TRICARE Prior Authorization Request Form for
upadacitinib **(Rinvoq ER)**

JOHNS HOPKINS
HEALTH PLANS
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.

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.

Step 1

Please complete patient and physician information (please print):

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

Step 2

Please complete clinical assessment:

1. Is the requested medication being used for non-radiographic axial spondyloarthritis, rheumatoid arthritis, psoriatic arthritis, ulcerative colitis, ankylosing spondylitis, or Crohn's disease?	☐ Yes proceed to question **2**	☐ No proceed to question **6**
2. Humira is the Department of Defense's preferred targeted biologic agent. Has the patient tried Humira?	☐ Yes proceed to question **3**	☐ No proceed to question **5**
3. Has the patient had an inadequate response to Humira?	☐ Yes proceed to question **6**	☐ No proceed to question **4**
4. Has the patient experienced an adverse reaction to Humira that is not expected to occur with the requested agent?	☐ Yes proceed to question **6**	☐ No **STOP** **Coverage not approved**
5. Does the patient have a contraindication to Humira (adalimumab)?	☐ Yes proceed to question **6**	☐ No **STOP** **Coverage not approved**

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6. What is the indication or diagnosis?	☐ Moderate to severe active rheumatoid arthritis – proceed to question **7** ☐ Moderate to severe atopic dermatitis - proceed to question **14** ☐ Active psoriatic arthritis (PsA) - proceed to question **9** ☐ Moderately to severely active ulcerative colitis - proceed to question **12** ☐ Moderately to severely active Crohn's disease - proceed to question **13** ☐ Ankylosing spondylitis – proceed to question **22** ☐ Non-radiographic axial spondyloarthritis – proceed to question **21** ☐ Other indication or diagnosis – **STOP: Coverage not approved**	
7. The provider acknowledges that for rheumatoid arthritis a trial of Xeljanz or Olumiant is required before Rinvoq.	_____ Proceed to question **8**	
8. Has the patient experienced an inadequate response or adverse reaction to Xeljanz OR Xeljanz XR OR Olumiant?	☐ Yes proceed to question **11**	☐ No proceed to question **10**
9. Has the patient experienced an inadequate response or adverse reaction to Xeljanz OR Xeljanz XR?	☐ Yes proceed to question **11**	☐ No proceed to question **10**
10. Does the patient have a contraindication to Xeljanz OR Xeljanz XR OR Olumiant?	☐ Yes proceed to question **11**	☐ No **STOP** **Coverage not approved**
11. Has the patient had an inadequate response or an intolerance to methotrexate or other nonbiologic disease-modifying antirheumatic drugs (DMARDs)?	☐ Yes proceed to question **13**	☐ No **STOP** **Coverage not approved**
12. Has the patient had an inadequate response to non-biologic systemic therapy (for example – methotrexate, aminosalicylates (e.g. sulfasalazine, mesalamine), corticosteroids, immunosuppressants (e.g. azathioprine), etc?	☐ Yes proceed to question **13**	☐ No **STOP** **Coverage not approved**
13. Is the patient 18 years of age or older?	☐ Yes proceed to question **27**	☐ No **STOP** **Coverage not approved**
14. Has the patient received this medication under the TRICARE benefit in the last 6 months? *Please choose "No" if the patient did not previously have a TRICARE approved PA for Rinvoq ER.*	☐ Yes (subject to verification) proceed to question **15**	☐ No proceed to question **16**

