

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

NC Medicaid Pharmacy Prior Approval Request

Immunomodulators: Remicade and Infliximab

Beneficiary Information					
1. Beneficiary Last Name:		2. Fi	rst Name:		
3. Beneficiary ID #:4. Benefic		ciary Date of Birth:			5. Beneficiary Gender:
Prescriber Information					
6. Prescribing Provider NPI #:					_
7. Requester Contact Information - Name:			Pnone #:		Ext
Drug Information					
8. Drug Name:		9. Strength:		10. Q	Quantity Per 30 Days:
11. Length of Therapy (in days):					
Other					
Clinical Information					
1. Does the beneficiary have a d 2. Is the beneficiary been cons 4. Has the beneficiary been tests 5. Has the beneficiary experience to receive treatment with NSAID disease? ☐ Yes ☐ No 6. Has the beneficiary had a trial Cosentyx, Enbrel or Humira? ☐ Yes ☐ No 1. Does the beneficiary have a d 2. Is the beneficiary not on anot 3. Has the beneficiary been tests 5. Has the beneficiary had a trial Request for Crohn's Disease (Pe 1. Does the beneficiary have a d 2. Is the beneficiary had a trial Request for Crohn's Disease (Pe 1. Does the beneficiary have a d 2. Is the beneficiary not on anot 3. Has the beneficiary have a d 2. Is the beneficiary have a d 3. Has the beneficiary been cons 4. Has the beneficiary been tests 5. Has the beneficiary had a trial	iagnosis of Anky her injectable bisidered and screed with Hep B Soled inadequate soled inadequate soled Mo Idult) Idult) Idultagnosis of modelidered and screed with Hep B Soled inadequate bisidered injectable bisidered and screed with Hep B Soled injectable bisidered and screed with Hep B Soled injectable bisidered and screed with Hep B Soled	ened for the AG and Core symptom reliations of Cosentyx, Endications of Cosentyx, Endicated for the AG and Core dumira or a complete to several	nomodulator presence of Ab?	or? Yes Nof latent tuberon Noother tuberon Yes Yes Yes Noother tuberon Noother tuberon Yes Yes Noother tuberon Yes Yes Noother tuberon Yes Noother tuberon Noo	culosis infection? Yes No t least two NSAIDS or is unable severe or rapidly progressing I reason beneficiary cannot try Solo culosis infection? Yes No cannot try Humira? Yes No lo culosis infection? Yes No culosis infection? Yes No culosis infection? Yes 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 (not required for Otezla)? ☐ 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 (OR 2 years or older for Simponi Aria)? 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 (not required
for Otezla)? Yes No
5. Has the beneficiary been tested with Hep B SAG and Core Ab (not required for Otezla)? \Box Yes \Box 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? \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? ☐ 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) ? \Box Yes \Box
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



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Request for Ulcerative Colitis (Adult)
1. Does the beneficiary have a diagnosis of ulcerative colitis? \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 had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? \square Yes \square No
Request for Ulcerative Colitis (Pediatric)
1. Does the beneficiary have a diagnosis of ulcerative colitis? \square Yes \square 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? \square Yes \square No
5. Has the beneficiary had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? \square Yes \square No
Signature of Prescriber: Date: 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.

Fax this form to 1-866-940-7328 01.02.2025

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