Blue Cross Blue Shield/Blue Care Network of Michigan Medication Authorization Request Form



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This form is to be used by participating physicians to obtain coverage for **drugs covered under the medical benefit**. For <u>commercial members only</u>, please complete this form and submit via fax to 1-877-325-5979. If you have any questions regarding this process, please contact BCBSM Provider Relations and Servicing or the Medical Drug Helpdesk at 1-800-437-3803 for assistance.

| PATIENT INFORMATION | PHYSICIAN INFORMATION | | | |
|---|--------------------------------|--|--|--|
| Name | Name | | | |
| ID Number | Specialty | | | |
| D.O.B/ MM/DD/YYYY | Address | | | |
| Diagnosis | City /State/Zip | | | |
| Drug Name Skyrizi IV | Phone: Fax: | | | |
| Dose and Quantity | NPI | | | |
| Directions | Contact Person | | | |
| Date of Service(s) | Contact Person Phone / Ext. | | | |
| STEP 1: DISEASE STATE INFORMATION | | | | |
| Required Demographic Information: Patient Weight:kg Patient Height:ftinches | | | | |
| Will the provider be administering the medication to the FEP member within the health plan's geographic service area? \[\sum \text{Yes} \sum \text{No} \text{ No} \text{if No, a prior authorization is not required through this process.} \] | | | | |
| Prior authorizations are required for FEP members that will be serviced by a provider within the health plan's geographic service area. If you are not a provider in the geographic service area, please contact the health plan for questions regarding the FEP member's benefit requirements. | | | | |
| Is this member's FEP coverage primary or secondary coverage? ☐ If primary, continue with questionset. ☐ If secondary, an authorization is not needed through this process. Please contact the member's primary coverage fo determination of benefit and additional information. | | | | |

| | | <u>Duestions:</u> | | | |
|--|--|--|--|--|--|
| 1. | Is this requ | est for initiation of therapy with Skyrizi? \(\sigma\)Yes \(\sigma\)No | | | |
| 2. | Will the par | ient's initial induction dosing be administered by intravenous in | fusion at week 0, week 4, and week 8? □Yes □No | | |
| 3. | | patient's diagnosis? disease (CD) | | | |
| | b. Doe | s the patient have moderately to severely active Crohn's disease is the prescriber agree to administer the initial induction IV infus e colitis (UC) | | | |
| | a. Doe b. Doe | s the patient have moderately to severely active ulcerative colities the prescriber agree to administer the initial induction IV infus gnosis (<i>please specify</i>): | ons within the dose of 1200 mg? □Yes □No | | |
| 4. | Does the p | rescriber agree to monitor liver enzymes and bilirubin levels for | hepatotoxicity? | | |
| 5. | | es the patient have an intolerance or contraindication or have they had an inadequate treatment response to at least one conventional rapy option? No | | | |
| 6. | Has the patient been tested for latent tuberculosis (TB)? □Yes* □No *If YES, was the result of the test positive or negative for TB infection? □Negative □Positive* *If POSITIVE, has the patient completed treatment or is the patient currently receiving treatment for latent TB? □Yes □No | | | | |
| 7. Does the patient have any active infections including tuberculosis (TB) or hepatitis B virus (HBV)? Yes No | | | | | |
| 8. Will the patient be given live vaccines while on Skyrizi? □Yes □No | | | | | |
| 9. Will Skyrizi be used in combination with another biologic *disease-modifying antirheumatic drug (DMARD) or targeted synthetic DMARD? □Yes* □No *If YES, please specify medication: *DMARDs: Actemra, Avsola, Cimzia, Cosentyx, Enbrel, Entyvio, Humira, Ilumya, Inflectra, Kevzara, Kineret, Olumiant, Orencia, Otezla, Remicade, Renflexis, Riabni, Rinvoq, Rituxan, Ruxience, Siliq, Simponi/Simponi Aria, Stelara, Taltz, Tremfya, Truxima, and Xeljanz/Xeljanz XR | | | | | |
| | | | | | |
| Chart notes are required for the processing of all requests. Please add any other supporting medical information necessary for our review (required) | | | | | |
| | Coverage will not be provided if the prescribing physician's signature and date are not reflected on this document. Request for expedited review: I certify that applying the standard review time frame may seriously jeopardize the life or health of the member or the member's ability to regain maximum function | | | | |
| | ıysician's Nan | | Date | | |
| | tep 2: | ☐ Form Completely Filled Out ☐ Provide chart notes | ☐ Attach test results | | |
| | tep 3: | By Fax: BCBSM Specialty Pharmacy Mailbox | By Mail: BCBSM Specialty Pharmacy Program | | |
| | ıbmit | 1-877-325-5979 | P.O. Box 312320, Detroit, MI 48231-2320 | | |