

## Pharmacy Request for Prior Approval – Renflexis

Beneficiary Information			
1. Beneficiary Last Name:			
3. Beneficiary ID #:	4. Beneficiary Date of Birth:	5. Beneficiar	y Gender:
Prescriber Information			
6. Prescriber Name:		NPI #:	
Mailing address:	City: _	State:	ZIP:
7. Requester Contact Information:			
Name:	Phone #:	Fax #:	
Drug Information			
	Dose:	10. Directions:	
11. Length of Therapy:up to 30 days60	days90 days120 days	_180 days365 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? YesNo			
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 Cosentyx, Enbrel or Humira or a clinical reason beneficiary cannot try Cosentyx, Enbrel or Humira? Yes No			
Request for Crohn's Disease: 1. Does the beneficiary have a diagnosis of moderate to severe Crohn's Disease? 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 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? YesNo			
6. Does the beneficiary have a body surface area (BSA) involvement of at least 3%? YesNo			
7. Does the beneficiary have involvement of the	ie palms, soles, head and neck,	or genitalia, causing disruption in n	ormal daily activities
and/or employment? YesNo		atath areas and ONE of the followin	a madications or
8. Has the beneficiary failed to respond to, or has been unable to tolerate phototherapy and <b>ONE</b> of the following medications, or beneficiary has contraindications to these treatments: Soriatane (acitretin), Methotrexate, and/or Cyclosporine? YesNo			
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	coscincyx, Elister of Hulling of	a chinear reason beneficiary carmot	try coscillyx, Elibrar of
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			
<u> </u>	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 heneficiary have a diagnosis of Rho	oumatoid Arthritis? Vas No	)	



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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/inadequate 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:			
Signature of Frescriber 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.