US Family Health Plan Prior Authorization Request Form for Mirikizumab-mrkz (Omvoh Pen and Syringe)

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

| | documentation may be required for review. | | | | | |
|------|---|------------------------------|--|--|--|--|
| | thorization does not expire. | | | | | |
| Step | Please complete patient and physician information (please print): | | | | | |
| 1 | Patient Name: Phys | sician Name: | | | | |
| | Address: | Address: | | | | |
| | O 15 " | | | | | |
| | Sponsor ID # S Date of Birth: S | Phone #: ecure Fax #: | | | | |
| Ston | | ecule rax #. | | | | |
| Step | Please complete the clinical assessment: | | | | | |
| 2 | 1. Is the patient 18 years of age or older? | ☐ Yes | □ No | | | |
| | | Proceed to question 2 | STOP | | | |
| | | | Coverage not approved | | | |
| | Does the patient have moderate to severely active ulcerative colitis? | ☐ Yes | □ No | | | |
| | | Proceed to question 3 | STOP | | | |
| | | · | Coverage not approved | | | |
| | 3. Humira is the Department of Defense's preferred targeted biologic agent for ulcerative colitis. Has the patient tried Humira? | □ Yes | □ No | | | |
| | | Proceed to question 4 | Proceed to question 6 | | | |
| | | Proceed to question 4 | Floceed to question o | | | |
| | | | | | | |
| | 4. Has the patient had an inadequate response to Humira? | ☐ Yes | □ No | | | |
| | | Proceed to question 7 | Proceed to question 5 | | | |
| | 5. Has the patient experienced an adverse reaction to Humira that is not expected to occur with the requested agent? | ☐ Yes | □ No | | | |
| | | Proceed to question 7 | STOP | | | |
| | | | Coverage not approved | | | |
| | 6. Does the patient have a contraindication to | □ Yes | □ No | | | |
| | Humira (adalimumab)? | Proceed to question 7 | STOP | | | |
| | | 1 locced to question 7 | Coverage not approved | | | |
| | 7. Has the patient had an inadequate response to non-biologic systemic therapy? (For example: methotrexate, aminosalicylates [such as, sulfasalazine, mesalamine], corticosteroids, immunosuppressants [for example, azathioprine], | | | | | |
| | | ☐ Yes | □ No | | | |
| | | Proceed to question 8 | STOP | | | |
| | | | Coverage not approved | | | |
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| | 8. | Does the patient have evidence of a negative TB test result in the past 12 months (or TB is adequately managed)? | ☐ Yes Proceed to question 9 | □ No STOP Coverage not approved | | |
|--------|---------|--|----------------------------------|---------------------------------|--|--|
| | 9. | Will the patient be receiving any other targeted immunomodulatory biologics with mirikizumab including but not limited to the following: certolizumab (Cimzia), etanercept (Enbrel), golimumab (Simponi), infliximab (Remicade), apremilast (Otezla), ustekinumab (Stelara), abatacept (Orencia), anakinra (Kineret), tocilizumab (Actemra), tofacitinib (Xeljanz/Xeljanz XR), rituximab (Rituxan), secukinumab (Cosentyx), ixekizumab (Taltz), brodalumab (Siliq), sarilumab (Kevzara), guselkumab (Tremfya), baricitinib (Olumiant), tildrakizumab (Ilumya), risankizumab (Skyrizi), upadacitinib (Rinvoq ER), or vedolizumab (Entyvio)? | ☐ Yes STOP Coverage not approved | □ No Sign and date below | | |
| Step 3 | l certi | tify the above is true to the best of my knowledge. Please sign and date: | | | | |
| | | Prescriber Signature | Date | [13 November 2024] | | |