

Cosentyx® Prior Authorization Request Form (Page 1 of 2)

Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information <small>(required)</small>					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information <small>(required)</small>					
Select the diagnosis below: <input type="checkbox"/> Active ankylosing spondylitis <input type="checkbox"/> Active psoriatic arthritis <input type="checkbox"/> Moderate to severe plaque psoriasis <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
For all diagnoses, answer the following: Is this request for continuation of prior Cosentyx therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if Cosentyx is prescribed by or in consultation with one of the following specialists: <input type="checkbox"/> Dermatologist <input type="checkbox"/> Rheumatologist Will Cosentyx be used in combination with a biologic DMARD [e.g., Enbrel (etanercept), Humira (adalimumab), Cimzia (certolizumab), Simponi (golimumab)]? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if the patient has history of failure, contraindication, or intolerance to the following, if applicable for the patient's diagnosis: <input type="checkbox"/> Cimzia (certolizumab) <input type="checkbox"/> Simponi (golimumab) <input type="checkbox"/> Taltz (ixekizumab) <input type="checkbox"/> Humira (adalimumab) <input type="checkbox"/> Stelara (ustekinumab)					
For active ankylosing spondylitis, also answer the following: Does the patient have history of failure, contraindication, or intolerance to two non-steroidal anti-inflammatory drugs (NSAIDs)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Reauthorization: If this is a reauthorization request, answer the following questions: Is there documentation the patient has had a positive clinical response to Cosentyx therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Will Cosentyx be used in combination with a biologic DMARD [e.g., Enbrel (etanercept), Humira (adalimumab), Cimzia (certolizumab), Simponi (golimumab)]? <input type="checkbox"/> Yes <input type="checkbox"/> No					

Cosentyx® Prior Authorization Request Form (Page 2 of 2)

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
Please fax this form to 1-844-712-8129 to initiate a prior authorization review for the member and medication above.
Please note: plan benefits may limit or exclude coverage of specific medications including those requested on this form.

I certify, to the best of my knowledge, the statements and information provided on this form are factual and correct.

Provider/Representative (and Title): _____ Date: _____

PROACT INTERNAL USE ONLY:

Clinical Review Decision

Approved, through

Denied (documentation attached, if necessary)

Tracking:

1 st Attempt		2 nd Attempt		Letter Mailed:	
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