

Pharmacy Prior Approval Request for Immunomodulators: Taltz

Beneficiary	/ Information
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1. Beneficiary Last Name:	2. First Name:	
3. Beneficiary ID #:	4. Beneficiary Date of Birth:	5. Beneficiary Gender:

Prescriber Information

Drug Information

8. Drug Name:	9. Strength:	_ 10. Quantity Per 30 Days:		
11. Length of Therapy (in days): 🗆 up to 30 Days 🗆 60 Days 🗆 90 Days 🗆 120 Days 🗆 180 Days 🗆 365 Days				
Other				

Clinical Information

Requests for Ankylosing Spondylitis:

- 1. Does the beneficiary have a diagnosis of Ankylosing Spondylitis? \Box Yes \Box No
- 2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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? \square Yes \square 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?
- 8. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or have a clinical reason they cannot try Cosentyx, Enbrel or Humira?
 Yes No

Requests for Plaque psoriasis (Pediatric):

1. Does the beneficiary have a diagnosis of plaque psoriasis and is a candidate for systemic therapy or phototherapy?
Yes
No

2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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? \Box Yes \Box No
- 5. Has the beneficiary experienced a therapeutic failure/inadequate response with or has a contraindication or intolerance to methotrexate?
 Yes
 No
- 6. Does the beneficiary have 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. For ages 6 and up has there been a trial and failure of Cosentyx, Enbrel or Humira or have a clinical reason they cannot try Cosentyx, Enbrel or Humira?
Yes
No

Fax this form to (833) 404-2393

Pharmacy PA Call Center: (833) 585-4309



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Requests 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? \Box Yes \Box No

3. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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? \Box Yes \Box No

6. Does the beneficiary have body surface area (BSA) involvement of at least 3%?

7. Does the beneficiary have involvement of the palms, soles, head and neck, or genitalia, causing disruption in normal daily activities and/or employment?

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 have a clinical reason they cannot try Cosentyx, Enbrel or Humira?

Yes
No

Requests for Psoriatic Arthritis:

1. Does the beneficiary have a documented definitive diagnosis of Psoriatic Arthritis?

Yes
No

- 2. Is the beneficiary 18 years of age or older? \Box Yes \Box No
- 3. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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? \Box Yes \Box No
- 6. Does the beneficiary have a documented inadequate response or inability to take methotrexate? \Box Yes \Box No

7. Has the beneficiary had a trial and failure of Cosentyx, Enbrel or Humira or have a clinical reason they cannot try Cosentyx, Enbrel or Humira?
Question Yes
No

Requests for Non-Radiographic Axial Spondylorarthritis:

- 1. Does the beneficiary have a diagnosis of Non-Radiographic Axial Spondyloarthritis?
- 2. Is the beneficiary not on another injectable biologic immunomodulator? \Box Yes \Box 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?
- 5. Has the beneficiary been tested with Hep B SAG and Core Ab? \Box Yes \Box No

6. Has the beneficiary had a trial and failure of Cosentyx or a clinical reason beneficiary cannot try Cosentyx? **Yes No**

Signature of Prescriber: _

(Prescriber Signature Mandatory)

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.

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Date: