

## CIGNA HealthCare Prior Authorization Form - Cimzia (Certolizumab pegol) -

Notice: Failure to complete this form in its entirety may result in delayed processing or an adverse determination for insufficient information.

PROVIDER INFORMATION			PATIENT INFORMATION		
* Provider Name:		**Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all			
Specialty:	* DEA or TIN:		asterisked (*) items on this form are completed**		
Office Contact Person:			* Patient Name:		
Office Phone:			* CIGNA ID:		
Office Fax:			* Date Of Birth:		
* Is your fax machine kept in a secure location?  * May we fax our response to your office?  Yes No Yes No Yes No			* Patient Street Address:		
Office Street Address:			City	State	Zip
City	State	Zip	Patient Phone:		
Medication requested:					
☐ Cimzia (certolizumab pegol) 200mg kit ☐ Cimzia (certolizumab pegol) 400mg/2ml syringe kit					
Dose and Quantity: Duration of therapy:			J-Code:		
Frequency of administration:					
Where will this medication be obtained?					
☐ CIGNA Tel-Drug (CIGNA's nationally preferred specialty pharmacy) ☐ Prescriber's office stock (billing on a medical claim form) ☐ Other (please specify):			☐ Retail pharmacy ☐ Home Health / Home Infusion vendor		
Diagnosis related to use:					
☐ Rheumatoid Arthritis ☐ Crohn's Disease ☐ Other (Please specify):					
Chron's Disease:					
Does the patient have a history of positive clinical response to Cimzia (certolizumab pegol) therapy?					
☐ Yes ☐ I	lo				
Did the patient have a failure, contraindication, or intolerance to conventional therapies (e.g. aminosalicylate, corticosteroids, or immumomodulators)?					
☐ Yes ☐ I	lo				
Please indicate if the patient has had evidence of failure, inadequate response, intolerance or contraindication to adalimumab (Humira) or infliximab (Remicade)?					
☐ Yes ☐ N	lo				

Rheumatoid Arthritis:				
Does the patient have a history of positive clinical response to Cimzia (certolizumab pegol) therapy?  ☐ Yes ☐ No				
Please indicate if the patient has had evidence of failure, inadequate response, intolerance or contraindication to any of the following tumor necrosis factor (TNF) antagonists. Please check all that apply:				
Please indicate if the patient has had evidence of failure, inadequate response, intolerance or contraindication to any of the following disease-modifying anti-rheumatic drugs (DMARDs). Please check all that apply:    Methotrexate				
Which of the following methods was used to measure the patient's disease progression PRIOR to therapy on Cimzia? (Check all that apply):    Health Assessment Questionnaire Disease Index (HAQ-DI)				
If this is a request for <b>CONTINUED THERAPY</b> (after at least 16 weeks of treatment), has the patient shown positive response to treatment with Cimzia based on any of the following measurements? (Check all that showed a positive response to Cimzia therapy):    Health Assessment Questionnaire Disease Index (HAQ-DI)				
Additional pertinent information:				
CIGNA HealthCare's coverage position on this and other medications may be viewed online at: <a href="http://www.cigna.com/customer-care/healthcare-professional/coverage-positions">http://www.cigna.com/customer-care/healthcare-professional/coverage-positions</a>				
Please fax completed form to (800)390-9745. Phone requests may be submitted by calling (800)244-6224.				
Our standard response time for prescription drug coverage requests is 2-4 business days. If your request is urgent, it is important that				

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you call Pharmacy Services to expedite the request. View our formulary on line at http://www.cigna.com.