

Pharmacy Request for Prior Approval – Cimzia

Beneficiary Information					
1. Beneficiary Last Name:	: 2. First Name:				
3. Beneficiary ID #:	4. Beneficiary Date of Birth:		5. Βε	5. Beneficiary Gender:	
Prescriber Information					
6. Prescriber Name:	NPI #:				
Mailing address:					
7. Requester Contact Information:					
Name:	Phone #:		Fax #: _		
Drug Information					
8. Drug Name:	9. Dose:		10. Directions:		
11. Length of Therapy:up to 30 da	ys60 days90 days	120 days	180 days365 days	Other:	
Clinical Information					
Request for Ankylosing Spondylitis:					
1. Does the beneficiary have a diagnosi	s of Ankylosing Spondylitis	s? Yes No_			
2. Is the beneficiary on another injectal	_				
3. Has the beneficiary been considered			tuberculosis infection? \	Yes No	
4. Has the beneficiary been tested with	•				
5. Has the beneficiary experienced inac				? 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? Yes No					
8. Has the beneficiary had a trial and fa				v cannot try Cosentyx Enhrel or	
Humira? Yes No	nare or coscileyx, Elibrer), , , , , , , , , , , , , , , , , , ,	similar reason beneficial y	y carmot try coscilityx, Elibrer of	
Request for Crohn's Disease (Adult):					
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? YesNo					
4. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No					
5. Has the beneficiary had a trial and fa	ílure of Humira, or a clinic	al reason bene	ficiary cannot try Humira	? Yes No	
Request for Non-Radiographic Axial Sp					
1. Does the beneficiary have a diagnosis of Non-Radiographic Axial Spondyloarthritis? Yes No					
2. Is the beneficiary on another injectable biologic immunomodulator? YesNo					
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? Yes No					
5. Has the beneficiary been tested with				res NO	
6. Has the beneficiary had a trial and fa	•		-		
Request for Plaque Psoriasis (Adult):					
	nted definitive diagnosis c	of moderate-to	-severe Chronic Plaque Ps	soriasis? Yes No	
 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 injectal		lator? Yes	No		
4. Has the beneficiary been considered	and screened for the pres	sence of latent	tuberculosis infection? `	Yes No	
5. Has the beneficiary been tested with	•		_		
6. Does the beneficiary have a body surface area (BSA) involvement of at least 3%? Yes No					
7. Does the beneficiary have involvement	nt of the palms, soles, hea	ad and neck, or	genitalia, causing disrup	tion in normal daily activities	
and/or employment? Yes No	to or has been unable to	tolerate photo	otherany and ONE of the	following medications or	
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	or obscury, embrer		oar reason beneficial y	, same a y sosemeyn, embrer or	



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North Carolina

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/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 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.