

NC Medicaid Pharmacy Prior Approval Request Immunomodulators: Inflectra

Beneficiary Information

1. Beneficiary Last Name:	2. F	irst Name:			
3. Beneficiary ID #: 4.	4. Beneficiary Date of Birth:			5. Beneficiary Gender:	
Prescriber Information					
6. Prescribing Provider NPI #:				_	
7. Requester Contact Information - Name:		Pho	one #:	Ext	
Drug Information					
8. Drug Name:	9. Strength:		10. Qı	uantity Per 30 Days:	
11. Length of Therapy (in days): up to 30					
Other					
Clinical Information					
Request for Ankylosing Spondylitis					
1. Does the beneficiary have a diagnosis of Ankylosing Spondylitis? Yes No					
2. Is the beneficiary not on another injectable biologic immunomodulator? Yes No					
3. Has the beneficiary been considered and		•		ulosis infection? Yes No	
4. Has the beneficiary been tested with He				least two NCAIDS or is upable	
5. Has the beneficiary experienced inadeque to receive treatment with NSAIDS due to co					
disease? ☐ Yes ☐ No	zirer ann area eronis e	or rias cirricar	evidence or s	reverse or rupidity progressing	
6. Has the beneficiary had a trial and failure	of Cosentyx, Enl	brel or Humir	a or a clinical	reason beneficiary cannot try	
Cosentyx, Enbrel or Humira? Yes No	, .			, ,	
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 not on another injecta	_				
3. Has the beneficiary been considered and		•		ulosis 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 					
5. Has the beneficiary had a trial and failur	e of Humira or a c	clinical reasor	n beneficiary o	cannot try Humira? 🗆 Yes 🗆 No	
Request for Crohn's Disease (Pediatric)					
1. Does the beneficiary have a diagnosis of moderate to severe Crohn's Disease? Yes 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 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 					
5. Has the beneficiary had a trial and failur	: OI HUIIIII OF A C	ınıncar reasor	i beneficiary (Cannot try number 1 Yes 11 No	
Request for_Plaque Psoriasis (Adult)					



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1. Does the beneficiary have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis? \Box
Yes □ No
2. Is the beneficiary 18 years of age or older? Yes No
3. Is the beneficiary not on another injectable biologic immunomodulator? Yes 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? 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 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
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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? \(\subseteq \text{Yes} \subseteq \text{No} \)
 3. Is the beneficiary not 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? \square Yes \square 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? \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? \square Yes \square 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) ? \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 Ulcerative Colitis (Adult)
1. Does the beneficiary have a diagnosis of ulcerative colitis? Yes No
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2. Is the beneficiary not on another injectable biologic immunomodulator? ☐ Y 3. Has the beneficiary been considered and screened for the presence of laten 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 had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical reason beneficiary had a trial and failure of Humira or a clinical re	t tuberculosis? 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.

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