

Pharmacy Request for Prior Approval – Taltz

Beneficiary Information					
1. Beneficiary Last Name:	2. First Name:				
3. Beneficiary ID #:	4.0 (:: 0 : (0::1				
Prescriber Information	·				
6. Prescriber Name:	NPI #:				
Mailing address:			State:	 _ ZIP:	
7. Requester Contact Information:		ity:	State		
-			Fax #:		
	1 Holic #.				
Drug Information					
11. Length of Therapy:up to 30 days60	days90 days120 da	ys180 days	365 daysOther:		
Clinical Information					
Request for Ankylosing Spondylitis:					
1. Does the beneficiary have a diagnosis of Ankylosing Spondylitis? Yes No					
2. Is the beneficiary on another injectable biologic immunomodulator? Yes No					
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No					
4. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No					
5. Has the beneficiary experienced inadequate symptom relief from treatment with at least two NSAIDS? Yes No					
6. Is the beneficiary unable to receive treatment with NSAIDS due to contraindications? Yes No					
7. Does the beneficiary have clinical evidence of severe or rapidly progressing disease? YesNo					
8. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason beneficiary cannot try Cosentyx, Enbrel or					
Humira? Yes No					
Request for Plaque Psoriasis (Pediatric): (Ages 6 and up)					
1. Does the beneficiary have a diagnosis of plaque psoriasis and is a candidate for systemic therapy or phototherapy? YesNo					
2. Is the beneficiary 6 years of age or older? Yes No					
3. Is the beneficiary on another injectable biologic immunomodulator? YesNo					
4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No					
5. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No					
6. Has the beneficiary experienced a therapeutic failure/inadequate response with or has a contraindication or intolerance to					
methotrexate? Yes No					
7. Does the beneficiary have a body surface area (BSA) involvement of at least 3%? YesNo					
8. Does the beneficiary have involvement of the palms, soles, head and neck, or genitalia, causing disruption in normal daily activities					
and/or employment? YesNo					
9. Has the beneficiary had a trial and failure of Cosentyx or Enbrel or a clinical reason beneficiary cannot try Cosentyx or Enbrel?					
Yes No	,			•	
Request for Plaque Psoriasis (Adult):					
1. Does the beneficiary have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis? Yes No					
2. Is the beneficiary 18 years of age or older? Yes No					
3. Is the beneficiary on another injectable biologic immunomodulator? Yes No					
4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? YesNo					
5. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No					
6. Does the beneficiary have a body surface area (BSA) involvement of at least 3%? Yes No					
7. Does the beneficiary have involvement of the palms, soles, head and neck, or genitalia, causing disruption in normal daily activities					
and/or employment? Yes No					
8. Has the beneficiary failed to respond to, or has been unable to tolerate phototherapy and ONE of the following medications, or					
beneficiary has contraindications to these treatments: Soriatane (acitretin), Methotrexate, and/or Cyclosporine? Yes No					
9. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason beneficiary cannot try Cosentyx, Enbrel or					
Humira? YesNo					
Request for Psoriatic Arthritis:					
Does the beneficiary have a documented de	efinitive diagnosis of Psoria	tic Arthritis?	'es No		
2. Is the beneficiary 18 years of age or older? Yes No					



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3. Is the beneficiary on another injectable biologic immunomodulator? Yes No				
4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No				
5. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No				
6. Does the beneficiary have a documented inadequate response or inability to take methotrexate? Yes No				
7. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason beneficiary cannot try Cosentyx, Enbrel or				
Humira? Yes No				
Request for Non-Radiographic Axial Spondyloarthritis:				
1. Does the beneficiary have a diagnosis of Non-Radiographic Axial Spondyloarthritis? Yes No				
2. Is the beneficiary on another injectable biologic immunomodulator? Yes No				
3. Has the beneficiary failed an adequate trial of a Non-Steroidal Anti-Inflammatory Drug (NSAID) unless contraindicated? Yes No				
4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No				
5. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No				
6. Has the beneficiary had a trial and failure of Cosentyx, or a clinical reason beneficiary cannot try Cosentyx? Yes No				
Request for FDA Approved Diagnosis Not Listed Above:				
1. Diagnosis:				
2. Is the beneficiary on another injectable biologic immunomodulator? Yes No				
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No				
4. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No				

*Prescriber signature mandatory

Signature of Prescriber: _____

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

Date: _____