

Pharmacy Request for Prior Approval – Humira

Beneficiary Information			
1. Beneficiary Last Name:	2. First Name:		
3. Beneficiary ID #:	4. Beneficiary Date of Birth: 5. Beneficiary Gender:		iary Gender:
Prescriber Information			
6. Prescriber Name:	NPI #:		
Mailing address:		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 days6	0 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? YesNo			
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	e of severe or rapidly progressin	g disease? 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			
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			
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 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			
	thesitis related artifitis? Fes	NO	
Request for Plaque Psoriasis (Adult):	d-finiti diaifdt-	ta anno Changia Blanca Bania	-:-2 V N-
 Does the beneficiary have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis? Yes No 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			_ 110
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	pa, 5555,	, or Berneama, causing and aption .	
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			
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 s	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			



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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? 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/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 Ulcerative Colitis:
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
Request for Hidradenitis Suppurativa: (ages 12 and older)
1. Does the beneficiary have a diagnosis of Hidradenitis Suppurativa (moderate to severe)? 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
Request for Non-infectious Intermediate Posterior Panuveitis: (ages 2 and older)
1. Does the beneficiary have a diagnosis of Non-infectious Intermediate Posterior Panuveitis? 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
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
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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.