

Pharmacy Request for Prior Approval – Enbrel

Beneficiary Information				
1. Beneficiary Last Name:	2. First Name:			
3. Beneficiary ID #:	4. Beneficiary Date of Birth:	4. Beneficiary Date 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:				
	30 days60 days90 days120 days180 days365 daysOther:			
Clinical Information				
Request for Ankylosing Spondylitis: 1. Does the baneficiary have a diagnosis of Ankylosing Spondylitis? Ves. No.				
Does the beneficiary have a diagnosis of Ankylosing Spondylitis? Yes No 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				
Request for Polyarticular Juvenile Idiopathic Arthritis (PJIA):				
1. Does the beneficiary have a diagnosis of Polyarticular Juvenile Idiopathic Arthritis? Yes No				
2. 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				
5. Has the beneficiary tried one systemic corticosteroid (e.g., prednisone, methylprednisolone) or methotrexate, leflunomide or				
sulfasalazine with inadequate response or is unable to take these therapies due to contraindications? Yes No				
6. Does the beneficiary have PJIA subtype enthesitis related arthritis? Yes No				
Request 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 on another injectable biologic immunomodulator? Yes No				
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection? YesNo				
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 or has a contraindication or intolerance to				
methotrexate? YesNo				
6. Does the beneficiary have a body surface a	area (BSA) involvement of at least 3%? Ye	es 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				
Request for Plaque Psoriasis (Adult):				
1. Does the beneficiary have a documented of	definitive diagnosis of moderate-to-severe	Chronic Plaque Psoriasis?	Yes No	
2. Is the beneficiary 18 years of age or older?		•		
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 tre	eatments: Soriatane (acitretin), Methotrex	ate, or Cyclosporine? Yes_	No	
Request for Psoriatic Arthritis:				
Does the beneficiary have a documented of t	definitive diagnosis of Psoriatic Arthritis?	Yes No		



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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
Request for Rheumatoid Arthritis:
1. Does the beneficiary have a diagnosis of Rheumatoid Arthritis? YesNo
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
Request for FDA Approved Diagnosis Not Listed Above:
1. Diagnosis:
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

*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: _____