

### **TYSABRI (natalizumab) PRIOR AUTHORIZATION FORM**

Prior authorization guidelines for **Tysabri** and **Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at <https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>.

<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 ( <u>submit documentation</u> ):			Dx code ( <u>required</u> ):
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 FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION**

<b>Prescriber Signature:</b>	<b>Date:</b>
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