



Prior Authorization Form

Patient Information

Patient's Name:

Insurance ID: Date of Birth: Height: Weight:

Address: Apartment #:

City: State: Zip:

Phone Number: Alternate Phone: Sex: Male Female

Provider Information

Provider's Name: Provider ID Number:

Address: City: State: Zip:

Suite Number: Building Number:

Phone Number: Fax number:

Provider's Specialty:

Medication Information

Medication: Quantity: ICD9 Code:

Directions: Diagnosis: Refills:

Will the physician supply this medication? Yes No

By providing the information it will only be used for coverage determination request administered by OptumRx.

Medication Instructions

Has the patient been instructed on how to Self-Administer? Yes No

Is this medication a New Start? Yes No

If NO please provide the following: Initiation Date: / / Date of Last Dose: / /

This is to notify you that your patient's request for this medication may be denied unless we receive supportive information, i.e., medications tried and failed, document improvement with medication(s). Please provide information to support this request. Please fax back at the number listed above or call at 1-800-711-4555.

Administration Instructions

Dispensing Location: Physician's Office Patient's Address Date medication is needed: / /

Medication Administered: Home Health Self Administered LTC Physician's Office

\*If you have any questions regarding your patient's plan drug limits you may call us at: 1-800-711-4555.

**Enbrel-Humira-Remicade-Orencia-Kineret-Simponi-Cimzia-Actemra-Stelara**

Patients Name: \_\_\_\_\_

**OptumRx**  
**Fax # 1-800-853-3844**

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Patients ID#: \_\_\_\_\_ DOB: \_\_\_\_\_

**OptumRx Specialty Prior Authorization (continued)**

Document the patient's diagnosis: \_\_\_\_\_ ICD-9 Code: \_\_\_\_\_

**Please Document all that applies to the Patient**

Has the patient been evaluated for tuberculosis and treated accordingly?  Yes  No  
Document date of last PPD test: \_\_\_\_\_  Negative  Positive

For Diagnosis of **Rheumatoid Arthritis or Ulcerative Colitis**: Does the Patient exhibit symptoms of MODERATE to SEVERE?  
 Yes  No

For Diagnosis of **Psoriasis**: Does the patient have failure, intolerance or contraindication to: **(Please circle all that apply)**  
**Ultraviolet Light B (UVB), Pulsed Dye Laser, Photochemotherapy, Psoralen and exposure to Ultraviolet light a (PUVA)**

For Diagnosis of **Crohn's Disease**: Does the patient exhibit symptoms of MODERATE to SEVERE?  Yes  No  
Has induction dose been prescribed?  Yes  No (if **NO** document reason why it has not been prescribed)

Document if the patient has tried, failed or had contraindication

- |  |   |
|--|---|
| <input type="checkbox"/> Methotrexate  | <input type="checkbox"/> 6-mercaptopurine (Purnethol) |
| <input type="checkbox"/> Imuran (azathioprine)   | <input type="checkbox"/> NSAIDs (e.g. Ibuprofen)      |
| <input type="checkbox"/> Cyclosporine (Sandimmune, Neoral)                             | <input type="checkbox"/> 6-thioguanine                |
| <input type="checkbox"/> Gold compounds (Myochrisine, Ridura, Aurolate, and Solganal)  | <input type="checkbox"/> Acitretin (soriatane)        |
| <input type="checkbox"/> Plaquenil (hydroxychloroquine)                                | <input type="checkbox"/> Hydroxyurea (hydreia)        |
| <input type="checkbox"/> Arava (leflunomide)   | <input type="checkbox"/> Mycophenolate (cellcept)     |
| <input type="checkbox"/> Cuprimine (penicillamine)                                     | <input type="checkbox"/> Corticosteroids              |
| <input type="checkbox"/> Aminosalicylates (e.g. sulfasalazine, azulfidine, mesalamine) |   |

Please Document Dates of therapies for medications selected: \_\_\_\_\_

Please document any clinical contraindications to these medications: \_\_\_\_\_

Has the patient had a trial, failure or contraindication to any of the following medications? (Please list dosage and/or contraindication)

	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<b>Dosage / Contraindication</b>
<b>Enbrel®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Humira®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Remicaid®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Orencia®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Kineret®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Simponi®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Cimzia®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Actemra®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____
<b>Stelara®</b>	<input type="checkbox"/>	<input type="checkbox"/>	_____

**Continuation of Therapy**

Has the patient utilized the medication in the past 45 days?  Yes  No

Has the patient had documented clinical improvement from ongoing therapy?  Yes  No

**(Please document dose reduction or reason for high dose [if applicable])**

**\*If the above information is not available, please attach the patient's chart notes documenting clinical improvement.**

\*If you have any questions regarding your patient's plan drug limits you may call us at: 1-800-711-4555.

**For UHC members:** Your patient's prescription benefit requires that we review certain requests for coverage with the prescriber. You have prescribed a medication for your patient that requires Prior Authorization before benefit coverage can be provided. Please complete the following questions then fax this form to the toll free number listed below. Upon receipt of the completed form, prescription benefit coverage will be determined based on the plan's rules