

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

Ilumya (tildrakizumab-asmn)

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 Birth:			
Office Fax:			* Patient Street Address:						
Office Street Address:		City	State Zip		Zip				
City	State	Zip	Patient Phone:						
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: ☐ Ilumya 100mg/ml									
Dose and Quantity:	y: J-Code:								
Frequency of administration:	ICD10:								
Is this a new start or continuation of therapy with the requested medication? If patient has been taking samples, please pick "new start".									
(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)									
Where will this medication be obtained? Accredo Specialty Pharmacy** Hospital Outpatient Retail pharmacy Other (please specify):			☐ Home Health / Home Infusion vendor ☐ 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 dispe	ensing and ad	ministering ı	medication:						
Facility Name: Address (City, State, Zip Code):	Sta	ate:		Tax ID#:					
Where will this drug be administered? ☐ Patient's Home ☐ Hospital Outpatient			☐ Physician's Office☐ 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? Yes No (provide medical necessity rationale):									
Is the requested medication for a the patient?	for which the pre	scription medic	ation may	be necess	sary for the life of ☐ Yes ☐ No				

What is the indication or diagnosis? plaque psoriasis other (please specify):							
Clinical Information:							
Will the requested medication be used in combination with a BIOLOGIC or with a targeted synthetic oral small molecule Biologic (an adalimumab product [Humira, biosimilar], Bimzelx, Cimzia, Cosentyx (IV or SC), etanercept SC product [Enbrel, biosimilar], Entyvio (IV or SC), infliximab IV products [Remicade, biosimilar], Kevzara, Kineret, Omvoh (IV or SC), Orencia [IV or SC], a rituximab IV product [Rituxan, biosimilar], Skyrizi (IV or SC), Siliq, Simponi [Aria or SC]), Taltz, a tocilizumab product [Actemra (IV or SC), biosimilar], Tremfya (IV or SC), an ustekinumab product [Stelara (IV or SC), biosimilar], or Zymfentra. Targeted synthetic oral small molecule drug (such as Cibinqo, Leqselvi, Litfulo, Sotyktu, Olumiant, Otezla, Rinvoq, Rinvoq LQ, 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 small molecule drug							
If Plaque psoriasis:							
Is the patient currently receiving the requested medication?	☐ Yes ☐ No						
(if currently receiving the requested medication) Has the patient already received at least 3 months of therapy with the medication? Please Note: Answer No if the patient has received less than 3 months of therapy or if the patient is rest the requested medication.							
(if not currently receiving the requested medication OR has received less than 3 months) Has the patient tried at least systemic agent for psoriasis for at least 3 months, unless intolerant? Please Note: Examples of one traditional systemethotrexate [MTX], cyclosporine, or acitretin tablets. A 3-month trial of psoralen plus ultraviolet A light (PUVA) also							
(if no) Has the patient already had a 3-month trial or previous intolerance to at least one biologic other than drug? Please Note: A biosimilar of the requested biologic does not count. Examples include Bimzelx, an eta [Enbrel, biosimilars], Cosentyx, an adalimumab product [Humira, biosimilars], Cimzia, an infliximab product Remicade, biosimilars], Siliq, Skyrizi, Taltz, Tremfya or an ustekinumab product [Stelara SC or IV, biosimila							
(if no) Does the patient have a contraindication to methotrexate, as determined by the prescriber?	☐ Yes ☐ No ☐ Yes ☐ No						
(if not currently receiving the requested medication OR has received less than 3 months) Is the requested medication by or in consultation with a dermatologist?	being prescribed ☐ Yes ☐ No						
(if currently receiving the requested medication) Has the patient experienced a beneficial clinical response, defined as impr from baseline (prior to initiating the requested medication) in at least one of the following: estimated body surface area, ery induration/thickness, and/or scale of areas affected by psoriasis?							
(if currently receiving the requested medication) Compared with baseline (prior to initiating the requested medication), experienced an improvement in at least one symptom, such as decreased pain, itching, and/or burning?	has the patient ☐ Yes ☐ No						
Has the patient tried ANY of the following? Check ALL that apply. Please Note: Medical documentation specific to you question must be attached to this case or your request could be denied. Documentation may include, but is not limited prescription claims records, and/or prescription receipts.							
an adalimumab product (Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimur adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yyusimry. A trial of multiple adalimumab products counts as ONE product.) Cosentyx Enbrel							
□ Otezla □ Skyrizi subcutaneous □ Sotyktu □ an ustekinumab product (Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otufli, Pyzch ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana and Yesintek. A trial of multiple ustekinumab produ ONE product.) □ Taltz □ Tremfya subcutaneous							
Is the prescriber verifying that the patient has been receiving Ilumya for at least 90 days?	☐ Yes ☐ No						
Is the prescriber verifying that the patient has been receiving Ilumya via paid claims (for example, patient has not been samples or coupons or other types of waivers in order to obtain access to Ilumya)?	n receiving □ Yes □ No						

Additional Information: Please provide clinical rationale for the use of this drug for your patient (pertinent patient history, lternatives tried, any inability to use alternatives above or standard therapy, etc). Please include drug name(s), date(s) taken and for ow long, and what the documented results were of taking each drug, including any intolerances or adverse reactions your patient xperienced.	
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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Our standard response time for prescription drug coverage requests is 5 business days. If your request is urgent, it is important that you call us to expedite the request. View our Prescription Drug List and Coverage Policies online at cigna.com.

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