

**USFHP Prior Authorization Request Form for
upadacitinib (Rinvoq)**

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

<https://www.usfamilyhealth.org/for-providers/pharmacy-information/>

For Atopic Dermatitis, prior authorization expires after 12 months. Renewal PA criteria will be approved indefinitely. For renewal of therapy an initial Tricare prior authorization approval is required.

For Rheumatoid Arthritis, Psoriatic Arthritis, Polyarticular Juvenile Idiopathic Arthritis, Ankylosing Spondylitis, Non-Radiographic Axial Spondyloarthritis, Ulcerative Colitis, Crohn's Disease, or Giant cell arteritis the prior authorization is approved indefinitely.

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 clinical assessment:

2	<p>1. Has the patient received this medication under the TRICARE benefit in the last 6 months for treatment of atopic dermatitis? Please choose "No" if the patient did not previously have a TRICARE approved PA for Rinvoq.</p> <p>2. For atopic dermatitis, has the patient's disease severity improved and stabilized to warrant continued therapy?</p> <p>3. What is the indication or diagnosis?</p>	<p><input type="checkbox"/> Yes (subject to verification) proceed to question 2</p> <p><input type="checkbox"/> Yes (subject to verification) Sign and date below</p> <p><input type="checkbox"/> Moderate To Severely Active Rheumatoid Arthritis – proceed to question 4</p> <p><input type="checkbox"/> Active Psoriatic Arthritis – proceed to question 4</p> <p><input type="checkbox"/> Ankylosing Spondylitis – proceed to question 4</p> <p><input type="checkbox"/> Non-Radiographic Axial Spondyloarthritis – proceed to question 4</p> <p><input type="checkbox"/> Active Polyarticular Juvenile Idiopathic Arthritis – proceed to question 4</p> <p><input type="checkbox"/> Moderately To Severely Active Ulcerative Colitis – proceed to question 4</p> <p><input type="checkbox"/> Moderately To Severely Active Crohn's Disease – proceed to question 4</p>	<p><input type="checkbox"/> No proceed to question 3</p> <p><input type="checkbox"/> No STOP Coverage not approved</p>
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	<input type="checkbox"/> Giant cell arteritis – proceed to question 8 <input type="checkbox"/> Moderate to severe atopic Dermatitis – proceed to question 8 <input type="checkbox"/> Other indication or diagnosis – STOP: Coverage not approved	
4. Humira is the Department of Defense's preferred targeted biologic agent. Has the patient tried Humira?	<input type="checkbox"/> Yes proceed to question 5	<input type="checkbox"/> No proceed to question 7
5. Has the patient had an inadequate response to Humira?	<input type="checkbox"/> Yes proceed to question 8	<input type="checkbox"/> No proceed to question 6
6. 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 8	<input type="checkbox"/> No STOP Coverage not approved
7. Does the patient have a contraindication to Humira (adalimumab)?	<input type="checkbox"/> Yes proceed to question 8	<input type="checkbox"/> No STOP Coverage not approved
8. What is the patient's age?	<input type="checkbox"/> 18 years of age or older –proceed to question 9 <input type="checkbox"/> 17 years of age or younger –proceed to question 10	
9. What is the indication or diagnosis for this adult patient?	<input type="checkbox"/> Moderate to severe active rheumatoid arthritis – proceed to question 11 <input type="checkbox"/> Active psoriatic arthritis - proceed to question 12 <input type="checkbox"/> Ankylosing spondylitis – proceed to question 14 <input type="checkbox"/> Non-radiographic axial spondyloarthritis– proceed to question 14 <input type="checkbox"/> Active polyarticular juvenile idiopathic arthritis– proceed to question 23 <input type="checkbox"/> Moderately to severely active ulcerative colitis- proceed to question 23 <input type="checkbox"/> Moderately to severely active Crohn's disease - proceed to question 23 <input type="checkbox"/> Giant cell arteritis - proceed to question 16 <input type="checkbox"/> Moderate to severe Atopic Dermatitis - proceed to question 19 <input type="checkbox"/> Other indication or diagnosis – STOP: Coverage not approved	
10. What is the indication or diagnosis for this pediatric patient?	<input type="checkbox"/> Active psoriatic arthritis - proceed to question 12 <input type="checkbox"/> Active polyarticular juvenile idiopathic arthritis– proceed to question 23 <input type="checkbox"/> Moderate to severe Atopic Dermatitis - proceed to question 18 <input type="checkbox"/> Other indication or diagnosis – STOP: Coverage not approved	

