Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

| MEMBER'S LAST NAME: | MEMBER'S FIRST NAME: | | | | | |
|---|--|---------------------------|----------------|--------------------------------|--|--|
| Instructions: Please fill out all important for the review (e.g., this form is Protected Health I | chart notes or lab data, to su | | • | | | |
| MEMBER INFORMATION | | | | | | |
| LAST NAME: | | FIRST NAME: | | | | |
| DUONE NUMBER | | | | | | |
| PHONE NUMBER: | | DATE OF BIRTH: | DATE OF BIRTH: | | | |
| STREET ADDRESS: | | | | | | |
| CITY: | | STATE: | ZIP CODE: | | | |
| PATIENT INSURANCE ID NUM | MBER: | | | | | |
| MALE FEMALE HEIGHT OF THE PRESCRIPTION OF THE | BER, YOU WILL NEED TO SUBMIT A PHI DISCI | OSURE AUTHORIZATION FORM | WITH THIS REQU | JEST WHICH CAN BE FOUND AT THE | | |
| PRESCRIBER INFORMATION | | | | | | |
| LAST NAME: | | FIRST NAME: | | | | |
| PRESCRIBER SPECIALTY: | | EMAIL ADDRESS: | | | | |
| NPI NUMBER: | DEA NUMBER: | | | | | |
| PHONE NUMBER: | | FAX NUMBER: | | | | |
| STREET ADDRESS: | | | | | | |
| CITY: | STATE: ZIP CODE: | | | | | |
| REQUESTOR (if different than prescriber): | | OFFICE CONTACT PERSON: | | | | |
| | | | | | | |
| MEDICATION OR MEDICAL I | DISPENSING INFORMATION | | | | | |
| MEDICATION NAME: | | | | | | |
| DOSE/STRENGTH: | FREQUENCY: | LENGTH OF THERAPY/REFILLS | : | QUANTITY: | | |
| ☐ NEW THERAPY | RENEWAL | IF RENEWAL: DATE | THERAPY | INITIATED: | | |
| DURATION OF THERAPY (SPE | CIFIC DATES): | | | | | |

Continued on next page.



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| MEMBER'S LAST NAME: | MEMBER'S FIRST NAME: | | | | | |
|---|--|---|--|--|--|--|
| 1. HAS THE PATIENT TRIED ANY OTHER | R MEDICATIONS FOR THIS CONDITION? | YES (if yes, complete below) NO | | | | |
| MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE): | DURATION OF THERAPY (SPECIFY DATES): | RESPONSE/REASON FOR FAILURE/ALLERGY: | | | | |
| 2. LIST DIAGNOSES: | | ICD-10: | | | | |
| □ Plaque psoriasis □ Psoriatic Arthritis □ Crohn's Disease □ Ulcerative Colitis | | | | | | |
| □ Other Diagnosis Code(s): | | | | | | |
| 3. REQUIRED CLINICAL INFORMATION PRIOR AUTHORIZATION. | : PLEASE PROVIDE ALL RELEVANT CLINIC | AL INFORMATION TO SUPPORT A | | | | |
| Clinical information: Initial Request for all diagnosis: Is drug being used as part of a clinical trial? Yes No Will the patient be using Skyrizi concurrently with another biologic or other immunomodulatory agents? Yes No | | | | | | |
| Has the patient had a 3-month trial and inadequate response to the biosimilar for Humira-adalimumab-aacf? No *Must provide documentation, including trial dates. | | | | | | |
| Does patient have a absolute contraindication to the biosimilar for the biosimilar for Humira-adalimumab-aacf? □ Yes □ No Please submit documentation. | | | | | | |
| | nadequate response to a 4- month No Please submit documentati | | | | | |
| Does patient have a absolute con ☐ Yes ☐ No Please submit docu | traindication to the biosimilar for mentation. | Stelara- <u>Otulfi(ustekinumb-aauz</u>)? | | | | |
| Initial Request for Plaque Psorias Is prescriber a dermatologist? | | | | | | |
| | overing at least 10% of their body s palms, soles, head and neck, or ge | | | | | |
| • | e response to topical therapy (e.g., s | • | | | | |
| Select if the patient has had a tria | al and inadequate response to the f JVA) □ UVB with coal tar | following phototherapy options: | | | | |
| Select if the nationt has had a tria | al and inadequate response to the f | following systemic theranies: | | | | |



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| MEMBER'S LAST NAI | ИЕ: | | MEMBER'S FIRST NAME: |
|---|---|-----------------------|--|
| □ Acitretin □ C dates. | yclosporine | □ Methotrexate | *Must provide documentation, including trial |
| Does the patient No *Must provide do | | ntation of a contrain | dication to all oral systemic therapies? □ Yes □ |
| • | ermatologist or e using Skyriz | r rheumatologist? | □ Yes □ No another biologic or other immunomodulatory |
| Does the patient | have documer | nted active disease? | P □ Yes □ No *Must provide documentation |
| agents (DMARDs cyclosporine)? | , e.g., methotro Yes □ No | | py with oral disease modifying anti-rheumatic (Azulfidine), leflunamide(Arava), or |
| Select if the patie following system Glucocortice Methotrexate Azathioprine 6-mercaptop 5-ASA/mesa | ent has tried ar ic therapies: oid therapy e ourine lamine | · | er the following: the response, intolerance, or contraindication to the the response intolerance, or contraindication to the |
| Initial Request fo | r Ulcerative Co | · | |
| | | i concurrently with a | another biologic or other immunomodulatory |
| agents ? — Yes Select if the patie following system Glucocortice Methotrexate | nt has tried ar ic therapies: oid therapy | nd had an inadequat | e response, intolerance, or contraindication to the |
| □ Azathioprine | | | |
| □ 6-mercapto | | | |
| | | umentation, includir | ng which agent(s) have been tried and trial dates: |
| Renewal Reques Is the prescriber gastroenterologist checked rheumatologist Will the patient b | one of the belo | | another biologic or other immunomodulatory |
| agents ? - Yes | | i John Griden William | another biologic of other initiationioudiatory |



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|--|---|-------------------|
| Is the patient continuing to have a positive clinical documentation | response? □ Yes □ No *Must provid | le |
| Please note: Not all drugs/diagnosis are covered on al required information is received. | I plans. This request may be denied unle | ess all |
| ATTESTATION: I attest the information provided is true understand that the Health Plan, insurer, Medical Grou request the medical information necessary to verify the | p or its designees may perform a routine | audit and |
| Prescriber Signature or Electronic I.D. Verification: | | Date: |
| CONFIDENTIALITY NOTICE: The documents accompanying this transmissi you are not the intended recipient, you are hereby notified that any discl of these documents is strictly prohibited. If you have received this inform and arrange for the return or destruction of these documents. | osure, copying, distribution, or action taken in reliance | e on the contents |

FAX THIS FORM TO: 800-424-7640

MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program

Attn: CP – 4201 P.O. Box 64811 St. Paul, MN 55164-0811

