

GILENYA Prior Authorization Form

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- Standard Request (72 hours)
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- Expedited Request (24 hours)

If you or your prescriber believe that waiting 72 hours for a standard decision could seriously harm your life, health, or ability to regain maximum function, you can request an expedited decision. For expedited requests you will receive a decision within 24 hours. You cannot request an expedited coverage determination if you are requesting reimbursement for a drug you already received.

Demographics

Patient Information		Prescriber Information	
Patient Name:		Prescriber Name:	
DOB:	Age:	NPI#:	Specialty:
Health Plan ID#:		Phone:	Fax:
Pharmacy Name:	Pharmacy Phone:	Office Contact:	Direct Phone # or Ext:

Medication Information

Drug Requested: Gilenya	Strength: 0.5 mg	Directions:	Quantity Dispensed:	Day Supply:
<input type="checkbox"/> New medication <input type="checkbox"/> Continuation of therapy	Start Date:	If this is continuation of therapy, please provide CHART DOCUMENTATION indicating the member showed improvement while on therapy.		

Clinical Information

Diagnosis:				Date of Diagnosis:	
Please indicate past medication(s) tried and failed					
Medication name	Start date	End date	Strength	Frequency	Reason for Failure or Discontinuation
<input type="checkbox"/> Avonex					
<input type="checkbox"/> Copaxone					
<input type="checkbox"/> Betaseron					
<input type="checkbox"/> Extavia					
<input type="checkbox"/> Rebif					
<input type="checkbox"/> Tecfidera					
Does the member have a relapsing form of Multiple Sclerosis?					<input type="checkbox"/> Yes <input type="checkbox"/> No
Will the member be observed for 6 hours for signs and symptoms of bradycardia?					<input type="checkbox"/> Yes <input type="checkbox"/> No
Does the member have evidence of active infection?					<input type="checkbox"/> Yes <input type="checkbox"/> No
Did the member have a recent (within the past 6 months) complete blood count (CBC)? If yes, please indicate date: _____					<input type="checkbox"/> Yes <input type="checkbox"/> No
Did the member have a recent (with in the past 6 months) transaminase and bilirubin level? If yes, please indicate date: _____					<input type="checkbox"/> Yes <input type="checkbox"/> No

Was the member vaccinated against varicella zoster virus (VZV)? If yes, please indicate date: _____		<input type="checkbox"/> Yes <input type="checkbox"/> No
Has the member demonstrated immunity to VZV by VZV antibody serology? If yes, please provide chart documentation of VZV antibody serology.		<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Chart Notes
Did the member have a baseline ophthalmologic evaluation of the macula? If yes, please indicate date: _____		<input type="checkbox"/> Yes <input type="checkbox"/> No
Did the member have a recent electrocardiogram (ECG)? If yes, please indicate date: _____		<input type="checkbox"/> Yes <input type="checkbox"/> No
Does the member have Mobitz Type II second- or third-degree atrioventricular (AV) block or sick sinus syndrome? If yes, does the member have a functioning pacemaker?		<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> No
Please provide the member's baseline QTc interval: _____		
Does the member have pre-existing lung disease, such as asthma or COPD? If yes, please provide a spirometric evaluation of respiratory function and evaluation of diffusion lung capacity for carbon monoxide		<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Chart Notes
Did the member recently (in the past 6 months) experience a myocardial infarction, unstable angina, stroke, TIA, decompensated heart failure requiring hospitalization, or Class III/IV heart failure?		<input type="checkbox"/> Yes <input type="checkbox"/> No
Is the member on concomitant therapy with antineoplastic, immunosuppressive therapy, or immune modulating therapies? If yes, please complete below:		<input type="checkbox"/> Yes <input type="checkbox"/> No
Medication	Dose/Strength	Frequency
Is the member on concomitant therapy with any Class I or Class III antiarrhythmic medications? If yes, please complete below:		<input type="checkbox"/> Yes <input type="checkbox"/> No
Medication	Dose/Strength	Frequency
<p>Is this request for a reauthorization? <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>If yes, please include the following documentation:</p> <ul style="list-style-type: none"> <input type="checkbox"/> Documentation showing member's disease has improved and/or stabilized <input type="checkbox"/> Documentation of no active infection <input type="checkbox"/> Documentation of 3-month follow-up ophthalmologic evaluation within 3 to 4 months of starting therapy, including date. <input type="checkbox"/> Documentation that the member CBC and transaminase/bilirubin levels are being monitored consistently. 		
Please provide any additional information which should be considered in the space below:		