



Fax completed form to: (855) 840-1678

If this is an URGENT request, please call (800) 882-4462
(800.88.CIGNA)

Cosentyx Intravenous (secukinumab)

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician's Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on this form are completed.*		
Specialty:	* DEA, NPI or TIN:				
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:		* Date of Birth:
Office Fax:			* Patient Street Address:		
Office Street Address:			City	State	Zip
City	State	Zip	Patient Phone:		
Urgency: <input type="checkbox"/> Standard <input type="checkbox"/> Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)					
Medication requested: <input type="checkbox"/> Cosentyx 125mg/5ml IV Dose and Quantity: Duration of therapy: J-Code: Frequency of administration: ICD10: What is your patient's current weight? Is this a new start or continuation of therapy with the requested medication? If patient has been taking samples, please pick "new start." <input type="checkbox"/> new start of therapy <input type="checkbox"/> continuation of therapy If continuation of therapy: (if continuation of therapy) Has the patient demonstrated a beneficial response to this medication? <input type="checkbox"/> Yes <input type="checkbox"/> No (if no) Please provide support for continued use in your patient. <i>(Please note: there are different preferred products depending on your patient's plan. Please refer to the applicable Cigna health care professional resource [e.g. cignaforhcp.com] to determine benefit availability and the terms and conditions of coverage)</i>					
Where will this medication be obtained? <input type="checkbox"/> Accredo Specialty Pharmacy** <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Retail pharmacy <input type="checkbox"/> Other (please specify): <input type="checkbox"/> Home Health / Home Infusion vendor <input type="checkbox"/> Physician's office stock (billing on a medical claim form) **Cigna's nationally preferred specialty pharmacy **Medication orders can be placed with Accredo via E-prescribe - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822 NCPDP 4436920), Fax 888.302.1028, or Verbal 866.759.1557					
Facility and/or doctor dispensing and administering medication: Facility Name: State: Tax ID#: Address (City, State, Zip Code): Where will this drug be administered? <input type="checkbox"/> Patient's Home <input type="checkbox"/> Physician's Office <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Other (please specify): NOTE: Per some Cigna plans, infusion of medication MUST occur in the least intensive, medically appropriate setting. Is this patient a candidate for re-direction to an alternate setting (such as alternate infusion site, physician's office, home) with					

assistance of a Specialty Care Options Case Manager?	<input type="checkbox"/> Yes <input type="checkbox"/> No (provide medical necessity rationale):
Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
What is the indication or diagnosis?	
<input type="checkbox"/> Ankylosing spondylitis (AS) <input type="checkbox"/> Crohn's disease (CD) <input type="checkbox"/> Entesitis-related arthritis <input type="checkbox"/> Non-radiographic axial spondyloarthritis (nr-axSpA) <input type="checkbox"/> Plaque psoriasis (PsO) <input type="checkbox"/> Psoriatic arthritis (PsA) <input type="checkbox"/> Rheumatoid Arthritis (RA) <input type="checkbox"/> other (please specify):	
Clinical Information:	
Will the requested medication be used in combination with a BIOLOGIC or with a targeted synthetic oral small molecule?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
If Ankylosing spondylitis:	
Is the patient currently receiving Cosentyx intravenous or subcutaneous?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient already received at least 6 months of therapy with Cosentyx intravenous or subcutaneous? Please Note: Answer No if the patient has received less than 6 months of therapy or if the patient is restarting therapy with Cosentyx intravenous or subcutaneous.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Is Cosentyx being prescribed by or in consultation with a rheumatologist?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
When assessed by at least one objective measure, has the patient experienced a beneficial clinical response from baseline (prior to initiating Cosentyx intravenous or subcutaneous)? Please Note: Examples of objective measures include Ankylosing Spondylitis Disease Activity Score (ASDAS), Ankylosing Spondylitis Quality of Life Scale (ASQoL), Bath Ankylosing Spondylitis Disease Activity Index (BASDAI), Bath Ankylosing Spondylitis Functional Index (BASFI), Bath Ankylosing Spondylitis Global Score (BAS-G), Bath Ankylosing Spondylitis Metrology Index (BASMI), Dougados Functional Index (DFI), Health Assessment Questionnaire for the Spondyloarthropathies (HAQ-S), and/or serum markers (for example, C-reactive protein, erythrocyte sedimentation rate).	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Compared with baseline (prior to receiving Cosentyx intravenous or subcutaneous), has the patient experienced an improvement in at least one symptom, such as decreased pain or stiffness, or improvement in function or activities of daily living?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
If Non-radiographic axial spondyloarthritis:	
Is the patient currently receiving Cosentyx intravenous or subcutaneous?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient already received at least 6 months of therapy with Cosentyx intravenous or subcutaneous? Please Note: Answer No if the patient has received less than 6 months of therapy or if the patient is restarting therapy with Cosentyx intravenous or subcutaneous.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Does the patient have objective signs of inflammation, defined as C-reactive protein elevated beyond the upper limit of normal for the reporting laboratory?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Does the patient have objective signs of inflammation, defined as sacroiliitis reported on magnetic resonance imaging?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Is Cosentyx being prescribed by or in consultation with a rheumatologist?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
If Psoriatic arthritis:	
Is the patient currently receiving Cosentyx intravenous or subcutaneous?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Has the patient already received at least 6 months of therapy with Cosentyx intravenous or subcutaneous? Please Note: Answer No if the patient has received less than 6 months of therapy or if the patient is restarting therapy with Cosentyx intravenous or subcutaneous.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Is Cosentyx being prescribed by or in consultation with a rheumatologist or a dermatologist?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
When assessed by at least one objective measure, has the patient experienced a beneficial clinical response from baseline (prior to initiating Cosentyx intravenous or subcutaneous)? Please Note: Examples of standardized measures of disease activity include Disease Activity Index for Psoriatic Arthritis (DAPSA), Composite Psoriatic Disease Activity Index (CPDAI), Psoriatic Arthritis Disease Activity Score (PsA DAS), Grace Index, Leeds Entesitis Score (LEI), Spondyloarthritis Consortium of Canada (SPARCC) entesitis score, Leeds Dactylitis Instrument Score, Minimal Disease Activity (MDA), Psoriatic Arthritis Impact of Disease (PsAID-12), and/or serum markers (for example, C-reactive protein, erythrocyte sedimentation rate).	
<input type="checkbox"/> Yes <input type="checkbox"/> No	

Compared with baseline (prior to receiving Cosentyx intravenous or subcutaneous), has the patient experienced an improvement in at least one symptom, such as less joint pain, morning stiffness, or fatigue; improved function or activities of daily living; decreased soft tissue swelling in joints or tendon sheaths)? ☐ Yes ☐ No

Additional Information: *Please provide clinical rationale for the use of this drug for your patient (pertinent patient history, alternatives tried, any inability to use alternatives above or standard therapy, etc). Please include drug name(s), date(s) taken and for how long, and what the documented results were of taking each drug, including any intolerances or adverse reactions your patient experienced.*

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature: _____ **Date:** _____

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