

US Family Health Plan

Prior Authorization Request Form for anakinra (**Kineret**)

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.

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 1 Please complete patient and physician information (please print):

1	Patient Name: _____	Physician Name: _____
	Address: _____	Address: _____
	Sponsor ID #: _____	Phone #: _____
	Date of Birth: _____	Secure Fax #: _____

Step 2 Please complete the clinical assessment:

2	1. How old is the patient?	<input type="checkbox"/> Pediatric patient (less than 18 years old) - Proceed to question 2 <input type="checkbox"/> Adult patient (18 years of age or older) - Proceed to question 3	
	2. What is the indication or diagnosis in this pediatric patient?	<input type="checkbox"/> Neonatal Onset Multisystem Inflammatory Disease (NOMID), a subset of Cryopyrin-Associated Periodic Syndrome (CAPS) - Proceed to question 9 <input type="checkbox"/> Systemic Juvenile Idiopathic Arthritis (sJIA) - Proceed to question 9 <input type="checkbox"/> Deficiency of Interleukin-1 Receptor Antagonist (DIRA) - Proceed to question 9 <input type="checkbox"/> Other - STOP Coverage not approved	
	3. Humira is the Department of Defense's preferred targeted biologic agent. Has the patient tried Humira?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No Proceed to question 6
	4. Has the patient had an inadequate response to Humira?	<input type="checkbox"/> Yes Proceed to question 7	<input type="checkbox"/> No Proceed to question 5
	5. Has the patient experienced an adverse reaction to Humira that is not expected to occur with the requested agent?	<input type="checkbox"/> Yes Proceed to question 7	<input type="checkbox"/> No STOP Coverage not approved
	6. Does the patient have a contraindication to Humira (adalimumab)?	<input type="checkbox"/> Yes Proceed to question 7	<input type="checkbox"/> No STOP Coverage not approved
	7. What is the indication or diagnosis in this adult patient?	<input type="checkbox"/> Moderate to severe active rheumatoid arthritis - Proceed to question 8 <input type="checkbox"/> Other - STOP Coverage not approved	

USFHP Prior Authorization Request Form for
anakinra (**Kineret**)

<p>8. Has the patient had an inadequate response to 1 or more non-biologic systemic therapy (for example: methotrexate, aminosalicylates [for example, sulfasalazine, mesalamine], corticosteroids, immunosuppressants [for example, azathioprine])?</p>	<p><input type="checkbox"/> Yes Proceed to question 9</p>	<p><input type="checkbox"/> No STOP Coverage not approved</p>
<p>9. Does the patient have a negative TB test result in the past 12 months (or is TB adequately managed)?</p>	<p><input type="checkbox"/> Yes Proceed to question 10</p>	<p><input type="checkbox"/> No STOP Coverage not approved</p>
<p>10. Will the patient be receiving other targeted immunomodulatory biologics with Kineret, including but not limited to the following: adalimumab (Humira), etanercept (Enbrel), certolizumab (Cimzia), golimumab (Simponi), infliximab (Remicade), apremilast (Otezla), ustekinumab (Stelara), abatacept (Orencia), tocilizumab (Actemra), tofacitinib (Xeljanz/Xeljanz XR), rituximab (Rituxan), secukinumab (Cosentyx), ixekizumab (Taltz), brodalumab (Siliq), sarilumab (Kevzara), guselkumab (Tremfya), baricitinib (Olumiant), tildrakizumab (Ilumya), risankizumab-rzaa (Skyrizi), or upadacitinib (Rinvoq ER)?</p>	<p><input type="checkbox"/> Yes STOP Coverage not approved</p>	<p><input type="checkbox"/> No Sign and date below</p>

Step 3 I certify the above is true to the best of my knowledge. Please sign and date:

_____ Prescriber Signature

_____ Date