

UnitedHealthcare Pharmacy Clinical Pharmacy Programs

Program Number	2025 P 1498-1
Program	Prior Authorization/Non-Formulary
Medications	Brand Humira
P&T Approval Date	10/2025
Effective Date	1/1/2026

1. Background:

This program requires the provider to validate that the member is not an appropriate candidate for Humira biosimilars.

2. Coverage Criteria^a:

A. Brand Humira

- 1. **Brand Humira** will be approved based on **ONE** of the following:
 - a. Submission of medical records documenting patient allergy or demonstrated intolerance to the inactive ingredients in Adalimumab-adaz (unbranded Hyrimoz), and Amjevita

-OR-

- b. **Both** of the following:
 - (a) Submission of medical records documenting patient has previously been successfully treated with brand Humira
 - (b) Submission of medical records documenting patient has tried Adalimumab-adaz (unbranded Hyrimoz), and Amjevita for 6-8 weeks ^b per product with a decrease in effectiveness

-OR-

- c. **Both** of the following:
 - (a) Prescriber is requesting continuation of therapy for an established member on Humira (as documented by a paid claim of at least 28 days supply in past 120 days)

-AND-

(b) Dose does not exceed the following limits based on indication:

For rheumatoid arthritis or hidradenitis suppurativa: 40mg per week or 80mg every other week

For Crohn's disease, ulcerative colitis, or uveitis: 40mg per week For polyarticular juvenile idiopathic arthritis, psoriatic arthritis, plaque psoriasis,



or ankylosing spondylitis: 40mg every other week

Authorization will be issued for 12 months.

^a State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

^b For Connecticut, Kentucky and Mississippi business, only a 30-day trial will be required

3. Additional Clinical Rules:

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.
- Supply limits may be in place.

4. References:

N/A

Program	Prior Authorization/Non-Formulary – Brand Humira
Change Control	
10/2025	New program