

Fax completed form to: (855) 840-1678
If this is an URGENT request, please call (800) 882-4462 (800.88.CIGNA)

## Skyrizi IV (risankizumab)

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					
Specialty:	* DEA, NPI or	TIN:	form are completed.*					
Office Contact Person:			* Patient Name:					
Office Phone:			* Cigna ID:		* Date of	* Date of Birth:		
Office Fax:			* Patient Street Address:					
Office Street Address:			City	State		Zip		
City	State	Zip	Patient Phone:			1		
Urgency:  ☐ Standard  ☐ 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:  ☐ Skyrizi 600mg/10ml vial ☐ other (please specify):								
Dose and Quantity: Duration of therapy: J-Code:								
Frequency of administration:	ICD10:							
Where will this medication  ☐ Accredo Specialty Pharmac ☐ Hospital Outpatient ☐ Retail pharmacy ☐ Other (please specify):  **Medication orders can be plan NCPDP 4436920), Fax 888.300	☐ Home Health / Home Infusion vendor ☐ Physician's office stock (billing on a medical claim form)  **Cigna's nationally preferred specialty pharmacy e - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822							
Facility and/or doctor dispensing and administering medication: Facility Name: State: Tax ID#: Address (City, State, Zip Code):								
Where will this drug be ad Patient's Home Hospital Outpatient	☐ Physician's Office ☐ Other (please specify):							
<b>NOTE:</b> Per some Cign	a plans, infusion	of medication I	MUST occur in the least intensiv	e, medicall	y appropri	ate 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?  Yes 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?								
What is the indication or diagnosis?								
☐ Crohn's disease (CD)	☐ Crohn's disease (CD) ☐ other (please specify):							

Clinical Information:						
Will the requested medication be used in combination with a BIOLOGIC or with a targeted synthetic oral small molecule drug use an inflammatory condition?  Biologic (an adalimumab product [Humira, biosimilar], Bimzelx, Cimzia, Cosentyx (IV or SC), etanercept SC product [Enbrel, biosimilar], Entyvio (IV or SC), Ilumya, infliximab IV products [Remicade, biosimilar], Kevzara, Kineret, Omvoh (IV or SC), Orencor SC], a rituximab IV product [Rituxan, biosimilar], Skyrizi (SC), Siliq, Simponi [Aria or SC]), an ustekinumab product [Stelara (IV						
SC), biosimlar], Taltz, a tocilizumab product [Actemra (IV or SC), biosimilar], Tremfya (IV or SC), or Zymfentra.  Targeted synthetic oral small molecule drug (such as Cibinqo, Leqselvi, Litfulo, Sotyktu, Olumiant, Otezla, Rinvoq Xeljanz, Xeljanz XR, Velsipity, or Zeposia.)  Conventional synthetic DMARD (such as methotrexate, leflunomide, sulfasalazine, hydroxychloroquine)  No, the requested medication will NOT be used in combination with another BIOLOGIC or targeted synthetic oral drug	•					
If Crohn's disease:						
Will the requested medication be used as induction therapy?	☐ Yes ☐ No					
Has the patient tried or is currently taking a systemic corticosteroid, or is a systemic corticosteroid contraindicated in this patient.						
Has the patient tried one other conventional systemic therapy for Crohn's disease? Please Note: Examples of convertherapy for Crohn's disease include azathioprine, 6-mercaptopurine, or methotrexate. A trial of mesalamine does not systemic agent for Crohn's disease.	ntional systemic					
Has the patient had a previous trial of a biologic used for Crohn's disease? Please Note: Examples of biologics used for Crohn's disease are an adalimumab product (Humira, biosimilars), Cimzia, an infliximab IV product (Remicade, biosimilars), Zymfentra, an ustekinumab product (Stelara [SC or IV], biosimilars), or Entyvio (IV or SC).						
Does the patient have enterocutaneous (perianal or abdominal) or rectovaginal fistulas?	☐ Yes ☐ No					
Has the patient had an ileocolonic resection (to reduce the chance of Crohn's disease recurrence)?	☐ Yes ☐ No					
Is the requested medication being prescribed by or in consultation with a gastroenterologist?	☐ Yes ☐ No					
If Ulcerative colitis:						
Will the requested medication be used as induction therapy?	☐ Yes ☐ No					
Is the requested medication being prescribed by or in consultation with a gastroenterologist?	☐ Yes ☐ No					
<b>Additional pertinent information:</b> 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:						
Save Time! Submit Online at: <a href="https://www.covermymeds.com/main/prior-authorization-forms/cigna/">www.covermymeds.com/main/prior-authorization-forms/cigna/</a> or via SureScri	pts in your EHR.					
Our standard response time for prescription drug coverage requests is 5 business days. If your request is urgent if	is important that					

you call us to expedite the request. View our Prescription Drug List and Coverage Policies online at cigna.com.