SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

Drug Requested: Simponi® (golimumab) SQ ONLY (Pharmacy)

MEMBER & PRESCRIBER INFORM	IATION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	Fax Number:
NPI #:	
DRUG INFORMATION: Authorization r	nay be delayed if incomplete.
Drug Name/Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight (if applicable):	Date weight obtained:
	ncomitant therapy with more than one biologic ira, Rinvoq, Stelara) prescribed for the same or different. Safety and efficacy of these combinations has NOT been
• Will the member be discontinuing a previousl	y prescribed biologic if approved for requested medication?
	□ Yes OR □ No
• If yes, please list the medication that will be d approval along with the corresponding effective	iscontinued and the medication that will be initiated upon ve date.
Medication to be discontinued:	Effective date:
Medication to be initiated:	Effective date:

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

	Diagnosis: Moderate-to-Severe Rheumatoid Arthritis Dosing: SubQ: 50 mg once a month (in combination with methotrexate)					
	1 M	Member has a diagnosis of moderate-to-severe rho	eumat	toid arthritis		
	P 1	rescribed by or in consultation with a Rheumato	logist			
		Member has tried and failed at least ONE of the following DMARD therapies for at least three (3) months				
		hydroxychloroquine				
		l leflunomide				
		1 methotrexate				
		l sulfasalazine				
	1 M	Tember meets ONE of the following:				
		Member tried and failed, has a contraindication biologics below (verified by chart notes or pl	-			
		☐ Preferred adalimumab product		Enbrel [®]		
		□ Rinvoq®/Rinvoq® LQ		Preferred tocilizumab product: Actemra® SC or Tyenne® SC		
		□ Xeljanz [®] /XR [®]				
		Member has been established on Simponi® SQ indicates at least a 90-day supply of Simponi (verified by chart notes or pharmacy paid cl	SQ v	vas dispensed within the past 130 days	story	
Diagnosis: Active Psoriatic Arthritis Dosing: SubQ: 50 mg once a month (either alone or in combination with methotrexate or other non-biologic DMARDs)						
C	1 M	Member has a diagnosis of active psoriatic arthri	tis			
	P 1	rescribed by or in consultation with a Rheumato	logist			
	☐ Member has tried and failed at least <u>ONE</u> of the following DMARD therapies for at least <u>three (3)</u> <u>months</u>					
		l cyclosporine				
		l leflunomide				
		1 methotrexate				
		ı sulfasalazine				

(Continued on next page)

		mber meets <u>ONE</u> of the following: Member tried and failed, has a contrai	ndication, or intoleran	ce to TWO of	the PREFERRED
		biologics below (verified by chart no			
			□ Enbrel [®]	□ Otezla [®]	□ Rinvoq®/ Rinvoq® LQ
		☐ Preferred adalimumab product	□ Skyrizi [®]	□ Stelara [®]	□ Taltz [®]
			□ Xeljanz®/XR®	□ Tremfya	®
		Member has been established on Simpindicates at least a 90-day supply of a (verified by chart notes or pharmac	Simponi SQ was disp		
Do	osin	nosis: Active Ankylosing Spone g: SubQ: 50 mg once a month (either tic DMARDs)	~	on with metho	trexate or other non-
	Me	mber has a diagnosis of active ankylos	sing spondylitis		
	Pre	scribed by or in consultation with a Rh	eumatologist		
	Me	mber tried and failed, has a contraindic	cation, or intolerance t	o <u>TWO</u> NSAI	Ds
		mber meets ONE of the following:			
		Member tried and failed, has a contrainabiologics below (verified by chart no			the PREFERRED
		☐ Preferred adalimumab product	□ Enbrel®		1 Rinvoq®
		□ Taltz [®]	□ Xeljanz [®] /XR	R	
		Member has been established on Simp			
		indicates at least a 90-day supply of some (verified by chart notes or pharmac)		ensed within	the past 130 days
			,		
□ Diagnosis: Moderate-to-Severe Active Ulcerative Colitis Dosing: SubQ: Induction: 200 mg at week 0, then 100 mg at week 2, followed by maintenance therapy of 100 mg every 4 weeks					
	Me	mber has a diagnosis of moderate-to-se	evere active Ulcerativ	e Colitis	
	☐ Prescribed by or in consultation with a Gastroenterologist				
		(Con	tinued on next page)		

ш	Μŧ	ember meets <u>ONE</u> of the following:		
		Member has tried and failed budesonide or high dose steroids (40-60 mg prednisone)		
		Member has tried and failed at least ONE of the following DMARD therapies for at least three (3)		
	<u>months</u>			
		☐ 5-aminosalicylates (balsalazide, olsalazine, sulfasalazine)		
		□ oral mesalamine (Apriso, Asacol/HD, Delzicol, Lialda, Pentasa)		
	Member meets ONE of the following:			
		Member tried and failed, has a contraindication, or intolerance to ONE preferred adalimumab product		
		Member has been established on Simponi® SQ for at least 90 days <u>AND</u> prescription claims history indicates <u>at least a 90-day supply of Simponi SQ was dispensed within the past 130 days</u>		
		(verified by chart notes or pharmacy paid claims)		

Medication being provided by a Specialty Pharmacy - Proprium Rx

Not all drugs may be covered under every Plan

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

**Use of samples to initiate therapy does not meet step edit/preauthorization criteria. **

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.