



Kevzara® (sarilumab) Medication Precertification Request

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(All fields must be completed and legible for Precertification Review.)

Aetna Precertification Notification
503 Sunport Lane, Orlando, FL 32809
Phone: 1-855-240-0535
FAX: 1-877-269-9916

For Medicare Advantage Part B:
FAX: 1-844-268-7263

Please indicate: Start of treatment: Start date ____ / ____ / ____
 Continuation of therapy: Date of last treatment ____ / ____ / ____

Precertification Requested By: _____ **Phone:** _____ **Fax:** _____

A. PATIENT INFORMATION			
First Name: _____		Last Name: _____	
Address: _____		City: _____	State: _____ ZIP: _____
Home Phone: _____		Work Phone: _____	Cell Phone: _____
DOB: _____	Allergies: _____	E-mail: _____	
Current Weight: _____ lbs or _____ kgs		Height: _____ inches or _____ cms	
B. INSURANCE INFORMATION			
Aetna Member ID #: _____		Does patient have other coverage? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Group #: _____		If yes, provide ID#: _____ Carrier Name: _____	
Insured: _____		Insured: _____	
Medicare: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____		Medicaid: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____	
C. PRESCRIBER INFORMATION			
First Name: _____		Last Name: _____ (Check One): <input type="checkbox"/> M.D. <input type="checkbox"/> D.O. <input type="checkbox"/> N.P. <input type="checkbox"/> P.A.	
Address: _____		City: _____	State: _____ ZIP: _____
Phone: _____	Fax: _____	St Lic #: _____	NPI #: _____ DEA #: _____ UPIN: _____
Provider E-mail: _____		Office Contact Name: _____ Phone: _____	
Specialty (Check one): <input type="checkbox"/> Rheumatologist <input type="checkbox"/> Other: _____			
D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION			
Place of Administration:		Dispensing Provider/Pharmacy: <i>Patient Selected choice</i>	
<input type="checkbox"/> Self-administered	<input type="checkbox"/> Physician's Office	<input type="checkbox"/> Physician's Office	<input type="checkbox"/> Retail Pharmacy
<input type="checkbox"/> Outpatient Infusion Center	Phone: _____	<input type="checkbox"/> Specialty Pharmacy	<input type="checkbox"/> Mail Order
Center Name: _____		<input type="checkbox"/> Other: _____	
<input type="checkbox"/> Home Infusion Center	Phone: _____	Name: _____	
Agency Name: _____		Phone: _____ Fax: _____	
<input type="checkbox"/> Administration code(s) (CPT): _____		Address: _____	
Address: _____		TIN: _____ PIN: _____	
E. PRODUCT INFORMATION			
Request is for Kevzara: Dose: _____		Frequency: _____	
F. DIAGNOSIS INFORMATION – Please indicate primary ICD Code and specify any other where applicable.			
Primary ICD Code: _____		Secondary ICD Code: _____ Other ICD Code: _____	
G. CLINICAL INFORMATION – Required clinical information must be completed in its entirety for all precertification requests.			
For Initiation Requests (Clinical documentation required for all requests):			
<input type="checkbox"/> Yes <input type="checkbox"/> No Will sarilumab (Kevzara) be used concomitantly with apremilast, tofacitinib, or other biologic DMARDs (e.g., adalimumab, infliximab)?			
<input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a documented TB test within 6 months of initiating biologic therapy?			
(check all that apply): <input type="checkbox"/> PPD test <input type="checkbox"/> interferon-gamma assay (IGRA) <input type="checkbox"/> chest x-ray			
Please enter the date and results of the TB test: Date: ____ / ____ / ____ Results: <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown			
If positive , does the patient have latent or active TB? <input type="checkbox"/> Latent <input type="checkbox"/> Active			
If latent TB , <input type="checkbox"/> Yes <input type="checkbox"/> No Will TB treatment be started before initiation of therapy with sarilumab (Kevzara)?			
Rheumatoid arthritis			
Please indicate the severity of the patient's rheumatoid arthritis: <input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe			
<input type="checkbox"/> Yes <input type="checkbox"/> No Is there evidence that the disease is active?			
<input type="checkbox"/> Yes <input type="checkbox"/> No Has a trial of a conventional DMARD been ineffective?			
<input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have an intolerance to conventional DMARDs?			
Please select DMARDs tried: <input type="checkbox"/> Cyclophosphamide <input type="checkbox"/> Cyclosporine <input type="checkbox"/> Hydroxychloroquine <input type="checkbox"/> Leflunomide			
<input type="checkbox"/> Methotrexate <input type="checkbox"/> Sulfasalazine <input type="checkbox"/> Other: please explain: _____			

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
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G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

For Continuation of Therapy Requests: (Clinical documentation required for all requests)

- Yes No Will sarilumab (Kevzara) be used concomitantly with apremilast, tofacitinib, or other biologic DMARDs (e.g., adalimumab, infliximab)?
- Yes No Has the patient received samples of sarilumab (Kevzara)? (Sampling of Kevzara does not guarantee coverage under the provisions of the pharmacy benefit)
- Yes No Is there clinical documentation of disease stability or improvement? Disease stability Improvement
- Yes No Does the patient have any risk factors for TB?
 - Yes No Has the patient had a TB test within the past 12 months?
 - (check all that apply): PPD test interferon-gamma assay (IGRA) chest x-ray
 - Please enter the date and results of the TB test: Date: ____ / ____ / ____
 - Results: Positive Negative Unknown

H. ACKNOWLEDGEMENT

Request Completed By (Signature Required): _____ **Date:** ____ / ____ / ____

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.