Trillium Health Resources Pharmacy Prior Approval Request for



Immunomodulators: Avsola

Member Information		
L. 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:		
Drug Information		
8. Drug Name: 9. Stren	gth:	10. Quantity Per 30 Days:
11. Length of Therapy (in days): \Box up to 30 Days \Box 60 D		
Other		
Clinical Information		
Request for Ankylosing Spondylitis 1. Does the member have a diagnosis of Ankylosing Spondylitis 2. Is the member not on another injectable biologic immode in the state of the second for the second f	unomodulator? Yes e presence of latent tue Ab? Yes No elief from treatment with DS due to contraindicate rapidly progressing disc	berculosis infection? Yes No th at least two NSAIDS? Yes No tions? Yes No
1. Does the member have a diagnosis of moderate to se	vere Crohn's Disease? [☐ Yes ☐ No
2. Is the member not on another injectable biologic imm		
3. Has the member been considered and screened for the 4. Has the member been tested with Hep B SAG and Cor		berculosis infection? ☐ Yes ☐ No
5. Has the member had a trial and failure of Humira or a		r cannot try Humira? 🗆 Yes 🗆 No
Request for Crohn's Disease (Pediatric) 1. Does the member have a diagnosis of moderate to se 2. Is the member not on another injectable biologic imm 3. Has the member been considered and screened for th 4. Has the member been tested with Hep B SAG and Cor 5. Has the member had a trial and failure of Humira or a	unomodulator? □ Yes e presence of latent tu e Ab? □ Yes □ No	☐ No berculosis infection? ☐ Yes ☐ No
Request for Plaque Psoriasis (Adult)		

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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? Yes No 3. Is the member not on another injectable biologic immunomodulator? Yes 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? Yes 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 (actiretin), 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 (OR 2 years or older for Simponi Aria)? ☐ Yes ☐ No 3. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ 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? ☐ Yes ☐ No 6. Does the member have a documented inadequate response or inability to take methotrexate? ☐ Yes ☐ 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? ☐ 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 disease 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 contraindications or intolerabilities? ☐ Yes ☐ No 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			
Request for Ulcerative Colitis (Adult)			
1. Does the member have a diagnosis of ulcerative colitis? ☐ Yes ☐ No			
2. Is the member not on another injectable biologic immunomodulator? Yes No			

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(Proscriber Signature Mandatory)				
Signature of Prescriber:	Date:			
5. Has the member had a trial and failure of Humira or a clinical				
3. Has the member been considered and screened for the pres4. Has the member been tested with Hep B SAG and Core Ab?				
2. Is the member not on another injectable biologic immunom				
1. Does the member have a diagnosis of ulcerative colitis? Y				
Request for Ulcerative Colitis (Pediatric)				
5. Has the member had a trial and failure of Humira or a clinical	al reason member cannot try Humira? Yes No			
4. Has the member been tested with Hep B SAG and Core Ab?	☐ Yes ☐ No			
3. Has the member been considered and screened for the pres	sence of latent tuberculosis? 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.