15. For atopic dermatitis, has the patient's disease severity improved and stabilized to warrant continued therapy?	☐ Yes (subject to verification) **Sign and date below**	☐ No **STOP** **Coverage not approved**
16. Is the patient greater than or equal to 12 year(s) of age?	☐ Yes proceed to question **17**	☐ No **STOP** **Coverage not approved**
17. Is the requested medication prescribed by a dermatologist, allergist, or immunologist?	☐ Yes proceed to question **18**	☐ No **STOP** **Coverage not approved**
18. Is the patient's disease adequately controlled with other systemic drug products including biologics (for example, Dupixent)?	☐ Yes **STOP** **Coverage not approved**	☐ No proceed to question **19**
19. Does the patient have a contraindication to, intolerability to, or have they failed treatment with ONE medication in EACH of the following two categories: • **Topical Corticosteroids AND** NOTE: **For patients 18 years of age or older,** high potency/class 1 topical corticosteroids (for example, clobetasol propionate 0.05% ointment/cream, fluocinonide 0.05% ointment/cream) is required. **For patients 12 to 17 years of age,** can be any topical corticosteroid. • **Topical calcineurin inhibitor (for example, pimecrolimus, tacrolimus)**	☐ Yes proceed to question **20**	☐ No **STOP** **Coverage not approved**
20. Does the patient have a contraindication to, intolerability to, inability to access treatment, or has failed treatment with Narrowband UVB phototherapy?	☐ Yes proceed to question **27**	☐ No **STOP** **Coverage not approved**
21. Does the patient have active non-radiographic axial spondyloarthritis?	☐ Yes proceed to question **22**	☐ No **STOP** **Coverage not approved**
22. Is the patient 18 years of age or older?	☐ Yes proceed to question **23**	☐ No **STOP** **Coverage not approved**
23. Has the patient experienced an inadequate response to Cosentyx?	☐ Yes proceed to question **26**	☐ No proceed to question **24**
24. Has the patient experienced an adverse reaction to Cosentyx that is not expected to occur with the requested agent?	☐ Yes proceed to question **26**	☐ No proceed to question **25**

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25. Does the patient have a contraindication to Cosentyx?	☐ Yes proceed to question **26**	☐ No **STOP** **Coverage not approved**
26. Has the patient experienced an inadequate response to at least TWO NSAIDs (for example: ibuprofen, naproxen, diclofenac) over a period of at least two months?	☐ Yes proceed to question **27**	☐ No **STOP** **Coverage not approved**
27. Is the provider aware of the FDA safety alerts AND Boxed Warnings?	☐ Yes proceed to question **28**	☐ No **STOP** **Coverage not approved**
28. Does the patient have a hemoglobin level LESS THAN 8 g/dL?	☐ Yes **STOP** **Coverage not approved**	☐ No proceed to question **29**
29. Does the patient have an absolute neutrophil count (ANC) LESS THAN 1,000/mm^3?	☐ Yes **STOP** **Coverage not approved**	☐ No proceed to question **30**
30. Does the patient have an absolute lymphocyte count (ALC) LESS THAN 500/mm^3?	☐ Yes **STOP** **Coverage not approved**	☐ No proceed to question **31**
31. Will the patient be receiving other targeted immunomodulatory biologics with Rinvoq ER, except for Otezla, including but not limited to the following: Actemra, Cimzia, Cosentyx, Enbrel, Humira, Ilumya, Kevzara, Kineret, Olumiant, Orencia, Remicade, Rituxan, Siliq, Stelara, Taltz, Xeljanz or Xeljanz XR or Tremfya and other potent immunosuppressant's (for example: azathioprine, cyclosporine)?	☐ Yes **STOP** **Coverage not approved**	☐ No proceed to question **32**
32. Does the patient have a history of venous thromboembolic (VTE) disease?	☐ Yes **STOP** **Coverage not approved**	☐ No proceed to question **33**
33. Does the patient have evidence of an active TB infection within the past 12 months?	☐ Yes **STOP** **Coverage not approved**	☐ No **Sign and date below**

Step 3

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

Prescriber Signature _____ Date _____

[06 December 2023]

For Internal Use Only	
☐ Approved:	Duration of Approval: _____month(s)
☐ Denied:	Authorized By:
☐ Incomplete/Other:	PA#:
Date Faxed to MD:	Date Decision Rendered: