

Policy Title:	Tysbari (natalizumab) (Intravenous)		
		Department:	PHA
Effective Date:	01/01/2020		
Review Date:	9/12/2018, 12/13/2019, 1/22/2020, 6/10/2021, 1/20/2022, 7/13/2023		

Purpose: To support safe, effective, and appropriate use of Tysbari (natalizumab) in the treatment of Multiple Sclerosis and Crohn's disease.

Scope: Medicaid, Commercial, Medicare-Medicaid Plan (MMP)

Policy Statement:

Tysbari (natalizumab) is covered under the medical benefit when used within the following guidelines. Use outside of these guidelines may result in non-payment unless approved under an exception process.

Procedure:

Coverage of Tysbari (natalizumab) will be reviewed prospectively via the prior authorization process based on criteria below.

Initial Criteria:

- Patient is at least 18 years of age; AND
- Patient has had anti-JCV antibody testing with ELISA prior to initiating treatment with natalizumab and annually thereafter; AND
- Cannot be used in combination with antineoplastic, immunosuppressant, or immunomodulating agents; AND
- Patient must not have a systemic medical condition resulting in significantly compromised immune system function
- MMP members who have previously received this medication within the past 365 days are not subject to Step Therapy Requirements

Multiple Sclerosis (MS):

- Patient is diagnosed with a relapsing form of multiple sclerosis and documented by laboratory report (i.e., MRI); AND
- Member has had a failure, intolerance, or contraindication to at least two drugs indicated for the treatment of relapsing MS; AND
- Must be prescribed by or in consultation with a neurologist; AND
- Used as monotherapy for the treatment of relapsing forms of MS; AND

Crohn's Disease:

- Patient is diagnosed with Crohn's disease by a gastroenterologist; AND
- Prescriber has assessed baseline disease severity utilizing an objective measure/tool; AND
- Patient has had failure, intolerance, or contraindication to at least two oral immunosuppressive therapy for at least 3 months, such as corticosteroids, methotrexate, azathioprine, and/or 6-mercaptopurine; AND
- Patient has had failure, intolerance, or contraindication to at least one TNF-Inhibitor therapy for at least 3 months; AND
- Patient is not taking in combination with another biologic drug or immunosuppressant (e.g., 6-mercaptopurine, azathioprine, cyclosporine, methotrexate, etc.) used for Crohn's disease; AND
- Tysabri is prescribed by or in consultation with a gastroenterologist

Continuation of therapy criteria:

- Multiple Sclerosis:
 - Patient is tolerating treatment
 - Patient has annual anti-JCV antibody testing with ELISA
 - Patient has experienced disease improvement or slowing of disease worsening (e.g., no decline in Expanded Disability Status Score [EDSS] or MRI findings) since initiating therapy
- Crohn's disease:
 - Patient is tolerating treatment
 - Patient has annual anti-JCV antibody testing with ELISA
 - For initial renewal only:
 - Patient must show clinical response and remission of disease by 12 weeks
 - For all subsequent renewals:
 - Patient does not require additional steroid use that exceeds three months in a calendar year
 - Disease response as indicated by improvement in signs and symptoms compared to baseline such as endoscopic activity, number of liquid stools, presence and severity of abdominal pain, presence of abdominal mass, body weight compared to IBW, hematocrit, presence of extra intestinal complications, use of anti-diarrheal drugs, and/or an improvement on a disease activity scoring tool [e.g., an improvement on the Crohn's Disease Activity Index (CDAI) score or the Harvey-Bradshaw Index score]

Coverage durations:

- Initial coverage criteria = 3 months for Crohn's disease and 6 months for MS
- Continuation of therapy = 12 months

*** Requests will also be reviewed to National Coverage Determination (NCD) and Local Coverage Determinations (LCDs) if applicable. ***

Dosage/Administration:

Indication	Dose	Maximum dose (1 billable unit = 1 mg)
All Indications	300 mg intravenously over one hour every four weeks	300 billable units every 28 days

Investigational use: All therapies are considered investigational when used at a dose or for a condition other than those that are recognized as medically accepted indications as defined in any one of the following standard reference compendia: American Hospital Formulary Service Drug information (AHFS-DI), Thomson Micromedex DrugDex, Clinical Pharmacology, Wolters Kluwer Lexi-Drugs, or Peer-reviewed published medical literature indicating that sufficient evidence exists to support use. Neighborhood does not provide coverage for drugs when used for investigational purposes.

Applicable Codes:

Below is a list of billing codes applicable to covered treatment options for multiple sclerosis. The below tables are provided for reference purposes and may not be all-inclusive. Requests received with codes from tables below do not guarantee coverage. Requests must meet all criteria are provided in the procedure section.

Codes:

The following HCPCS/CPT codes are:

HCPCS/CPT Code	Description
J2323	Injection, natalizumab, 1mg

References:

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2. Fox RJ, Cutter G, Chan A, et al. Comparative Effectiveness Using A Matching-Adjusted Indirect Comparison Between Delayed-Release Dimethyl Fumarate and Fingolimod for The Treatment of Relapsing-Remitting Multiple Sclerosis. *Value Health.* 2015 Nov;18(7):A750. Epub 2015 Oct 20. [PubMed](#)
3. Metin H, Huppertz H. Adjusted Indirect Comparison of Oral Multiple Sclerosis Agents. *Value Health.* 2015 Nov;18(7):A750. Epub 2015 Oct 20. [PubMed](#)
4. Tramacere I, Del Giovane C, Salanti G, et al. Immunomodulators and immunosuppressants for relapsing-remitting multiple sclerosis: a network meta-analysis. *Cochrane Database Syst Rev.* 2015. [PubMed](#)
5. Tolley K, Hutchinson M, You X, et al. A Network Meta-Analysis of Efficacy and Evaluation of Safety of Subcutaneous Pegylated Interferon Beta-1a versus Other Injectable Therapies for the Treatment of Relapsing-Remitting Multiple Sclerosis. *PLoS One.* 2015;10(6):e0127960.
6. Bainbridge JL, Miravalle A, Corboy JR. Multiple Sclerosis. In: DiPiro JT, Talbert RL, Yee GC, Matzke GR, Wells BG, Posey L, eds. *Pharmacotherapy: A Pathophysiologic Approach.* 9th ed. New York, NY: McGraw-Hill; 2014. <http://accesspharmacy.mhmedical.com/content.aspx?bookid=689&Sectionid=45310489>. Accessed May 18, 2016.
7. Goodin DS, Frohman EM, Garmany GP, et al. Disease modifying therapies in multiple sclerosis: report of the Therapeutics and Technology Assessment Subcommittee of the American Academy of Neurology and the MS Council for Clinical Practice Guidelines. *Neurology.* 2002;58(2):169-178.
8. Hauser SL, Bar-Or A, Comi G, Giovannoni G, Hartung HP, Hemmer B, Lublin F, Montalban X, Rammohan KW, Selmaj K, et al. Ocrevus versus Interferon Beta-1a in Relapsing Multiple Sclerosis. *N Eng J Med.* 2016;376(3):221–234. doi: 10.1056/NEJMoa1601277.
9. Montalban X, et al. Ocrevus versus Placebo in Primary Progressive Multiple Sclerosis. *N Engl J Med.* 2017;376:209–220. doi: 10.1056/NEJMoa1606468
10. Tysabri prescribing information. South San Francisco, CA: Elan Pharmaceuticals, Inc.; June 2020.