



HEALTH PARTNERS PLANS
PRIOR AUTHORIZATION REQUEST FORM

Health Partners Plans

Tysabri

Phone: 215-991-4300

Fax back to: 866-240-3712

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL DELAY the review process.

Form with fields: Patient Name, Prescriber Name, HPP Member Number, Date of Birth, Patient Primary Phone, Address, City, State ZIP, Line of Business, Drug Name, Quantity, Directions, Diagnosis Code, Diagnosis, etc.

HPP's maximum approval time is 12 months but may be less depending on the drug.

Please attach any pertinent medical history including labs and information for this member that may support approval.

Please answer the following questions and sign.

Q1. Does the patient have the diagnosis of multiple sclerosis?

Yes No

Q2. Is the patient 18 years of age or older?

Yes No

Q3. Is the patient receiving chronic immunosuppressant or immunomodulatory therapy?

Yes No

Q4. Did the patient have a magnetic resonance imaging (MRI) scan prior to initiating Tysabri therapy to help differentiate multiple sclerosis (MS) symptoms from progressive multifocal leukoencephalopathy (PML)?

Yes No

Q5. Did the patient have baseline testing for anti-JC virus antibodies?

Yes No

Q6. Was the baseline test for anti-JC positive?

Yes No

Q7. Was the testing for anti-JC virus antibodies repeated?

Yes No

Q8. Is this a request for a continuation of therapy?

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Patient Name: Prescriber Name:

Yes No

Q9. Has the patient's disease course improved or stabilized after starting Tysabri therapy?

Yes No

Q10. Does the patient have the diagnosis of moderately-to-severely active Crohn's disease with inflammation?

Yes No

Q11. Is the patient 18 years of age or older?

Yes No

Q12. Is the patient receiving chronic immunosuppressant or immunomodulatory therapy?

Yes No

Q13. Does the patient have a documented history of therapeutic failure of a trial to one of the following aminosaliclates: a) mesalamine for 3 months or b) sulfasalazine for 3 months?

Yes No

Q14. Does the patient have a contraindication or intolerance to one of the following aminosaliclates: A) mesalamine, B) sulfasalazine?

Yes No

Q15. Does the patient have a documented history of therapeutic failure of a trial to one the following immunomodulators: A) azathioprine for 3 months, B) methotrexate for 3 months, or C) 6-mercaptopurine for 3 months?

Yes No

Q16. Does the patient have a contraindication or intolerance to one of the following immunomodulators: A) azathioprine, B) methotrexate, OR C) 6-mercaptopurine?

Yes No

Q17. Does the patient have a documented history of therapeutic failure of a trial to one of the following TNF inhibitors: A) Cimzia for 6 weeks, B) Humira for 3 months, OR C) Remicade for 10 weeks?

Yes No

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Patient Name: Prescriber Name:

Q18. Does the patient have a contraindication or intolerance to one of the following TNF inhibitors: A) Cimzia, B) Humira, OR c) Remicade? Q19. Did the patient have baseline testing for anti-JC virus antibodies? Q20. Is this a request for a continuation of therapy? Q21. Is the patient taking chronic oral corticosteroids prior to starting Tysabri? Q22. Does the patient require additional steroid use that exceeds 3 months in a calendar year to control their Crohn's disease? Q23. Did the patient experience therapeutic benefit after 3 months of therapy? Q24. Was the patient able to discontinue concomitant corticosteroid use within 6 months of starting therapy? Q25. Was the baseline anti-JC virus antibody testing negative? Q26. Did the provider repeat testing for anti-JC antibodies? Q27. Additional Information:

Prescriber Signature

Date

Updated for 2021