

**Entyvio (for Maryland only)
Prior Authorization Request**

Send completed form to: Case Review Unit, CVS Caremark Prior Authorization Fax: 1-866-249-6155

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-866-814-5506**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

The recipient of this fax may make a request to opt-out of receiving telemarketing fax transmissions from CVS Caremark. There are numerous ways you may opt-out: The recipient may call the toll-free number at 877-265-2711, at any time, 24 hours a day/7 days a week. The recipient may also send an opt-out request via email to do_not_call@cvscaremark.com. An opt out request is only valid if it (1) identifies the number to which the request relates, and (2) if the person/entity making the request does not, subsequent to the request, provide express invitation or permission to CVS Caremark to send facsimile advertisements to such person/entity at that particular number. CVS Caremark is required by law to honor an opt-out request within thirty days of receipt.

Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____ **NPI#:** _____
Specialty: _____ **Physician Office Fax:** _____
Physician Office Telephone: _____
Request Initiated For: _____

- Has the patient been diagnosed with any of the following?
 - Moderately to severely active Crohn's disease (CD)
 - Moderately to severely active ulcerative colitis (UC)
 - Other _____
- What is the ICD-10 code? _____

Section A: Preferred Product

- These are the primary preferred products for which coverage is provided for treatment of the following conditions:
 - a) Crohn's disease (CD): **Humira (primary); Cimzia (secondary)***
 - b) Ulcerative colitis (UC): **Humira (primary); Simponi (secondary)***

**Note: Secondary preferred products for CD and UC are Cimzia and Simponi, respectively. These preferred product options only apply to members who have had a documented inadequate response or intolerable adverse event with Humira.*

Can the patient's treatment be switched to a preferred product?

 - Yes - Please specify: _____ *If Yes, please call 1-866-814-5506 to have the updated form faxed to your office OR you may complete the PA electronically (ePA). You may sign up online via CoverMyMeds at: www.covermymeds.com/epa/caremark/ or call 1-866-452-5017.*
 - No
 - Not applicable - Requested for condition not listed above, skip to Section B: All Requests
- Is this request for continuation of therapy with the requested product? Yes No *If No, skip to #6*
- Is the patient currently receiving the requested product through samples or a manufacturer's patient assistance program? If unknown, answer Yes. Yes No *If No, skip to Section B: All Requests*
- Has the patient had a documented inadequate response or intolerable adverse event with any of the following preferred products? Please indicate ALL that apply. **ACTION REQUIRED: If Yes, attach supporting chart note(s).**
 - Cimzia: Inadequate response Intolerable adverse event

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Entyvio MD Step, VF, ACSF SGM - 3/2018.

CVS Caremark is an independent company that provides pharmacy benefit management services to CareFirst BlueCross BlueShield and CareFirst BlueChoice, Inc. members.

CareFirst BlueCross BlueShield is the shared business name of CareFirst of Maryland, Inc. and Group Hospitalization and Medical Services, Inc. CareFirst of Maryland, Inc., Group Hospitalization and Medical Services, Inc., CareFirst BlueChoice, Inc., The Dental Network and First Care, Inc. are independent licensees of the Blue Cross and Blue Shield Association. In the District of Columbia and Maryland, CareFirst MedPlus is the business name of First Care, Inc. In Virginia, CareFirst MedPlus is the business name of First Care, Inc. of Maryland (used in VA by: First Care, Inc.). © Registered trademark of the Blue Cross and Blue Shield Association

- Humira: Inadequate response Intolerable adverse event
- Simponi: Inadequate response Intolerable adverse event
- None of the above, *complete this form in its entirety and also complete Maryland State Step Therapy Section*

7. Does the patient have one of the following documented clinical reasons to avoid TNF inhibitors (Humira, Cimzia or Simponi)? **ACTION REQUIRED: If Yes, attach supporting chart note(s).**
- Yes - History of demyelinating disorder Yes - Autoantibody formation/lupus-like syndrome
 - Yes - History of congestive heart failure Yes - Risk of lymphoma
 - Yes - History of hepatitis B virus infection
 - No - none of the above, *complete this form in its entirety and also complete Maryland State Step Therapy Section*

Section B: All Requests

8. Is this request for continuation of therapy? Yes No *If No, skip to #12*
9. Is the patient currently receiving Entyvio through samples or a manufacturer's patient assistance program?
 Yes No Unknown *If Yes or Unknown, skip to #12*
10. How long has the patient been receiving the requested medication? _____ months
If less than 4 months, no further questions.
11. Has the patient achieved or maintained positive clinical response to treatment as evidenced by low disease activity or improvement in signs and symptoms? *If Yes, no further questions* Yes No
12. Has the patient received any of the following medications?
If Yes, please indicate the most recent medication and skip to diagnosis section.
- Cimzia Humira Inflectra Remicade Renflexis Simponi Stelara Tysabri No
13. Has the patient undergone pretreatment screening for latent tuberculosis (TB) infection with either a TB skin test or an interferon gamma release assay (e.g., QFT-GIT, T-SPOT.TB)? Yes No

