

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.

**This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.
Allow at least 24 hours for review.**

Section A – Member Information

First Name:	Last Name:	Member ID:
Address:		
City:	State:	ZIP Code:
Phone:	DOB:	Allergies:
Primary Insurance Information (if any):		
Is the requested medication: <input type="checkbox"/> New or <input type="checkbox"/> Continuation of Therapy? If continuation, list start date: _____		
Is this patient currently hospitalized? <input type="checkbox"/> Yes <input type="checkbox"/> No If recently discharged, list discharge date: _____		

Section B - Provider Information

First Name:	Last Name:	M.D./D.O.
Address:	City:	State: ZIP code:
NPI #:	Phone:	Fax: Specialty:
Office Contact Name / Fax attention to:		

Section C - Medical Information

Medication:	Strength:
Directions for use:	Quantity:
Diagnosis (Please be specific & provide as much information as possible):	ICD-10 CODE:
Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____	

Section D – Previous Medication Trials

Medication Name	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation

**Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs:
Please refer to the patient's PDL for a list of preferred alternatives**

Member First name:	Member Last name:	Member DOB:
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Clinical and Drug Specific Information

ALL REQUESTS:

- **What is the patient's diagnosis? (check which applies)**

- Moderate to Severe Plaque Psoriasis Active Ankylosing Spondylitis
 Active Psoriatic Arthritis Other, **List:** _____

- **Is the patient currently on Cosentyx therapy?** Yes No

- **Did the patient have a history of failure, contraindication, or intolerance to either of the following:** Yes No
(check which applies) Humira (adalimumab) Enbrel (etanercept)
(If yes, complete Section D above with medication information, including dose, date of trial, and reason for discontinuation)

- **Is the patient receiving Cosentyx in combination with any of the following:** Yes No

- Biologic DMARD [e.g., Humira (adalimumab), Cimzia (certolizumab), Simponi (golimumab), Taltz (ixekizumab), Orencia (abatacept)]
 Janus kinase inhibitor [e.g., Xeljanz (tofacitinib)]
 Phosphodiesterase 4 (PDE4) inhibitor [e.g., Otezla (apremilast)]

Requests for CONTINUATION OF THERAPY:

- **Is there documentation of a positive clinical response to Cosentyx therapy?** Yes No
If yes, list positive response: _____

Provider Signature: _____ **Date:** _____

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