



## MEDICARE FORM

### Cimzia® (certolizumab pegol) Injectable Medication Precertification Request

Page 1 of 3

(All fields must be completed and legible for precertification review.)

For Medicare Advantage Part B:  
For other lines of business:  
Please use commercial form.

Note: Cimzia is non-preferred.  
Preferred products vary based on  
indication. See section G below.

Would you like to use electronic prior authorization? Consider using **Availity**, our electronic prior authorization portal. Learn more about **Availity** from the links in the table below.

For phone or fax requests, refer to the table below for routing information. To determine which box to use, refer to the patient's Aetna ID card. State specific special needs and Medicare-Medicaid Plans may be designated on the front of the ID card or in the website URL on the back of the card. If you don't see your specific plan listed, call the number on the back of the member's ID card to confirm routing information.

For **Aetna Medicare Advantage** and **Allina Health Aetna Medicare** members send request to:

**Phone:** [1-866-503-0857](tel:1-866-503-0857) (TTY: [711](tel:1-866-503-0857))

**Fax:** [1-844-268-7263](tel:1-844-268-7263)

**Availity:** <https://www.aetna.com/health-care-professionals/resource-center/availability.html>

For Aetna Medicare Advantage **Virginia Dual Eligible Special Needs Plans** (HMO D-SNP) send request to:

**Phone:** [1-855-463-0933](tel:1-855-463-0933)

**Fax:** [1-833-280-5224](tel:1-833-280-5224)

**Availity:** <https://www.aetnabetterhealth.com/virginia-hmosnp/providers/portal>

For Aetna Assure Premier Plus Medicare Advantage **New Jersey Dual Eligible Special Needs Plans** (HMO D-SNP) send request to:

**Phone:** [1-844-362-0934](tel:1-844-362-0934)

**Fax:** [1-833-322-0034](tel:1-833-322-0034)

**Availity:** <https://www.aetnabetterhealth.com/new-jersey-hmosnp/providers/portal.html>

For Aetna Better Health of **Illinois Premier Medicare Medicaid Plan** (MMP) send request to:

**Phone:** [1-866-600-2139](tel:1-866-600-2139)

**FAX:** [1-855-320-8445](tel:1-855-320-8445)

**Availity:** <https://www.aetnabetterhealth.com/illinois/providers/portal>

For Aetna Better Health of **Ohio Premier Medicare Medicaid Plan** (MMP) send request to:

**Phone:** [1-855-364-0974](tel:1-855-364-0974)

**Fax:** [1-855-734-9389](tel:1-855-734-9389)

**Availity:** <https://www.aetnabetterhealth.com/ohio/providers/portal>

For Aetna Better Health of **Michigan Premier Medicare Medicaid Plan** (MMP) send request to:

**Phone:** [1-855-676-5772](tel:1-855-676-5772)

**Fax:** [1-844-241-2495](tel:1-844-241-2495)

**Availity:** <https://www.aetnabetterhealth.com/michigan/providers/portal.html>



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(All fields must be completed and legible for precertification review.)

For Medicare Advantage Part B:  
For other lines of business:  
Please use commercial form.

Note: Cimzia is non-preferred.  
Preferred products vary based on  
indication. See section G below.

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:		DOB:	
Address:		City:		State:	ZIP:
Home Phone:	Work Phone:	Cell Phone:		Email:	
Patient Current Weight: ____ lbs or ____ kgs Patient Height: ____ inches or ____ cms				Allergies:	
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 Email:		Office Contact Name:		Phone:	
Specialty (Check one): <input type="checkbox"/> Gastroenterologist <input type="checkbox"/> Rheumatologist <input type="checkbox"/> Dermatologist <input type="checkbox"/> Other: _____					
D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION					
<b>Place of Administration:</b> <input type="checkbox"/> Self-administered <input type="checkbox"/> Physician's Office <input type="checkbox"/> Home <input type="checkbox"/> Outpatient Infusion Center Phone: _____ Center Name: _____ <input type="checkbox"/> Home Infusion Center Phone: _____ Agency Name: _____ Address: _____ City: _____ State: _____ ZIP: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____ NPI: _____			<b>Dispensing Provider/Pharmacy: Patient Selected choice</b> <input type="checkbox"/> Physician's Office <input type="checkbox"/> Retail Pharmacy <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Mail Order <input type="checkbox"/> Other: _____ Name: _____ Address: _____ City: _____ State: _____ ZIP: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____ NPI: _____		
E. PRODUCT INFORMATION					
Request is for: <input type="checkbox"/> Cimzia (certolizumab pegol) HCPCS Code: _____					
F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other any other where applicable (*).					
Primary ICD Code: _____		Secondary ICD Code: _____		Other ICD Code: _____	
G. CLINICAL INFORMATION - Required clinical information must be completed for ALL precertification requests.					
<b>For All Requests (clinical documentation required for all requests):</b> What is the patient's diagnosis? <input type="checkbox"/> Moderately to severely active Crohn's disease <input type="checkbox"/> Moderately to severely active rheumatoid arthritis <input type="checkbox"/> Psoriatic arthritis <input type="checkbox"/> Ankylosing spondylitis <input type="checkbox"/> Non-radiographic axial spondyloarthritis <input type="checkbox"/> Polyarticular juvenile idiopathic arthritis <input type="checkbox"/> Moderate to severe plaque psoriasis <input type="checkbox"/> Immune checkpoint inhibitor-related toxicity – inflammatory arthritis <input type="checkbox"/> Other					
<b>For Initiation Requests (clinical documentation required for all requests):</b> Note: Cimzia is non-preferred. Entyvio IV, Inflectra, Renflexis, Simponi Aria and Tremfya IV are preferred for MA plans. For MAPD plans, Cosentyx SC, Enbrel, Humira, Idacio, Rinvoq, Skyrizi, Sotyktu, Stelara, Tremfya, Tyenne SC and Xeljanz/Xeljanz XR are preferred. Preferred products vary based on indication. <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient had prior therapy with Cimzia (certolizumab pegol) within the last 365 days? <input type="checkbox"/> No Has the patient had a trial and failure of any of the following? (if yes, select all that apply below) <input type="checkbox"/> Entyvio IV (vedolizumab) <input type="checkbox"/> Inflectra (infliximab-dyyb) <input type="checkbox"/> Renflexis (infliximab-abda) <input type="checkbox"/> Simponi Aria (golimumab) <input type="checkbox"/> Tremfya IV (guselkumab) → When was the member's trial and failure of the preferred drug? _____ → Please describe the nature of the failure of the preferred drug _____					

