriasis requests: If Yes or Unknown, skip to #13.			
	n B: All Requests this request for continuation of therapy? □ Yes □ No or plaque psoriasis requests: If No, skip to #13 or psoriatic arthritis requests: If No, no further questions.			
8.	Does the patient have one of the following documented exclusions to therapy with the preferred products?  **ACTION REQUIRED: If Yes, attach supporting chart note(s).  **Description of the preferred products of the preferred products?  **Description of the preferred products.  **De			
7.	oes the patient have one of the following documented clinical reasons to avoid Humira?  **CTION REQUIRED: If Yes, attach supporting chart note(s).**  1 Yes - History of demyelinating disorder			
	Taltz: ☐ Inadequate response ☐ Intolerable adverse event  No - none of the above ☐ Intolerable adverse event			