SENTARA COMMUNITY PLAN (MEDICAID)

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request</u>. 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>

<u>Drug Requested</u>: Taltz[®] SQ (ixekizumab) (self-administered, Pharmacy) (Non-Preferred)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.					
Member Name:					
Member Sentara #:	Date of Birth:				
Prescriber Name:					
Prescriber Signature:					
Office Contact Name:					
Phone Number:					
NPI #:					
DRUG INFORMATION: Authorization may be	be delayed if incomplete.				
Drug Name/Form/Strength:					
Dosing Schedule:	Length of Therapy:				
Diagnosis:	ICD Code, if applicable:				
Weight (if applicable):	Date weight obtained:				
Recommended Dose:					
Indication	Dosage				
☐ Moderate-to-Severe Plaque Psoriasis (Adults) — who are candidates for systemic therapy or phototherapy	 Two 80mg injections initially for 14 days, Then, two 80mg/28days until week 12 Then, one 80mg injection every 28 days 				
☐ Moderate-to-Severe Plaque Psoriasis (Children ≥ 6 years) - who are candidates for systemic therapy or phototherapy	In pediatrics weight based: • >50 kg: Two 80 mg injections/28 days initially, then one 80 mg injection/28 days • 25-50 kg: Two 40 mg injections/28 days initially, then one 40 mg injection/28 days • <25kg: Two 20 mg injections/28 days initially, then one 20 mg injection/28 days				
☐ Active Psoriatic Arthritis (PsA)	Two 80mg injections/28days, then one 80mg injection every 4 weeks				

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□ A	ctive Ankylosing Spondylitis		• Two 80mg inject injection every 4		s/28days, then one 80mg	
sp	ctive non-radiographic axial pondyloarthritis (nr-axSpA) with ogns of inflammation (Adults)	bjective			subcutaneously every 4 weeks	
suppo	NICAL CRITERIA: Check belort each line checked, all documentated ded or request may be denied.					
DIAGNOSIS: (Check below the applicable diagnosis or authorization will be delayed or denied)						
□ Moderate to Severe Plaque Psoriasis						
	Moderate-to-severe active Chronic Plaque Psoriasis who are candidates for systemic therapy or phototherapy					
	Member is at least 6 years of age or older					
	Trial and failure of at least <u>TWO (2)</u> topical treatments, such as corticosteroids, calcipotriene, coal tar, tazarotene, or anthralin. List drugs below:					
	1		2			
	Trial and failure of, contraindication	n, or adverse	reaction to methotrex	ate		
	Trial and failure of TWO (2) of the PREFERRED drugs below:					
	☐ Humira [®]	□ Enbrel®			Infliximab	
	□ Active Psoriatic Arthritis (PsA)					
	Trial and failure of, contraindication, or adverse reaction to methotrexate					
	Trial and failure of TWO (2) of the PREFERRED drugs below:					
	□ Humira [®]	□ Enbrel®			Infliximab	
	□ Ankylosing Spondylitis					
	Trial and failure of, contraindication, or adverse reaction to methotrexate					
	Trial and failure of TWO (2) of the PREFERRED drugs below:					
	☐ Humira [®]	□ Enbrel®			Infliximab	

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□ Non-Radiographic Axial Spondyloarthritis					
	Member has objective signs of inflammation				
	Trial and failure of, contraindication, or adverse reaction to methotrexate				
	Trial and failure of TWO (2) of the PREFERRED drugs below:				
	□ Humira [®]	□ Enbrel [®]			
Medication being provided by Specialty Pharmacy - PropriumRx					

^{**}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. *