

Beneficiary Information

	2 First News
1. Beneficiary Last Name:	2. First Name:
3. Beneficiary ID #: 4. Beneficiary Data	te of Birth: 5. Beneficiary Gender:
Prescriber Information	
6. Prescriber Name:	NPI #:
Mailing address:	City: State: ZIP:
7. Requester Contact Information:	
Name: Phone #:	Fax #:
Drug Information	
8. Drug Name: 9. Dose:	10. Directions:
11. Length of Therapy:up to 30 days60 days90 days	120 days180 days365 daysOther:
Clinical Information	
 Does the beneficiary have a diagnosis of Ankylosing Spondylitis? YesNo Is the beneficiary on another injectable biologic immunomodulator? YesNo Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? YesNo Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No Has the beneficiary experienced inadequate symptom relief from treatment with at least two NSAIDS? Yes No Is the beneficiary unable to receive treatment with NSAIDS due to contraindications? Yes No Is the beneficiary have clinical evidence of severe or rapidly progressing disease? Yes No 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 Does the beneficiary have a diagnosis of moderate to severe Crohn's Disease? Yes No Is the beneficiary on another injectable biologic immunomodulator? Yes No Is the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? Yes No Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No 	
5. Has the beneficiary had a trial and failure of Humira, or a clinical reason beneficiary cannot try Humira? 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? Yes No 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? Yes No	
Request 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? 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? 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 Rheumatoid Arthritis:

- 1. Does the beneficiary have a diagnosis of Rheumatoid Arthritis? 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 a therapeutic failure/inadeguate response with methotrexate or at least one disease modifying antirheumatic drug (e.g., leflunomide, hydroxychloroquine, minocycline, sulfasalazine)? Yes No

6. Is the beneficiary unable to receive methotrexate or disease modifying antirheumatic drug due to contraindications or intolerabilities? Yes No

7. Does the beneficiary have clinical evidence of severe or rapidly progressing disease? Yes____No___

8. Has the beneficiary had a trial and failure of Enbrel or Humira or a clinical reason beneficiary cannot try Enbrel or Humira? Yes No_

Request for Ulcerative Colitis (Adult):

- 1. Does the beneficiary have a diagnosis of ulcerative colitis? 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 had a trial and failure of Humira, or a clinical reason beneficiary cannot try Humira? 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

Signature of Prescriber:

Date:

*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.