



**Request for Prior Authorization
BIOLOGICALS FOR
PLAQUE PSORIASIS**

(PLEASE PRINT – ACCURACY IS IMPORTANT)

FAX Completed Form To
1 (877) 733-3195
Provider Help Desk
1 (844) 236-1464

IA Medicaid Member ID # _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _		Patient name		DOB	
Patient address					
Provider NPI _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _			Prescriber name		Phone
Prescriber address				Fax	
Pharmacy name		Address			Phone
Prescriber must complete all information above. It must be legible, correct, and complete or form will be returned.					
Pharmacy NPI _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _			Pharmacy fax		NDC _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _ _

Prior authorization (PA) is required for biologicals used for plaque psoriasis. Request must adhere to all FDA approved labeling for requested drug and indication, including age, dosing, contraindications, warnings and precautions, drug interactions, and use in specific populations. Payment for non-preferred biologicals for plaque psoriasis will be considered only for cases in which there is documentation of previous trials and therapy failures with two preferred biological agents. Payment will be considered under the following conditions:

- 1. Patient has a diagnosis of moderate to severe plaque psoriasis; and**
- 2. Patient has documentation of an inadequate response to phototherapy, systemic retinoids, methotrexate, or cyclosporine.**

The required trials may be overridden when documented evidence is provided that use of these agents would be medically contraindicated.

Preferred

- | | |
|--|---|
| <input type="checkbox"/> Adalimumab-aacf | <input type="checkbox"/> Simlandi |
| <input type="checkbox"/> Adalimumab-adbm | <input type="checkbox"/> Skyrizi Auto-Injector |
| <input type="checkbox"/> Adalimumab-fkjp | <input type="checkbox"/> Skyrizi Cartridge |
| <input type="checkbox"/> Amjevita 40mg/0.4mL | <input type="checkbox"/> Skyrizi Prefilled Syringe |
| <input type="checkbox"/> Amjevita 80mg/0.8mL | <input type="checkbox"/> Taltz (step through one preferred TNF) |
| <input type="checkbox"/> Enbrel | <input type="checkbox"/> Tremfya |
| <input type="checkbox"/> Humira | <input type="checkbox"/> Yusimry |

Non-Preferred

- | |
|---|
| <input type="checkbox"/> Bimzelx |
| <input type="checkbox"/> Cimzia |
| <input type="checkbox"/> Cosentyx |
| <input type="checkbox"/> Siliq |
| <input type="checkbox"/> Stelara |
| <input type="checkbox"/> Other Humira Biosimilar: _____ |

Strength _____	Dosage Instructions _____	Quantity _____	Days Supply _____
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Diagnosis: _____

Treatment failure with a preferred oral therapy: Trial Drug Name: _____

Trial start date: _____ Trial end date: _____

Failure reason: _____

Non-Pharmacological Treatments Tried: _____

Trial start date: _____ Trial end date: _____

Failure reason: _____

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Medical or contraindication reason to override trial requirements: _____

Other medical conditions to consider: _____

Possible drug interactions/conflicting drug therapies: _____

Attach lab results and other documentation as necessary.

Prescriber signature (Must match prescriber listed above.)	Date of submission
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IMPORTANT NOTE: *In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Health and Human Services, that the member continues to be eligible for Medicaid.*