

TY SABRI (natalizumab) PRIOR AUTHORIZATION FORM *(form effective 01/03/2022)*

Prior authorization guidelines for **Tysabri** and **Quantity Limits/Daily Dose Limits** are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	total pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	MA Provider ID#:
LTC facility contact/phone:			Street address:	
Beneficiary Name:			Suite #:	City/state/zip:
Beneficiary ID#:	DOB:	Phone:	Fax:	

CLINICAL INFORMATION

Drug requested:	<input type="checkbox"/> Tysabri (natalizumab) 300 mg/15 ml	Quantity: _____ vials	Refills:
Directions:	<input type="checkbox"/> 300 mg SQ every 4 weeks	<input type="checkbox"/> other: _____	
Diagnosis (<i>submit documentation</i>):			Dx code (<i>required</i>):
Is the beneficiary currently being treated with Tysabri?		<input type="checkbox"/> Yes – date of last dose: _____ <i>Submit documentation.</i> <input type="checkbox"/> No	
Is Tysabri being prescribed by or in consultation with a neurologist or gastroenterologist?		<input type="checkbox"/> Yes <i>Submit documentation of consultation if applicable.</i> <input type="checkbox"/> No	
Is the beneficiary receiving chronic immunosuppressive or immune modulating therapies?		<input type="checkbox"/> Yes <i>Submit complete medication list.</i> <input type="checkbox"/> No	

INITIAL requests

Check all of the following that apply to the beneficiary and this request and SUBMIT DOCUMENTATION for each item.

- Is being treated for multiple sclerosis
 - Has a relapsing form of MS
- Is being treated for Crohn's disease
 - Has moderate-to-severe disease
 - Has disease that is associated with high-risk or poor prognostic features
 - Failed to achieve remission with an induction course of corticosteroids
 - Has a contraindication or intolerance to an induction course of corticosteroids
 - Failed to maintain remission with an immunomodulator (e.g., AZA, 6-MP, MTX)
 - Has a contraindication or intolerance to immunomodulators (e.g., AZA, 6-MP, MTX)
 - Tried and failed or has a contraindication or intolerance to a TNF-inhibitor (e.g., Cimzia, Humira, Remicade)
 - Tried and failed or has a contraindication or intolerance to ustekinumab (Stelara)
 - Tried and failed or has a contraindication or intolerance to vedolizumab (Entyvio)

RENEWAL requests

Check all of the following that apply to the beneficiary and this request and SUBMIT DOCUMENTATION for each item.

- For a diagnosis of multiple sclerosis, experienced improvement or stabilization of the MS disease course since starting Tysabri
- For a diagnosis of Crohn's disease:
 - Experienced therapeutic benefit within 3 months of starting Tysabri
 - Was able to discontinue concomitant steroid use within 6 months of starting Tysabri (if applicable)
 - Has not required concomitant steroid use for disease control for more than 3 months in the past 12 months (if >1 year since starting Tysabri)

Please submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

GHP Family Pharmacy Customer Service
100 N. Academy Ave.
Danville, PA 17822
Tel. • 855•552•6028 PA Relay 711 GeisingerHealthPlan.com



Prescriber Signature:	Date:
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