Trillium Health Resources Pharmacy Prior Approval Request for



Immunomodulators: Simponi

Member Information								
1. Member Last Name:		2. First Name:						
3. Member ID #:	4. Member Date of Birth:					5. Member Gender:		
Prescriber Information								
6. Prescribing Provider NPI #:								
7. Requester Contact Information - Name:		Phone #:				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 Da	ys □ 365 Days □ Other		
_								
Clinical Information								
1. Does the member have a did 2. Is the member not on anoth 3. Has the member been considered. Has the member been tested 5. Has the member experience 6. Is the member unable to reconsidered. To Does the member have clinical 8. Has the member had a trial Cosentyx, Enbrel or Humira? Experience Cosentyx, Enbrel or Humira?	ner injectable biologidered and screen and screen and with Hep B SAG and inadequate symples treatment whical evidence of seand failure of Cos Yes \(\simeq\) No	ogic immunored for the pro and Core Ab aptom relief for the NSAIDS devere or rapid	modulator? esence of la ?	P Yes Natent tuberone No	culosis infed least two I s? Yes	NSAIDS? □ Yes □ No No		
1. Does the member have a do		ive diagnosis	of Psoriati	c Arthritis? l	□ Yes □ N	0		
2. Is the member 18 years of a		_						
	3. Is the member not on another injectable biologic immunomodulator? \square Yes \square No							
4. Has the member been consi	dered and screen	ed for the pro	esence of la	atent tubero	culosis infe	ction (not required for		
Otezla?	nd with Hen B SAG	and Core Ah	? □ V es □	No				
6. Does the member have a do 7. Has the member had a trial Cosentyx, Enbrel or Humira? [ocumented inadeq and failure of Cos	quate respons	se or inabil	ity to take m				
Request for Rheumatoid Arth	ritis							
1. Does the member have a di	agnosis of Rheum	atoid Arthriti	s? 🗆 Yes 🗆] No				

(Prescriber Signature Mandatory)	
Signature of Prescriber: Date:	
 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 	
Poquest for Ulcorative Colitic (Adult)	
7. Does the member have clinical evidence of severe or rapidly progressing disease? ☐ Yes ☐ 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	
2. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No 3. Has the member been considered and screened for the presence of latent tuberculosis? ☐ Yes ☐ 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 dise modifying antirheumatic drug (e.g. leflunomide, hydroxychloroquine, minocycline, sulfasalazine)? ☐ Yes ☐ No 6. Is the member unable to receive methotrexate or disease modifying antirheumatic drug due to contraindication intolerabilities? ☐ Yes ☐ No	

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.