



## Tysabri

### Prior Authorization Request

CVS Caremark administers the medical drug prior authorization program on behalf of Harvard Pilgrim Health Care. Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. **Please complete the information requested on the form below and fax this form to CVS Caremark, toll-free at 1-844-851-0882** to initiate the review process. If you have questions regarding the prior authorization please contact CVS Caremark at **1-844-387-1435**.

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**Patient's Name:** \_\_\_\_\_ **Date:** \_\_\_\_\_  
**Patient's ID:** \_\_\_\_\_ **Patient's Date of Birth:** \_\_\_\_\_  
**Physician's Name:** \_\_\_\_\_ **NPI#:** \_\_\_\_\_  
**Specialty:** \_\_\_\_\_ **HPHC Provider ID#:** \_\_\_\_\_  
**Physician Office Telephone:** \_\_\_\_\_ **Physician Office Fax:** \_\_\_\_\_

**Rendering Provider Info:**  Same as Requesting Provider

**Name:** \_\_\_\_\_ **NPI#:** \_\_\_\_\_  
**Fax:** \_\_\_\_\_ **Phone:** \_\_\_\_\_ **HPHC Provider ID:** \_\_\_\_\_

*Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.*

**Additional Demographic Information:**

Please indicate the place of service for the requested drug:

- Ambulatory Surgical  Home  Inpatient Hospital  Off Campus Outpatient Hospital  
 On Campus Outpatient Hospital  Office  Pharmacy

**Drug Information:**

Strength/Measure \_\_\_\_\_ Units  ml  Gm  mg  ea  Un  
Directions(sig) \_\_\_\_\_ Route of administration \_\_\_\_\_  
Dosing frequency \_\_\_\_\_

**Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-844-851-0882**

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**Criteria Questions:**

1. What is the diagnosis?  
 Crohn's disease  
 Relapsing form of multiple sclerosis  
 Primary progressive multiple sclerosis (PPMS)  
 Other \_\_\_\_\_
2. What is the ICD-10 code? \_\_\_\_\_

***Complete the following section based on the patient's diagnosis, if applicable.***

**Section A: Crohn's Disease**

3. Has the patient been diagnosed with moderately to severely active Crohn's disease (CD)?  Yes  No
4. Has the patient received Tysabri or any other biologic indicated for the treatment of Crohn's disease in a paid claim through a pharmacy or medical benefit in the previous 120 days?  
 Yes  No *If No, skip to #9*
5. Has the patient received any of the following medications? *If yes, please specify the most recent medication.*  
 Cimzia                       Humira                       Inflectra                       Remicade *Skip to #7*  
 Renflexis                       Entyvio                       Stelara  
 No *Skip to #9*
6. Has the patient previously received treatment with Remicade for CD?  Yes  No *If No, skip to #8*
7. Has the patient experienced any of the following during treatment with Remicade? **ACTION REQUIRED: *If yes, please provide supporting documentation***  
 Yes – Inadequate response *Skip to #16*  
 Yes – Intolerable adverse event (e.g. hypersensitivity reaction) *Skip to #16*  
 No
8. Does the patient have a contraindication to Remicade? **ACTION REQUIRED: *If yes, please provide supporting documentation***  Yes  No *If Yes or No, skip to #16*
9. Has the patient previously received treatment with Remicade for CD?  Yes  No *If No, skip to #11*
10. Has the patient experienced any of the following during treatment with Remicade? **ACTION REQUIRED: *If yes, please provide supporting documentation***  
 Yes – Inadequate response *Skip to #12*  
 Yes – Intolerable adverse event (e.g. hypersensitivity reaction) *Skip to #12*  
 No
11. Does the patient have a contraindication to Remicade? **ACTION REQUIRED: *If yes, please provide supporting documentation***  Yes  No
12. Has the patient tried and had an inadequate response to at least **ONE** conventional therapy options (e.g., azathioprine [Azasan, Imuran], budesonide [Entocort EC], ciprofloxacin [Cipro], mesalamine [Asacol, Delzicol, Pentasa, Lialda, Rowasa], Balsalazide [Colazal], olsalazine [Dipentum], mercaptopurine [Purinethol], methylprednisolone [Solu-Medrol], methotrexate, metronidazole [Flagyl], prednisone, prednisolone, sulfasalazine [Azulfidine, Sulfazine], rifaximin [Xifaxan])?  Yes  No *If No, skip to #14*
13. Please indicate the previous treatment regimen and *skip to #16.*  
  
\_\_\_\_\_

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14. Does the patient have a contraindication or intolerance to at least **ONE** conventional therapy options (e.g., azathioprine [Azasan, Imuran], budesonide [Entocort EC], ciprofloxacin [Cipro], mesalamine [Asacol, Delzicol, Pentasa, Lialda, Rowasa], Balsalazide [Colazal], olsalazine [Dipentum], mercaptopurine [Purinethol], methylprednisolone [Solu-Medrol], methotrexate, metronidazole [Flagyl], prednisone, prednisolone, sulfasalazine [Azulfidine, Sulfazine], rifaximin [Xifaxan])?  Yes  No

15. Please indicate the contraindication or intolerance.

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16. Is this a request for a continuation of therapy with Tysabri?  Yes  No *If No, no further questions*

17. How long has the patient been receiving the requested medication? \_\_\_\_\_ months

18. Has the patient achieved or maintained positive clinical response to treatment as evidenced by low disease activity or improvement in signs and symptoms?  Yes  No *No further questions*

Section B: Multiple Sclerosis

19. Has the patient been diagnosed with relapsing form of multiple sclerosis (MS)?  Yes  No

20. Has the patient tried and had an inadequate response, intolerance or contraindication to at least one alternative medication indicated for the treatment of multiple sclerosis (e.g., interferon beta-1b [Betaseron], glatiramer acetate injection [Copaxone], interferon beta-1a [Rebif], peginterferon beta-1a [Plegridy], teriflunomide [Aubagio], fingolimod [Gilenya], dimethyl fumarate [Tecfidera])?  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)

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