Trillium Health Resources Pharmacy Prior Approval Request for



Immunomodulators: Actemra

iviember information				
1. Member Last Name:	2. First Name:			
3. Member ID #:	4. Member Date of Birth	:	5. Membe	er Gender:
Prescriber Information				
6. Prescribing Provider NPI #:				
7. Requester Contact Information -				Ext
Drug Information				
8. Drug Name:	9. Strength:		_ 10. Quantity Per	30 Days:
11. Length of Therapy (in days):				
Days Other				
Clinical Information				
Request for Polyarticular Juvenile 1. Does the member have a diagno 2. Is the member not on another in 3. Has the member been considere 4. Has the member been tested wit 5. Has the member tried one syste leflunomide or sulfasalazine with contraindications? Yes No 6. Does the member have PJIA sub 7. Has the member had a trial and the Humira? Yes No	sis of Polyarticular Juvenile ijectable biologic immunomod id and screened for the preser th Hep B SAG and Core Ab? emic corticosteroid (e.g. pre h inadequate response or is	Iulator?	□ No perculosis infection hylprednisolone) te these therapie No	n? □ Yes □ No or methotrexate, s due to
Request for Systemic Onset Juveni 1. Does the member have a diagno 2. Is the member not on another in 3. Has the member been considere 4. Has the member been tested wit 5. Does the member have systemic determined by the prescribing position of the member have a diagnostic street the member have a diagnostic street the member not on another in 3. Has the member been considered	sis of Systemic Juvenile Idio ejectable biologic immunomod d and screened for the preser th Hep B SAG and Core Ab? ic arthritis with active syste hysician (e.g. arthritis of the osis of Rheumatoid Arthritis ejectable biologic immunomod	Iulator?	□ No Derculosis infection Ind features of poor In phic damage)? □ No	oor prognosis, as ☐ Yes ☐ No

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(Prescriber Signature Mandatory)				
Signature of Prescriber:	Date:			
4. Has the member been tested with Hep B SAG and Core A	Ab? □ Yes □ No			
3. Has the member been considered and screened for the \mid	presence of latent tuberculosis infection? \square Yes \square No			
2. Is the member not on another injectable biologic immur	nomodulator? 🗆 Yes 🗆 No			
1. Does the member have a diagnosis of Systemic Scleros				
Request for Systemic Sclerosis-Associated Interstitial Lung	g Disease (SSc-ILD)			
4. Has the member been tested with Hep B SAG and Core A	Ab? □ Yes □ No			
3. Has the member been considered and screened for the				
2. Is the member not on another injectable biologic immur	nomodulator? 🗆 Yes 🗆 No			
1. Does the member have a diagnosis of Cytokine Release	e Syndrome? 🗆 Yes 🗆 No			
Request for Cytokine Release Syndrome:				
4. Has the member been tested with Hep B SAG and Core A	Ab □ Yes □ No			
3. Has the member been considered and screened for the	presence of latent tuberculosis infection? Yes No			
2. Is the member not on another injectable biologic immur	nomodulator? Yes No			
 Does the member have a diagnosis of Giant Cell Arter 	itis? 🗆 Yes 🗆 No			
Request for Giant Cell Arteritis:				
Enbrel or Humira? ☐ Yes ☐ No				
8. Has the member had a trial and failure of Enbrel or Hur	mira or a clinical reason member cannot try either			
7. Does the member have clinical evidence of severe or				
contraindications or intolerabilities? \square Yes \square No				
6. Is the member unable to receive methotrexate or dis	ease modifying antirheumatic drug due to			
Yes □ No				
disease modifying antirheumatic drug (e.g. leflunomic	de, hydroxychloroquine, minocycline sulfasalazine)?			
5. Has the member experienced a therapeutic failure/inadequate response with methotrexate or at least one				
4. Has the member been tested with Hep B SAG and Core A	Ab? □ 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.