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Preferred products vary based  
on indication. See section G.

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 Initiation Requests continued (clinical documentation required for all requests):

- ☐ No Has the patient had an adverse reaction to any of the following? (if yes, select all that apply below)
- ☐ Entyvio IV (vedolizumab) ☐ Inflectra (infliximab-dyyb) ☐ Renflexis (infliximab-abda)
- ☐ Simponi Aria (golimumab) ☐ Tremfya IV (guselkumab)
- When was the member's adverse reaction to the preferred drug? \_\_\_\_\_
- Please describe the nature of the adverse reaction to the preferred drug \_\_\_\_\_
- ☐ No Has the patient had a trial and failure of any of the following? (if yes, select all that apply below)
- ☐ Cosentyx SC (secukinumab) ☐ Enbrel (etanercept) ☐ Humira (adalimumab) ☐ Idacio (adalimumab-aacf) ☐ Rinvoq (upadacitinib)
- ☐ Skyrizi (risankizumab-rzaa) ☐ Sotyktu (deucravacitinib) ☐ Stelara (ustekinumab) ☐ Tremfya (guselkumab)
- ☐ Tyenne SC (tocilizumab-aazg) ☐ Xeljanz/Xeljanz XR (tofacitinib)
- When was the member's trial and failure of the preferred drug? \_\_\_\_\_
- Please describe the nature of the failure of the preferred drug \_\_\_\_\_
- ☐ No Has the patient had an adverse reaction to any of the following? (if yes, select all that apply below)
- ☐ Cosentyx SC (secukinumab) ☐ Enbrel (etanercept) ☐ Humira (adalimumab) ☐ Idacio (adalimumab-aacf) ☐ Rinvoq (upadacitinib)
- ☐ Skyrizi (risankizumab-rzaa) ☐ Sotyktu (deucravacitinib) ☐ Stelara (ustekinumab) ☐ Tremfya (guselkumab)
- ☐ Tyenne SC (tocilizumab-aazg) ☐ Xeljanz/Xeljanz XR (tofacitinib)
- When was the member's adverse reaction to the preferred drug? \_\_\_\_\_
- Please describe the nature of the adverse reaction to the preferred drug \_\_\_\_\_

Please explain if there are any contraindications or other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis (select all that apply)

- ☐ Entyvio IV (vedolizumab) ☐ Inflectra (infliximab-dyyb) ☐ Renflexis (infliximab-abda) ☐ Simponi Aria (golimumab) ☐ Tremfya IV (guselkumab)

Please explain if there are any contraindications or other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis (select all that apply)

- ☐ Cosentyx SC (secukinumab) ☐ Enbrel (etanercept) ☐ Humira (adalimumab) ☐ Idacio (adalimumab-aacf) ☐ Rinvoq (upadacitinib)
- ☐ Skyrizi (risankizumab-rzaa) ☐ Sotyktu (deucravacitinib) ☐ Stelara (ustekinumab) ☐ Tremfya (guselkumab)
- ☐ Tyenne SC (tocilizumab-aazg) ☐ Xeljanz/Xeljanz XR (tofacitinib)

#### ☐ Crohn's disease:

- ☐ Yes ☐ No Has the patient been diagnosed with moderately to severely active Crohn's disease?

#### ☐ Rheumatoid arthritis:

- ☐ Yes ☐ No Has the patient been diagnosed with moderately to severely active rheumatoid arthritis?

#### ☐ Psoriatic arthritis:

- ☐ Yes ☐ No Has the patient been diagnosed with active psoriatic arthritis (PsA)?

#### ☐ Ankylosing spondylitis:

- ☐ Yes ☐ No Has the patient been diagnosed with active ankylosing spondylitis?

#### ☐ Non-radiographic axial spondyloarthritis:

- ☐ Yes ☐ No Has the patient been diagnosed with active non-radiographic axial spondyloarthritis?

#### ☐ Polyarticular juvenile idiopathic arthritis:

- ☐ Yes ☐ No Has the patient been diagnosed with active polyarticular juvenile idiopathic arthritis?

#### ☐ Plaque psoriasis:

- ☐ Yes ☐ No Has the patient been diagnosed with active moderate to severe plaque psoriasis?

#### ☐ For immune checkpoint inhibitor-related toxicity

- ☐ Yes ☐ No Has the patient been diagnosed with severe immunotherapy-related inflammatory arthritis?

#### For Continuation Requests (clinical documentation required for all requests):

- ☐ Yes ☐ No Is the patient receiving benefit from therapy with the requested drug?

### 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.