## Prior Authorization Request Form for tocilizumab subcutaneous ( Actemra SC )



## JOHNS HOPKINS HEALTHCARE

7231 Parkway Drive, Suite 100, Hanover, MD 21076

FAX Completed Form and Applicable Progress Notes to: (410) 424-4037

## **USFHP Pharmacy Prior Authorization Form**

To be completed by Requesting provider				
Drug Name:	Strength:			
Dosage/Frequency (SIG):	Duration of Therapy:			

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step	Please complete patient and physician information (please print):					
1	Patient Name:	otient Name:				
	Address		Addross:			
	Sponsor ID #		Phone #:			
	Date of Birth:	Secure Fax #:				
Step 2	Please complete clinical assessment:					
	Humira is the Department of Defe immune biologic. Has the patient		☐ Yes proceed to question 3	☐ No proceed to question 2		
	Is the patient an adult (18 years of diagnosis of giant cell arteritis?	age or older) with a	☐ Yes proceed to question 13	☐ No proceed to question 5		
	3. Has the patient had an inadequate	e response to Humira?	☐ Yes proceed to question 6	☐ No proceed to question 4		
	Has the patient experienced an acthat is not expected to occur with		☐ Yes proceed to question 6	□ No STOP Coverage not approved		
	5. Does the patient have a contraind (adalimumab)?	ication to Humira	☐ Yes proceed to question 6	□ No STOP Coverage not approved		
	6. Is the patient 18 years of age or o	lder?	☐ Yes proceed to question <b>7</b>	☐ No proceed to question 8		
	7. What is the indication or diagnosis?	□ moderate to severely active <b>rheumatoid arthritis</b> – proceed to question <b>10</b> □ other indication or diagnosis – <b>STOP: Coverage not approved</b> .				

	8. Is the patient 2 years of age o	r older?	☐ Yes proceed to question 9	□ No STOP Coverage not approved		
,	9. What is the indication or	□ active polyarticular Juvenile Idiopathic Arthritis (pJIA) – proceed to question 12				
	diagnosis?	☐ active systemic Juvenile	□ active systemic Juvenile Idiopathic Arthritis (sJIA) – proceed to question 12			
		☐ other indication or diagnosis – <b>STOP: coverage not approved.</b>				
	10. Has the patient had an inadeq disease modifying anti-rheum		☐ Yes proceed to question 11	□ No STOP Coverage not approved		
	<ul> <li>11. Does the patient have platelets less than 100,000/mm³ or liver transaminases above 1.5 time UNL?</li> <li>12. Does the patient have evidence of a negative TB test result in the past 12 months (or TB is adequately managed)?</li> <li>13. Will the patient be receiving other targeted immunomodulatory biologics with Actemra, including but not limited to the following: Cimzia, Cosentyx, Enbrel, Humira, Ilumya, Kevzara, Kineret, Olumiant, Orencia, Otezla, Remicade, Rituxan, Siliq, Simponi, Stelara, Taltz, Tremfya or Xeljanz/Xeljanz XR?</li> <li>Note: Subcutaneous Actemra is not approved for use in cytokine re</li> </ul>		☐ Yes STOP Coverage not approved	□ No proceed to question 12		
			☐ Yes proceed to question 13	☐ No STOP Coverage not approved		
			☐ Yes STOP Coverage not approved	□ No Sign and date below		
			release syndrome.			
Step 3	I certify the above is true to the	lease sign and date:				
	Prescriber Sig	gnature	Date			
				[24 April 2019]		
For Inte	ernal Use Only					
Appr	Approved:		Duration of Approval:month(s)			
Deni	ied:	Authorized By:				
Incor	mplete/Other:	PA#:				
Date Fa	axed to MD:	Date Decision Rendered:				