## HEALTHY U MEDICAID

### PRIOR AUTHORIZATION REQUEST FORM Non-Radiographic Axial Spondyloarthritis (nrx-SpA)

Avsola<sup>®</sup>, Cimzia<sup>®</sup>, Cosentyx<sup>®</sup>, Inflectra<sup>®</sup>, infliximab, Remicade<sup>®</sup>, Renflexis<sup>®</sup>, Taltz<sup>®</sup>

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department.

- For Medical Pharmacy please fax requests to: 801-213-1547
- For **Retail Pharmacy** please fax requests to: 385-425-4052

#### Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for Pharmacy Customer Service for assistance at 385-425-5094

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

| Date:         | Member Name: | ID#:            |
|---------------|--------------|-----------------|
| DOB:          | Gender:      | Physician:      |
| Office Phone: | Office Fax:  | Office Contact: |

#### Height/Weight:

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

#### Preferred/Non-Formulary:

- 1. 1st Line Preferred agents:
  - A. Infliximab products: Avsola<sup>®</sup> (infliximab-axxq), Inflectra<sup>®</sup> (infliximab-dyyb), infliximab, Remicade (infliximab), Renflexis<sup>®</sup> (infliximab-abda)
- 2. 2<sup>nd</sup> line preferred agents with single step; after trial and failure of 1 preferred first line agent:
  - A. Cimzia<sup>®</sup> (certolizumab), Taltz<sup>®</sup> (ixekizumab)
- 3. Non-Formulary agents with a triple step; after trial and failure of 1 preferred first line agent and 2 second line agents:
  - A. Cosentyx<sup>®</sup> (secukinumab)

#### Product being requested: \_\_\_\_\_

| Dosing/Frequency:_ |
|--------------------|
|--------------------|

| If the request is for reauthorization, proceed to reauthorization section.   |  |    |                              |  |
|--|--|----|------------------------------|--|
| Questions  |  | No | Comments/Notes               |  |
| 1. Is the member 18 years of age or older with Non-Radiographic Axial Spondyloarthritis?   |  |    | Please provide documentation |  |
| 2. Is the requesting provider a rheumatologist or in consultation with one?  |  |    |                              |  |
| 3. Does documentation show an adequate trial and failure of at<br>least one prescription strength nonsteroidal anti-inflammatory<br>drug (NSAID) at the maximally tolerated dose, unless<br>contraindicated? |  |    | Please provide documentation |  |
| 4. Has the provider performed tuberculosis (TB) screening prior to therapy initiation?   |  |    | Please provide documentation |  |

| 5. For tumor necrosis factor inhibitors (TNFIs), has the provider preformed Hepatitis B screening prior to therapy initiation? |  |  | Please provide documentation |  |  |  |
|--|--|--|------------------------------|--|--|--|
|  |  |  |                              |  |  |  |
| 1. Is the request for reauthorization of therapy?  |  |  |                              |  |  |  |
| 2. Does updated documentation show that the member has a continued medical need?   |  |  | Please provide documentation |  |  |  |
| 3. Has the provider performed continued tuberculosis screening during therapy?   |  |  | Please provide documentation |  |  |  |
| 4. Has the provider performed continued Hepatitis B monitoring in HBV carriers?  |  |  | Please provide documentation |  |  |  |
| name of treatment, reason for failure, treatment dates, etc.<br>Additional information:  |  |  |                              |  |  |  |
| Physician Signature:   |  |  |                              |  |  |  |

# \*\* Failure to submit clinical documentation to support this request will result in a dismissal of the request.\*\*

Policy: PHARM-HU-142 Origination Date: 07/28/2022 Reviewed/Revised Date: 08/24/2022 Next Review Date: 08/24/2023 Current Effective Date: 09/01/2022

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