## US Family Health Plan Prior Authorization Request Form for baricitinib (**Olumiant**)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD. Humira is the Department of Defense's preferred targeted biologic agent for FDA approved indications.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

Step	Please complete patient and physician information (please	print):		
1		hysician Name:		
. •	Address: Address:			
	Sponsor ID #	Phone #:		
	Date of Birth:	Secure Fax #:		
Step 2	Please complete clinical assessment:			
	Humira is the Department of Defense's preferred targeted biologic agent. Has the patient tried Humira?	☐ Yes proceed to question 2	□ No proceed to question 4	
	Has the patient had an inadequate response to Humira?	☐ Yes	□ No	
		proceed to question 5	proceed to question 3	
	3. Has the patient experienced an adverse reaction to Humira that is not expected to occur with the requested agent?	☐ Yes	□ No	
		proceed to question 5	STOP	
			Cov erage not approved	
	4. Does the patient have a contraindication to Humira (adalimumab)?	☐ Yes	□ No	
		Proceed to question 5	STOP	
			Cov erage not approved	
	5Is the patient 18 years of age or older?	□ Yes	□ No	
		proceed to question 6	STOP	
			Cov erage not approved	
	6. Does the patient have a diagnosis of moderately to severely active rheumatoid arthritis?	□ Yes	□ No	
		proceed to question 7	STOP	
			Cov erage not approved	
	7. Has the patient had an inadequate response to non-	☐ Yes	□ No	
	biologic systemic therapy? (For example: methotrexate, aminosalicylates [for example,	proceed to question 8	STOP	
	sulfasalazine, mesalamine], corticosteroids,		Cov erage not approved	
	immunos uppres sants [for example, azathioprine], etc.)?			
	8. Will the patient be receiving other biologic DMARDs or potent immunosuppressants (for example, azathioprine and cyclosporine) at the same time	□ Yes	□ No	
		STOP	proceed to question 9	
	(concomitantly)?	Cov erage not approved		

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9. Does the patient have a history of throm boembolic disease?	□ Yes	□ No		
	STOP	proceed to question 10		
	Cov erage not approved			
Is the provider aware of the FDA safety alerts AND Boxed Warnings?	□ Yes	□ No		
	proceed to question 11	STOP		
		Cov erage not approved		
11. Does the patient have a hemoglobin (Hgb) less than 8 g/dL?	□ Yes	□ No		
	STOP	proceed to question 12		
	Cov erage not approved			
12. Does the patient have an absolute neutrophil count (ANC) LESS	□ Yes	□ No		
THAN 1,000/mm3?	STOP	proceed to question 13		
	Cov erage not approved			
13. Does the patient have an absolute lymphocyte count (ALC) LESS	□ Yes	□ No		
THAN 500/mm3?	STOP	proceed to question <b>14</b>		
	Cov erage not approved			
Does the patient have evidence of a negative TB test result in past	□ Yes	□ No		
12 months (or TB is adequately managed)?	proceed to question <b>15</b>	STOP		
		Cov erage not approved		
15. Will the patient be receiving other targeted immunomodulatory	□ Yes	□ No		
biologics, with Olumiant, including but not limited to the following:	STOP	Sign and date below		
Actemra, Cimzia, Cosentyx, Enbrel, Humira, Ilumya, Kevzara, Kineret, Orencia, Remicade, Rituxan, Siliq, Simponi, Stelara, Taltz,	Cov erage not approved			
Tremfya, Skyrizi, Rinvoq, or Xeljanz/Xeljanz XR? (Note: does not				
apply to Otezla)				
Step				
3 I certify the above is true to the best of my knowledge. Please sign and date:				
i dorany and above to the booter my anomought house sign and date.				
Prescriber Signature	Date			
i resember dignature	Dato	[16 March 2022]		