Trillium Health Resources Pharmacy Prior Approval Request for



Immunomodulators: Remicade and Infliximab

Member Information				
1. Member Last Name:	2. First Na	ame:		
3. Member ID #:4. Member Date of Birth: 5.			_ 5. Member Gender:	
Prescriber Information				
6. Prescribing Provider NPI #:			_	
	n - Name:		Ext	
Drug Information				
8. Drug Name:	9. Strength:	10. Qu	antity Per 30 Days:	
11. Length of Therapy (in days):	□ up to 30 Days □ 60 Days □ 90 Days	□ 120 Days □ 180 Days □	365 Days	
Clinical Information				
 2. Is the member not on anoth 3. Has the member been cons 4. Has the member been teste 5. Has the member experience receive treatment with NSAID disease? ☐ Yes ☐ No 	iagnosis of Ankylosing Spondylitis? Iner injectable biologic immunomod idered and screened for the presented with Hep B SAG and Core Ab? ded inadequate symptom relief from S due to contraindications or has claim and failure of Cosentyx, Enbrel or Hereingtons	ulator?	st two NSAIDS or is unable to ere or rapidly progressing	
2. Is the member not on anoth3. Has the member been cons4. Has the member been teste	Adult) lagnosis of moderate to severe Croher injectable biologic immunomod idered and screened for the presented with Hep B SAG and Core Ab? and failure of Humira or a clinical r	ulator?	sis infection? Yes No	
Is the member not on anoth Has the member been cons	Pediatric) agnosis of moderate to severe Croher injectable biologic immunomod idered and screened for the presened with Hep B SAG and Core Ab?	ulator? Yes No nce of latent tuberculos		

5. Has the member had a trial and failure of Humira or a clinical reason member cannot try Humira? Yes No			
Request for_Plaque Psoriasis (Adult)			
1. Does the member have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis? \Box Yes			
□ No			
2. Is the member 18 years of age or older? Yes No			
3. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No4. Has the member been considered and screened for the presence of latent tuberculosis infection (not required for			
Otezla)? ☐ Yes ☐ No			
5. Has the member been tested with Hep B SAG and Core Ab (not required for Otezla)? \square Yes \square No			
6. Does the member have a body surface area (BSA) involvement of at least 3%? \square Yes \square No			
7. Does the member have involvement of the palms, soles, head and neck, or genitalia, causing disruption in normal daily activities and/or employment? \square Yes \square No			
8. Has the member failed to respond to, or has been unable to tolerate phototherapy and ONE of the following			
medications or member has contraindications to these treatments: Soriatane (acitretin), Methotrexate, and/or			
Cyclosporine? ☐ Yes ☐ No			
9. Has the member had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason member cannot try			
Cosentyx, Enbrel or Humira? Yes No			
Request for Psoriatic Arthritis			
1. Does the member have a documented definitive diagnosis of Psoriatic Arthritis? \square Yes \square No			
2. Is the member 18 years of age or older (OR 2 years or older for Simponi Aria)? \Box Yes \Box No			
3. Is the member not on another injectable biologic immunomodulator? \square Yes \square No			
4. Has the member been considered and screened for the presence of latent tuberculosis infection (not required for			
Otezla)? ☐ Yes ☐ No			
5. Has the member been tested with Hep B SAG and Core Ab (not required for Otezla)? \square Yes \square No			
6. Does the member have a documented inadequate response or inability to take methotrexate? \Box Yes \Box No			
7. Has the member had a trial and failure of Cosentyx, Enbrel or Humira or a clinical reason member cannot try			
Cosentyx, Enbrel or Humira? Yes No			
Request for Rheumatoid Arthritis			
1. Does the member have a diagnosis of Rheumatoid Arthritis? \square Yes \square No			
2. Is the member not on another injectable biologic immunomodulator? \square Yes \square No			
3. Has the member been considered and screened for the presence of latent tuberculosis? \Box Yes \Box No			
4. Has the member been tested with Hep B SAG and Core Ab? \square Yes \square No			
5. Has the member 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 member unable to receive methotrexate or disease modifying antirheumatic drug due to contraindications or			
intolerabilities? Yes No			
7. Does the member have clinical evidence of severe or rapidly progressing disease? Yes \(\subset \) No			
8. Has the member had a trial and failure of Enbrel or Humira or a clinical reason member cannot try Enbrel or			
Humira? ☐ Yes ☐ No			

Request for Ulcerative Colitis (Adult)
1. Does the member have a diagnosis of ulcerative colitis? \square Yes \square No
2. Is the member not on another injectable biologic immunomodulator? \square Yes \square No
3. Has the member been considered and screened for the presence of latent tuberculosis? \square Yes \square No
4. Has the member been tested with Hep B SAG and Core Ab? \square Yes \square No
5. Has the member had a trial and failure of Humira or a clinical reason member cannot try Humira? \Box Yes \Box No
Request for Ulcerative Colitis (Pediatric)
1. Does the member have a diagnosis of ulcerative colitis? \square Yes \square No
2. Is the member not on another injectable biologic immunomodulator? \square Yes \square No
3. Has the member been considered and screened for the presence of latent tuberculosis? \square Yes \square No
4. Has the member been tested with Hep B SAG and Core Ab? \square Yes \square No
5. Has the member had a trial and failure of Humira or a clinical reason member cannot try Humira? \square Yes \square 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.