Trillium Health Resources Pharmacy Prior Approval Request for



Immunomodulators: Inflectra

Member Information				
1. Member Last Name:	2. First Name:			
	4. Member Date of Birth:			
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6. Prescribing Provider NPI #:				
7. Requester Contact Information - Name:	Pho	ne #:	Ext	
Drug Information				
8. Drug Name:	9. Strength:	10. Quantity P	er 30 Days:	
11. Length of Therapy (in days): ☐ up to 30 Days	☐ 60 Days ☐ 90 Days	☐ 120 Days ☐ 180 D	ays 🗌 365 Days 🗌	
Other				
Clinical Information				
Request for Ankylosing Spondylitis				
1. Does the member have a diagnosis of Ankylo	sing Spondylitis? \square Yes \square	l No		
2. Is the member not on another injectable biol	ogic immunomodulator? [□ Yes □ No		
3. Has the member been considered and screen	ed for the presence of lat	ent tuberculosis infec	ction? 🗆 Yes 🗆 No	
4. Has the member been tested with Hep B SAG	and Core Ab? \square Yes \square I	No		
5. Has the member experienced inadequate syr	-			
receive treatment with NSAIDS due to contrain	dications or has clinical ev	idence of severe or ra	apidly progressing	
disease? ☐ Yes ☐ No		1		
6. Has the member had a trial and failure of Cos	entyx, Enbrei or Humira o	or a clinical reason me	ember cannot try	
Cosentyx, Enbrel or Humira? ☐ Yes ☐ No				
Request for Crohn's Disease (Adult)				
1. Does the member have a diagnosis of moder				
2. Is the member not on another injectable biol	-			
3. Has the member been considered and screen	•		ction? 🗆 Yes 🗆 No	
4. Has the member been tested with Hep B SAG				
5. Has the member had a trial and failure of Hu	mira or a clinical reason m	ember cannot try Hu	mira? Yes No	
Request for Crohn's Disease (Pediatric)				
1. Does the member have a diagnosis of moder	ate to severe Crohn's Dise	ase? 🗆 Yes 🗆 No		
2. Is the member not on another injectable biol	ogic immunomodulator? [□ Yes □ No		
3. Has the member been considered and screen	ed for the presence of lat	ent tuberculosis infec	ction? 🗆 Yes 🗆 No	
4. Has the member been tested with Hep B SAG	and Core Ab? \square Yes \square N	lo		
5. Has the member had a trial and failure of Hu	mira or a clinical reason m	ember cannot try Hu	mira? 🗆 Yes 🗆 No	

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Request for Plaque Psoriasis (Adult) 1. Does the member have a documented definitive diagnosis of moderate-to-severe Chronic Plaque Psoriasis?

Yes □ No 2. Is the member 18 years of age or older? \square Yes \square 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? \square Yes \square No 6. Does the member have a body surface area (BSA) involvement of at least 3%? ☐ Yes ☐ 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? ☐ Yes ☐ 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? ☐ Yes ☐ No 2. Is the member 18 years of age or older? \square Yes \square 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? ☐ Yes ☐ No 5. Has the member been tested with Hep B SAG and Core Ab? \square Yes \square No 6. Does the member have a documented inadequate response or inability to take methotrexate? \square Yes \square 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? \square Yes \square No 4. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ 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? \square Yes \square 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

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