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11. Has the patient had an inadequate response or adverse reaction or a contraindication to Xeljanz or Xeljanz XR or Olumiant?	<input type="checkbox"/> Yes proceed to question 13	<input type="checkbox"/> No STOP Coverage not approved
12. Has the patient experienced an inadequate response or adverse reaction or contraindication to Xeljanz or Xeljanz XR?	<input type="checkbox"/> Yes proceed to question 13	<input type="checkbox"/> No STOP Coverage not approved
13. Has the patient had an inadequate response, intolerance, or contraindication to non-biologic systemic therapy (for example, methotrexate, sulfasalazine, mesalamine, corticosteroids, immunosuppressants)?	<input type="checkbox"/> Yes proceed to question 23	<input type="checkbox"/> No STOP Coverage not approved
14. Has the patient experienced an inadequate response to at least TWO nonsteroidal anti-inflammatory drugs (e.g. ibuprofen, naproxen, diclofenac) over a period of at least two months?	<input type="checkbox"/> Yes proceed to question 15	<input type="checkbox"/> No STOP Coverage not approved
15. Has the patient experienced an inadequate response or adverse reaction or contraindication to Taltz?	<input type="checkbox"/> Yes proceed to question 23	<input type="checkbox"/> No STOP Coverage not approved
16. Is the requested medication being prescribed by or in consultation with a rheumatologist?	<input type="checkbox"/> Yes proceed to question 17	<input type="checkbox"/> No STOP Coverage not approved
17. Has the patient experienced an inadequate response or adverse reaction or contraindication to BOTH of the following: (a) glucocorticoid AND (b) tocilizumab (Tysen)?	<input type="checkbox"/> Yes proceed to question 23	<input type="checkbox"/> No STOP Coverage not approved
18. Is the patient 12 years of age or older?	<input type="checkbox"/> Yes proceed to question 19	<input type="checkbox"/> No STOP Coverage not approved
19. Is the requested medication prescribed by a dermatologist, allergist, or immunologist?	<input type="checkbox"/> Yes proceed to question 20	<input type="checkbox"/> No STOP Coverage not approved
20. Does the patient have a contraindication to, intolerance to, or have they failed treatment with ONE medication in EACH of the following two categories: <ul style="list-style-type: none"> • Topical Corticosteroids: Patients 12 to 17 years of age, can be any topical corticosteroid. Patients 18 years or older, high potency/class 1 topical corticosteroids (e.g., clobetasol propionate 0.05% ointment/cream) is required. • Topical calcineurin inhibitor (for example, pimecrolimus, tacrolimus)? 	<input type="checkbox"/> Yes proceed to question 21	<input type="checkbox"/> No STOP Coverage not approved
21. Is the patient's disease adequately controlled with other systemic drug products including biologics (for example, Dupixent)?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No proceed to question 22
22. Does the patient have a contraindication to, intolerance to, inability to access treatment, or has failed treatment with Narrowband UVB phototherapy?	<input type="checkbox"/> Yes proceed to question 23	<input type="checkbox"/> No STOP Coverage not approved

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23. Will the patient be receiving other targeted immunomodulatory biologics with Rinvoq, except for Otezla, including but not limited to the following: TNF inhibitors, IL-1, IL-6, IL-17, IL-23, IL-36, S1p, or JAK inhibitors?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Sign and date below
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Step 1 I certify the above is true to the best of my knowledge. Please sign and date:

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Prescriber Signature

Date

[29 October 2025]