

Pharmacy Prior Approval Request for Immunomodulators: Simponi

Beneficiary Information					
1. Beneficiary Last Name:	2. Fir	st Name:			
3. Beneficiary ID #:4.	4. Beneficiary Date of Birth:			5. Beneficiary Gender:	
Prescriber Information					
6. Prescribing Provider NPI #:				_	
7. Requester Contact Information - Name:	n - Name: Phone #:		one #:	Ext	
Drug Information					
8. Drug Name:	9. Strength:			Quantity Per 30 Days:	
11. Length of Therapy (in days): \Box up to 30	Days 🗆 60 Days	☐ 90 Days	\square 120 Days	\square 180 Days \square 365 Days \square	
Other					
Clinical Information					
Request for Ankylosing Spondylitis 1. Does the beneficiary have a diagnosis of 2. Is the beneficiary not on another injecta 3. Has the beneficiary been considered and 4. Has the beneficiary been tested with He 5. Has the beneficiary experienced inadeq 6. Is the beneficiary unable to receive trea 7. Does the beneficiary have clinical evider 8. Has the beneficiary had a trial and failur Cosentyx, Enbrel or Humira? Yes No Request for Psoriatic Arthritis	able biologic immund screened for the ep B SAG and Core a wate symptom relied the the threat with NSAIDS ance of severe or rap	nomodulato presence of Ab?	r?	culosis infection? Yes No t least two NSAIDS? Yes No No	
1. Does the beneficiary have a documente 2. Is the beneficiary 18 years of age or olde 3. Is the beneficiary not on another injecta 4. Has the beneficiary been considered and for Otezla? ☐ Yes ☐ No 5. Has the beneficiary been tested with He 6. Does the beneficiary have a documente 7. Has the beneficiary had a trial and failur Cosentyx, Enbrel or Humira? ☐ Yes ☐ No	er? Yes No able biologic immust d screened for the p B SAG and Core a d inadequate respo	nomodulato presence of Ab? Yes [onse or inab	r?	No culosis infection (not required nethotrexate? Yes No	

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Request for Rheumatoid Arthritis
1. Does the beneficiary have a diagnosis of Rheumatoid Arthritis? \square Yes \square No
2. Is the beneficiary not on another injectable biologic immunomodulator? \square Yes \square No
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis? \Box Yes \Box No
4. Has the beneficiary been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
5. Has the beneficiary 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 beneficiary unable to receive methotrexate or disease modifying antirheumatic drug due to contraindications
or intolerabilities? ☐ Yes ☐ No
7. Does the beneficiary have clinical evidence of severe or rapidly progressing disease? \Box Yes \Box No
8. Has the beneficiary had a trial and failure of Enbrel or Humira or a clinical reason beneficiary cannot try Enbrel or
Humira? ☐ Yes ☐ No
Request for Ulcerative Colitis (Adult)
1. Does the beneficiary have a diagnosis of ulcerative colitis? Yes No
 Is the beneficiary not on another injectable biologic immunomodulator? ☐ Yes ☐ No
3. Has the beneficiary been considered and screened for the presence of latent tuberculosis? Yes No
4. Has the beneficiary been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
5. Has the beneficiary had a trial and failure of Humira or a clinical reason beneficiary cannot try Humira? Yes No
5. Has the beneficiary flad a trial and failure of numina of a chilical reason beneficiary calmot try numina? — 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

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any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.