

Pharmacy PA Call Center: 1-855-258-1593

NC Medicaid Pharmacy Prior Approval Request Immunomodulators: Inflectra

Beneficiary Information

beneficially information				
1. Beneficiary Last Name:	2. First Name:			
3. Beneficiary ID #:	4. Beneficiary Date of Birth:		5. Beneficiary Gender:	
Prescriber Information				
6. Prescribing Provider NPI #:				
7. Requester Contact Information - I	Name:	Phone #:	Ext	
Drug Information				
8. Drug Name:	9. Strength:		10. Quantity Per 30 Days:	
11. Length of Therapy (in days):	up to 30 Days ☐ 60 Days	□ 90 Days □ 120 I	Days □ 180 Days □ 365 Days □	
Other				
Clinical Information				
Request for Ankylosing Spondyl	tis			
1. Does the beneficiary have a di		•		
2. Is the beneficiary not on anoth	•			
•		•	uberculosis infection? Yes No	
4. Has the beneficiary been teste	•		When he are a NGAIRG and a salely	
7 .			vith at least two NSAIDS or is unable ce of severe or rapidly progressing	
disease? ☐ Yes ☐ No	o due to contrainalcations o	i ilas cillical evideli	ce of severe of rapidly progressing	
	and failure of Cosentyx Enl	orel or Humira or a c	linical reason beneficiary cannot try	
Cosentyx, Enbrel or Humira?	•	, e. e	initial reason selfendially earmorely	
Request for Crohn's Disease (Ad	ult)			
1. Does the beneficiary have a di	agnosis of moderate to seve	ere Crohn's Disease?	☐ Yes ☐ No	
2. Is the beneficiary not on anoth	ier injectable biologic immu	nomodulator? 🗆 Ye	s □ No	
3. Has the beneficiary been cons	dered and screened for the	presence of latent t	uberculosis infection? Yes No	
4. Has the beneficiary been teste	d with Hep B SAG and Core	Ab? □ Yes □ No		
5. Has the beneficiary had a trial	and failure of Humira or a c	linical reason benefi	ciary cannot try Humira? ☐ Yes ☐ No	
Request for Crohn's Disease (Pe	diatric)			
1. Does the beneficiary have a di	agnosis of moderate to seve	ere Crohn's Disease?	☐ Yes ☐ No	
2. Is the beneficiary not on anoth	er injectable biologic immu	nomodulator? 🗆 Ye	s □ No	
3. Has the beneficiary been cons	dered and screened for the	presence of latent t	uberculosis infection? 🗆 Yes 🗆 No	
4. Has the beneficiary been teste	d 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? \Box Yes \Box No				



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Request for Plaque Psoriasis (Adult)
1. Does the beneficiary have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis?
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
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? ☐ Yes ☐ 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? 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? \Box Yes \Box 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 not on another injectable biologic immunomodulator? \Box Yes \Box 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? ☐ 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? \square Yes \square 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



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Request for Ulcerative Colitis (Adult)	
1. Does the beneficiary have a diagnosis of ulcerative colit	is? □ Yes □ No
2. Is the beneficiary not on another injectable biologic imr	
3. Has the beneficiary been considered and screened for t	he presence of latent tuberculosis? \square Yes \square No
4. Has the beneficiary been tested with Hep B SAG and Co 5. Has the beneficiary had a trial and failure of Humira or a	re AD? LI Yes LI NO a clinical reason beneficiary cannot try Humira? 🗌 Yes 🗆 N o
Signature of Prescriber:	Date:
	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.