

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc.
Cimzia (certolizumab pegol) Prior Authorization (PA)
Pharmacy Benefits Prior Authorization Help Desk
Length of Authorizations: Initial- 6 months; Continuation- 12 months

Instructions:

This form is used by Kaiser Permanente and/or participating providers for coverage of Cimzia (certolizumab pegol). Please complete and fax this form back to Kaiser Permanente within 24 hours [fax: <u>1-866-331-2104</u>]. If you have any questions or concerns, please call <u>1-866-331-2103</u>. Requests will not be considered unless this form is complete.

KP-MAS Formulary can be found at: Pharmacy | Community Provider Portal | Kaiser Permanente

| 1 – Patient Information | | | |
|--|---|----------------|--|
| Patient Name: | Kaiser Medical ID#: | Date of Birth: | |
| | 2 – Prescriber Information | | |
| Is the prescriber a Rheumatologist, Der | matologist, or Gastroenterologist? No Y | 'es | |
| If consulted with a specialist, specialist | name and specialty: | | |
| Prescriber Name: | Specialty: | NPI: | |
| Prescriber Address: | | | |
| Prescriber Phone #: | Prescriber Fax #: | | |
| Do you have an approved provider refe ☐ Yes — please provide your provider re | erral number from Kaiser Permanente? eferral number here: | | |
| | 3 – Pharmacy Information | | |
| Pharmacy Name: | Pharmacy NPI: | | |
| Pharmacy Phone # | Pharmacy Fax #: | | |
| | 4 – Drug Therapy Requested | | |
| Drug 1: Name/Strength/Formulation: _ | | | |
| Sig: | | | |
| Drug 2: Name/Strength/Formulation: | | | |
| | | | |
| | | | |

| 5-Diagnosis/Clinical Criteria |
|---|
| 1. Is this request for initial or continuing therapy? □ Initial therapy □ Continuing therapy, state start date: |
| 2. Indicate the patient's diagnosis for the requested medication: |
| Clinical Criteria: |
| Rheumatology: 1. Does the patient have a diagnosis of rheumatoid arthritis, psoriatic arthritis, or spondyloarthropathy? □ No □ Yes |
| 2. If of childbearing potential, is the patient pregnant, attempting to conceive, and/or breastfeeding? □ No □ Yes □ N/A, patient not of childbearing potential |
| 3. Has the patient had an inadequate response, contraindication, or intolerance to at least one of the preferred anti-TNF agents [i.e. adalimumab-atto (Amjevita) or infliximab-dyyb (Inflectra)]? □ No □ Yes |
| Gastroenterology: 1. Does the patient have a diagnosis of Crohn's disease? □ No □ Yes |
| 2. If of childbearing potential, is the patient pregnant, attempting to conceive, and/or breastfeeding? □ No □ Yes □ N/A, patient not of childbearing potential |
| 3. Has the patient had an inadequate response, contraindication, or intolerance to at least one of the preferred anti-TNF agents [i.e. adalimumab-atto (Amjevita) or infliximab-dyyb (Inflectra)]? □ No □ Yes |
| For continuation of therapy, please respond to <u>additional questions</u> below: |
| If of childbearing potential, is the patient still pregnant, attempting to conceive, and/or breastfeeding? □ No □ Yes □ N/A, patient not of childbearing potential |
| 2. Has the patient had a clinically significant benefit from medication (i.e. asymptomatic or in clinical remission)? □ No □ Yes |
| 3. Has specialist follow-up occurred in the past 12 months since last review? □ No □ Yes |
| 6 – Prescriber Sign-Off |
| Additional Information – Please submit chart notes/medical records for the patient that are applicable to this request. If member has not tried preferred agent(s) please provide rationale/explanation and any additional supporting information that should be taken into consideration for the requested medication: |
| I certify that the information provided is accurate. Supporting documentation is available for State audits. |

| Prescriber Signature: | Date: |
|--|-------|
| | |
| Place Note: This document contains confidential information, including protected health information, intended for a specific individual and purpose. The | |

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