

MULTIPLE SCLEROSIS AGENTS PRIOR AUTHORIZATION FORM (form effective 1/8/2024)

Prior authorization guidelines for **Multiple Sclerosis Agents** 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	# of pages: _____	Prescriber name:	
Name of office contact:		Specialty:	
Contact's phone number:		NPI:	State license #:
LTC facility contact/phone:		Street address:	
Beneficiary name:		City/state/zip:	
Beneficiary ID#:	DOB:	Phone:	Fax:

CLINICAL INFORMATION

Drug requested:	Dosage form:	Strength:	
Directions:		Quantity:	Refills:
Diagnosis (<i>submit documentation</i>):	Dx code (<i>required</i>):	Beneficiary's weight:	
Is the beneficiary currently being treated with the requested medication?	<input type="checkbox"/> Yes – date of last dose: _____ <i>Submit documentation.</i> <input type="checkbox"/> No		
Is the requested medication being prescribed by or in consultation with a neurologist (or, for Ampyra/dalfampridine, a neurologist or physical medicine and rehabilitation (PM&R) specialist)?	<input type="checkbox"/> Yes <i>Submit documentation of consultation if applicable.</i> <input type="checkbox"/> No		

Complete all sections that apply to the beneficiary and this request.

Check all that apply and submit documentation for each item.

INITIAL requests

<input type="checkbox"/> Has a relapsing form of MS (<i>specify</i>) → <input type="checkbox"/> clinically isolated syndrome <input type="checkbox"/> relapsing remitting disease <input type="checkbox"/> active secondary progressive disease
<input type="checkbox"/> Has primary progressive MS
<input type="checkbox"/> Request is for a NON-PREFERRED Multiple Sclerosis Agent: <input type="checkbox"/> Tried and failed or has a contraindication or an intolerance to the preferred drugs in this class approved for the beneficiary's diagnosis (<i>Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.</i>)
<input type="checkbox"/> Request is for AMPYRA (dalfampridine): <input type="checkbox"/> Has motor dysfunction on a continuous basis that impairs the ability to complete activities of daily living (ADLs) or instrumental ADLs <input type="checkbox"/> Has results of recent kidney function tests <input type="checkbox"/> Has a history of seizure
<input type="checkbox"/> Request is for AUBAGIO (teriflunomide): <input type="checkbox"/> Has results of recent liver function tests
<input type="checkbox"/> Request is for GILENYA (fingolimod):

☐ Has a comorbid heart condition – describe: _____

☐ Experienced any of the following in the past 6 months:

☐ Myocardial infarction

☐ Transient ischemic attack

☐ Unstable angina

☐ Decompensated heart failure requiring hospitalization

☐ Stroke

☐ Class III or IV heart failure

☐ **Request is for KESIMPTA (ofatumumab):**

☐ Does not have active hepatitis B virus infection

☐ **Request is for LEMTRADA (alemtuzumab):** Dates of previous treatment course(s): _____

☐ **Request is for MAVENCLAD (cladribine):** Dates of previous treatment course(s): _____

☐ Has results of a recent lymphocyte count AND:

☐ Lymphocyte count is within normal limits prior to initiating first treatment course

☐ **Request is for MAYZENT (siponimod):**

☐ Has been tested for CYP2C9 variants to determine CYP2C9 genotype

☐ Has a comorbid heart condition – describe: _____

☐ Experienced any of the following in the past 6 months:

☐ Myocardial infarction

☐ Transient ischemic attack

☐ Unstable angina

☐ Decompensated heart failure requiring hospitalization

☐ Stroke

☐ Class III or IV heart failure

☐ **Request is for OCREVUS (ocrelizumab):**

☐ Does not have active hepatitis B virus infection

☐ **Request is for ZEPOSIA (ozanimod):**

☐ Has severe untreated sleep apnea

☐ Will be taking a monoamine oxidase (MAO) inhibitor while taking Zeposia (e.g., selegiline, phenelzine)

☐ Has a comorbid heart condition – describe: _____

☐ Experienced any of the following in the past 6 months:

☐ Myocardial infarction

☐ Transient ischemic attack

☐ Unstable angina

☐ Decompensated heart failure requiring hospitalization

☐ Stroke

☐ Class III or IV heart failure

RENEWAL requests

☐ **For AMPYRA (dalfampridine):**

☐ Experienced an improvement in motor function since starting the requested medication

☐ Has a history of seizure

☐ **For all MS drugs OTHER THAN Ampyra (dalfampridine):**

☐ Has a relapsing form of MS AND:

☐ Experienced improvement or stabilization of the MS disease course since starting the requested medication

☐ Has primary progressive MS AND:

☐ Continues to benefit from the requested medication

☐ **Request is for AUBAGIO (teriflunomide):**

☐ Has results of recent liver function tests

☐ **Request is for GILENYA (fingolimod):**

☐ Has a comorbid heart condition – describe: _____

☐ Experienced any of the following in the past 6 months:

☐ Myocardial infarction

☐ Transient ischemic attack

☐ Unstable angina

☐ Decompensated heart failure requiring hospitalization

☐ Stroke

☐ Class III or IV heart failure

☐ **Request is for KESIMPTA (ofatumumab):**

<input type="checkbox"/> Does not have active hepatitis B virus infection	
<input type="checkbox"/> Request is for LEMTRADA (alemtuzumab): Dates of previous treatment course(s): _____	
<input type="checkbox"/> Request is for MAVENCLAD (cladribine): Dates of previous treatment course(s): _____	
<input type="checkbox"/> Has results of a recent lymphocyte count AND:	
<input type="checkbox"/> Lymphocyte count is at least 800 cells/microliter before initiating second treatment course	
<input type="checkbox"/> Request is for MAYZENT (siponimod):	
<input type="checkbox"/> Has a comorbid heart condition – describe: _____	
<input type="checkbox"/> Experienced any of the following in the past 6 months:	
<input type="checkbox"/> Myocardial infarction	<input type="checkbox"/> Transient ischemic attack
<input type="checkbox"/> Unstable angina	<input type="checkbox"/> Decompensated heart failure requiring hospitalization
<input type="checkbox"/> Stroke	<input type="checkbox"/> Class III or IV heart failure
<input type="checkbox"/> Request is for OCREVUS (ocrelizumab):	
<input type="checkbox"/> Does not have active hepatitis B virus infection	
<input type="checkbox"/> Request is for ZEPOSIA (ozanimod):	
<input type="checkbox"/> Has severe untreated sleep apnea	
<input type="checkbox"/> Will be taking a monoamine oxidase (MAO) inhibitor while taking Zeposia (e.g., selegiline, phenelzine)	
<input type="checkbox"/> Has a comorbid heart condition – describe: _____	
<input type="checkbox"/> Experienced any of the following in the past 6 months:	
<input type="checkbox"/> Myocardial infarction	<input type="checkbox"/> Transient ischemic attack
<input type="checkbox"/> Unstable angina	<input type="checkbox"/> Decompensated heart failure requiring hospitalization
<input type="checkbox"/> Stroke	<input type="checkbox"/> Class III or IV heart failure
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.	
Prescriber Signature:	Date:

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