Section C: Crohn's Disease

14. Has the patient tried and had an inadequate response to at least one conventional therapy option?
If Yes, indicate below and no further questions.
- | | |
|---|---|
| <input type="checkbox"/> Yes - Sulfasalazine (Azulfidine, Sulfazine) | <input type="checkbox"/> Yes - Azathioprine (Azasan, Imuran) |
| <input type="checkbox"/> Yes - Mesalamine, oral (Asacol, Pentasa, Delzicol, Lialda) | <input type="checkbox"/> Yes - Mercaptopurine (Purinethol) |
| <input type="checkbox"/> Yes - Metronidazole (Flagyl) | <input type="checkbox"/> Yes - Methotrexate |
| <input type="checkbox"/> Yes - Ciprofloxacin (Cipro) | <input type="checkbox"/> Yes - Methylprednisolone (Solu-Medrol) |
| <input type="checkbox"/> Yes - Prednisone | <input type="checkbox"/> Yes - Rifaximin (Xifaxan) |
| <input type="checkbox"/> Yes - Budesonide (Entocort EC) | <input type="checkbox"/> No |
15. Does the patient have a contraindication or intolerance to at least one conventional therapy option (e.g., azathioprine [Azasan, Imuran], budesonide [Entocort EC], ciprofloxacin [Cipro], mesalamine [Asacol, Delzicol, Lialda, Pentasa], mercaptopurine [Purinethol], methylprednisolone [Solu-Medrol], methotrexate, metronidazole [Flagyl], prednisone, sulfasalazine [Azulfidine, Sulfazine], rifaximin [Xifaxan])? *If Yes, no further questions*
 Yes No
16. Has the patient had an inadequate response to a TNF-alpha inhibitor indicated for the treatment of CD (e.g., Cimzia, Humira, Remicade)? *If Yes, no further questions* Yes No
17. Does the patient have a contraindication or intolerance to a TNF-alpha inhibitor indicated for the treatment of CD (e.g., Cimzia, Humira, Remicade)? Yes No

Section D: Ulcerative Colitis

18. Has the patient tried and had an inadequate response to at least one conventional therapy option?
If Yes, indicate below and no further questions.
- Yes - Azathioprine (Azasan, Imuran)
 - Yes - Corticosteroid (e.g., budesonide [Entocort, Uceris], hydrocortisone [Cortifoam, Colocort, Solu-Cortef, Cortef], methylprednisolone [Medrol, Solu-Medrol], prednisone)
 - Yes - Cyclosporine (Sandimmune)

- Yes - Mesalamine (e.g., Asacol, Lialda, Pentasa, Canasa, Rowasa)
- Yes - Mercaptopurine (Purinethol)
- Yes - Sulfasalazine
- Yes - Tacrolimus (Prograf)
- Yes - Metronidazole (Flagyl) or Ciprofloxacin (Cipro) (for pouchitis only)
- No

19. Does the patient have a contraindication or intolerance to at least one conventional therapy option (e.g., azathioprine [Azasan, Imuran], corticosteroid [e.g., budesonide [Entocort, Uceris], hydrocortisone [Cortifoam, Colocort, Solu-Cortef, Cortef], methylprednisolone [Medrol, Solu-Medrol], prednisone], cyclosporine [Sandimmune], mesalamine [Asacol, Lialda, Pentasa, Canasa, Rowasa], mercaptopurine [Purinethol], sulfasalazine, tacrolimus [Prograf], metronidazole/ciprofloxacin [for pouchitis only])? *If Yes, no further questions* Yes No
20. Has the patient had an inadequate response to a TNF-alpha inhibitor indicated for the treatment of UC (e.g., Humira, Remicade, Simponi)? *If Yes, no further questions* Yes No
21. Does the patient have a contraindication or intolerance to a TNF-alpha inhibitor indicated for the treatment of UC (e.g., Humira, Remicade, Simponi)? Yes No

Maryland State Step Therapy

1. Is the requested drug being used to treat stage four advanced metastatic cancer?
 Yes No *If No, skip to #3*
2. Is the requested drug's use consistent with the FDA-approved indication or the National Comprehensive Cancer Network Drugs & Biologics Compendium indication for the treatment of stage four advanced metastatic cancer and is supported by peer-reviewed medical literature? Yes No
3. Is the requested drug being used for an FDA-approved indication OR an indication supported in the compendia of current literature (examples: AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)? Yes No
4. Does the prescribed quantity fall within the manufacturer's published dosing guidelines or within dosing guidelines found in the compendia of current literature (examples: package insert, AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)? Yes No
5. Do patient chart notes document the requested drug was ordered with a paid claim at the pharmacy, the pharmacy filled the prescription and delivered to the patient or other documentation that the requested drug was prescribed for the patient in the last 180 days? Yes No
6. Has the prescriber provided proof documented in the patient chart notes that in their opinion the requested drug is effective for the patient's condition? Yes No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)