## TYSABRI (NATALIZUMAB) [PREFERRED] PRIOR AUTHORIZATION FORM







(form effective 1/6/2025)

Prescriber signature:

Fax to PerformRx $^{\text{SM}}$  at **1-855-851-4058**, or to speak to a representative call **1-888-674-8720**.

PRIOR AUTHORIZATION REQUEST	NEORMATION							
□ New request □ Renewal request Total # pages: Name of office contact:								
Contact's phone number:	P.O.	LTC facility	cility contact/phone:					
PATIENT INFORMATION								
Patient name:			Patient ID #:			DOB:		
Street address: Apt.								
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PRESCRIBER INFORMATION Prescriber name:			Specialty:					
State license #:	NPI:		MA Provider ID #					
Street address:	IVI I.	Suite	e #·	City/state				
Phone:		Cuit	Fax:	Oity/otate	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			
			Tun					
CLINICAL INFORMATION  Medication requested: Tysabri (natalizumab) 300 m	ng/15 ml			Quantity:	vial	Refills:		
Directions: ☐ 300 mg SQ every 4 weeks ☐ other				Quality.	Viai	Dx code (required):		
Diagnosis:     relapsing multiple sclerosis – Submit documentation of diagnosis and disease pattern.								
☐ moderately to severely active Crohn's disease with inflammation — Submit documentation of diagnosis and disease severity.								
□ other: − Submit documentation supporting the use of Tysabri for the patient's condition.								
PHARMACY INFORMATION (Prescrib	er to identify the ph	narmacy	that is to d	ispense	the medication,	f applicable):		
Deliver to: ☐ Patient's Home ☐ Physician's Office	□ Patient's Preferred Ph	narmacy Nar	ne:					
				Pharmacy Fax #:				
☐ I acknowledge that the patient agrees with the pharmacy chosen for delivery of this medication.								
HCPCS (HEALTHCARE COMMON PROCEDURE CODING SYSTEM) INFORMATION (if applicable):								
Treatment setting:   Infusion Center   Home   Provider's Office   Hospital Outpatient Facility								
Facility name:			Facility NPI:					
J-code:			Number of ur	iits:		Date of service (MM/DD/YYYY):		
INITIAL REQUESTS								
1. Is Tysabri (natalizumab) being prescribed by or in consultation with an appropriate specialist?  ☐ Yes list specialty:								
☐ Yes, list specialty:								
2. Is patient receiving chronic immunosuppressant or immunomodulator therapy?								
☐ Yes, list medications:								
3. For the treatment of Crohn's disease, does at least one of the following apply to the patient?								
☐ moderate to severe Crohn's disease and one of the following:								
☐ failed to achieve remission with or has a contraindication or intolerance to an induction course of corticosteroids ☐ failed to maintain remission or has a contraindication or intolerance to immunomodulators								
☐ has one or more high-risk or poor prognostic features								
□ has achieved remission with the requested medication and will be using the requested medication as maintenance therapy to maintain remission								
4. For the treatment of Crohn's disease, select all that apply to the patient.  □ history of trial and failure of at least one tumor necrosis factor (TNF) inhibitor OR contraindication or intolerance to TNF inhibitors;								
list medications tried OR provide explanation for contraindication/intolerance:								
☐ history of therapeutic failure, contraindication, or intolerance to ustekinumab (Stelara) ☐ history of therapeutic failure, contraindication, or intolerance to vedolizumab (Entyvio)								
☐ current history (within the past 90 days) of being prescribed Tysabri								
RENEWAL REQUESTS								
1. Is Tysabri (natalizumab) being prescribed by or in	consultation with an appropri	iate speciali	st? 🗆 Yes, list	specialty:_				
2. For the treatment of multiple sclerosis, did the patient experience disease improvement or stabilization since starting Tysabri?   No								
	atient experience disease im,	provement		J				
Submit documentation of response to therapy.		provement						
Submit documentation of response to therapy.  3. For the treatment of Crohn's disease, select all the experienced therapeutic benefit within 3 months.	nat apply to the patient. ns of starting therapy							
Submit documentation of response to therapy.  3. For the treatment of Crohn's disease, select all the experienced therapeutic benefit within 3 monther was able to discontinue concomitant corticoste	nat apply to the patient. ns of starting therapy roid use within 6 months of s	starting ther						
Submit documentation of response to therapy.  3. For the treatment of Crohn's disease, select all the experienced therapeutic benefit within 3 months.	nat apply to the patient. s of starting therapy roid use within 6 months of s than 3 months in a calendar	starting ther	ару					

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Date: