

NC Medicaid Pharmacy Prior Approval Request Immunomodulators: Cimzia

illillullollloudators.

Beneficiary Information			
Beneficiary Last Name:	2. First Name:		
3. Beneficiary ID #:	4. Beneficiary Date of Birth:		5. Beneficiary Gender:
Prescriber Information			
6. Prescribing Provider NPI #:			
			Ext
Drug Information			
8. Drug Name: 11. Length of Therapy (in days): Other			0. Quantity Per 30 Days: vs
Clinical Information			
 4. Has the beneficiary been test 5. Has the beneficiary experience 6. Is the beneficiary unable to re 7. Does the beneficiary have clir 	liagnosis of Ankylosing Spondyl ther injectable biologic immund sidered and screened for the properties ed with Hep B SAG and Core Akted inadequate symptom relief eceive treatment with NSAIDS of hical evidence of severe or rapid I and failure of Cosentyx, Enbre	omodulator? Yes cresence of latent tube o? Yes No from treatment with due to contraindication dly progressing disea	erculosis infection? Yes No at least two NSAIDS? Yes No ons? Yes No
4. Has the beneficiary been test	liagnosis of moderate to severe ther injectable biologic immund sidered and screened for the pi ed with Hep B SAG and Core Al	omodulator?	
Yes □ No 2. Is the beneficiary 18 years of 3. Is the beneficiary not on anot 4. Has the beneficiary been confor Otezla)? □ Yes □ No 5. Has the beneficiary been test	locumented definitive diagnosist age or older? Yes No her injectable biologic immunous sidered and screened for the properties of the p	omodulator? Yes resence of latent tube o? Yes No	erculosis infection (not required
6. Does the beneficiary have a b	ody surrace area (BSA) involve	ment of at least 3%?	⊔ Yes ⊔ NO



NC Medicaid Pharmacy Prior Approval Request

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
10. Are the beneficiaries, Providers, and Pharmacies utilizing Siliq registered appropriately in the Siliq Risk Evaluation
and Mitigation Strategy Program (REMS program)? Yes No
Request for Psoriatic Arthritis
1. Does the beneficiary have a documented definitive diagnosis of Psoriatic Arthritis? \square Yes \square No
2. Is the beneficiary 18 years of age or older (OR 2 years or older for Simponi Aria)? \Box Yes \Box No
3. Is the beneficiary not on another injectable biologic immunomodulator? \square Yes \square No
4. Has the beneficiary been considered and screened for the presence of latent tuberculosis infection (not required
for Otezla? Yes No
5. Has the beneficiary been tested with Hep B SAG and Core Ab (not required for Otezla? \square Yes \square No
6. Does the beneficiary have a documented inadequate response or inability to take methotrexate? \square Yes \square 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? \square Yes \square No
2. Is the beneficiary not on another injectable biologic immunomodulator? \square Yes \square No
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis? \Box Yes \Box No
4. Has the beneficiary been tested with Hep B SAG and Core Ab? \square Yes \square 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)? \square Yes \square 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 Non-Radiographic Axial Spondyloarthritis
1. Does the beneficiary have a diagnosis of Non-Radiographic Axial Spondyloarthritis?□ Yes □ No
2. Is the beneficiary not on another injectable biologic immunomodulator? Ves No
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? \Box Yes \Box No
5. Has the beneficiary been tested with Hep B SAG and Core Ab? Yes No
6. Has the beneficiary had a trial and failure of Cosentyx? Yes No

Pharmacy PA Call Center: 1-866-799-5318



NC Medicaid

WellCare	Pharmacy Prior Approval Request			
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.

Pharmacy PA Call Center: 1-866-799-5318