

Fax completed prior authorization request form to 855-799-2551 or submit Electronic Prior Authorization through CoverMyMeds[®] or SureScripts.

Aetna Better Health®

All requested data must be provided. Incomplete forms or forms without the chart notes will be returned

Pharmacy Coverage Guidelines are available at www.aetnabetterhealth.com/michigan/providers/medicaid/pharmacy

Multiple Sclerosis Agents Pharmacy Prior Authorization Request Form

Do not copy for future use. Forms are updated frequently.

REQUIRED: Office notes, labs and medical testing relevant to request showing medical justification are required to support diagnosis Member Information

Member Name (first & last):		Date of Birth:		Gender:			Height:				
Member ID:		O'thu				Male	🗆 Fe	emale) A (= : = : = : = :		
Member ID.		City:			State:			Weight:			
Prescribing Provider Information	on										
Provider Name (first & last):	Specialty:	Specialty:			NPI#		DEA#				
Office Address:	City:			State:				Zip Code:			
Office Contact:	Office Contact:			Office Phone			Office	Office Fax:			
Dispensing Pharmacy Informat	ion						Ι				
Pharmacy Name:			Pharmacy Phone: P			Pharm	Pharmacy Fax:				
Requested Medication Informa	tion										
🗆 Aubagio®	🗆 Bafierta	am™		🗆 Cop	axone®	40 mg □ Extavia®					
□ Glatiramer 20mg/ml and 40 mg/ml	🗆 Gilenya	®		□ Glatopa®				□ Kesimpta®			
□ Mavenclad®	🗆 Mayzer	Mayzent®			Plegridy [®]			Ponvory [®]			
□ Rebif®/ Rebif Rebidose®	🗆 Tascen	so ODT®	□ Tecfidera®				□ Vumerity®				
□ Zeposia®	Zeposia® D Other, please specify:										
Medication request is NOT for an FDA approved, or compend diagnosis (circle one): Yes No				a-supported ICD-10 Code: [Diagno	Diagnosis:			
What medication(s) have been tr	ied and failed	for diagnosis? (p	olease	specify):	•						
Has the member had a therapeut	Has the member had a therapeutic failure after one-month trial with one preferred medication?					□ Yes	🗆 No				
Does the member have any of the following to the preferred medication(s): (check all that apply)				□ Con	Allergy Contraindication or drug interactions History of unacceptable side effects						
Directions for Use: Strength:						Dosage Form:					
Qua		Quantity:	Quantity: Day		y Supply:		Duration of Therapy/Use:				
Turn-Around Time for Review											
h			Irgent – If waiting 24 hours for a standard decision could seriously harm life, ealth, or ability to regain maximum function, you can ask for an expedited								
		decision.	•								

		Signature:				_			
Bafiertam									
Does the member have a diagnosis of relapsing forms of multiple sclerosis (MS) to include clinically isolated syndrome (CIS), relapsing-remitting disease (RRMS) and active secondary progressive disease (SPMS)?							Yes		No
Prescriber attests that Bafiertam will be used as single agent monotherapy						Yes		No	
RENEWAL of Bafiertam:		Attestation of tolerance to maintenance dose Attestation of a CBC, including lyn serum aminotransferase, ALP, and							vels
Kesimpta									
Does the member have a diagnosis of relapsing forms of multiple sclerosis (MS) to include clinically isolated syndrome (CIS), relapsing-remitting disease (RRMS) and active secondary progressive disease (SPMS)?							Yes		No
Prescriber attests that Kesimpta will be used as single agent monotherapy							Yes		No
Prescriber attests that the first inj	ection will be n	nonitored by a healt	hcare	professional			Yes		No
Plegridy, Mavenclad, Mayzent,	Vumerity, Zep	osia							
Has the member had a therapeut	ic failure of on	e-month trial of at le	ast tv	vo preferred medications?			Yes		No
Ponvory						<u> </u>			
Does the member have a diagnor syndrome (CIS), relapsing-remitti		• ·			•		Yes		No
 Member has obtained a baseline electrocardiogram (ECG) 	Prescrib first-dos	er attests that e monitoring, as indicated, will		Member does NOT have an active infection, including clinically important localized infections	□ Merr for a varic or ha imm	ntiboo ella z s cor uniza prior	nas bee dies to oster v npleteo tion sei to begi	the irus (d the ries fo	VZV) or
For members with a history of uveitis and/or diabetes ONLY: A baseline ophthalmic evaluation of the fundus, including the macula, before starting treatment?							Yes		No
Prescriber attests that ponesimod will NOT be used in combination with anti-neoplastic, immunosuppressive, or							Yes		No
immune-modulating therapies, or, if therapy is unavoidable, the member will be monitored closely for adverse reactions and/or dose modifications?									
Has the member had a therapeutic failure of one-month trial of at least two preferred medications?							Yes		No
Mavenclad									
Does the member have a diagnosis of relapsing forms of multiple sclerosis (MS) to include relapsing-remitting disease and active secondary progressive disease?						Yes		No	
Mayzent						<u> </u>			
Does the member have a diagno							Yes		No
Member CYP2C9 variant status has been tested to determine genotyping (required for dosing)	ed to baseline for antibodies to the uveit ng electrocardiogram (ECG) varicella zoster virus (VZV) ONLY or has completed the ophth immunization series for the fu VZV prior to beginning macu					bers with a history of is and/or diabetes Y; a baseline halmic evaluation of undus, including the ula, before starting ment			
Tascenso ODT									
Does the member have a diagnos							Yes		No
syndrome (CIS), relapsing-remitting disease (RRMS) or active secondary progressive disease (SPMS)? Is the member unable to use generic fingolimod capsules or brand Gilenya capsules due to swallowing difficulties?						Yes		No	
Vumerity									
Does the member have a diagnosis of relapsing forms of multiple sclerosis (MS) to include clinically isolated syndrome (CIS), relapsing-remitting disease (RRMS) and active secondary progressive disease (SPMS)?						Yes		No	
Zeposia									
Does the member have a diagnosis of moderately or severely active ulcerative colitis (UC) and is prescribed by or in consultation with a gastroenterologist?						Yes		No	

-		e sclerosis (MS) to include clinical	-	□ Yes	🗆 No			
		ondary progressive disease (SPMS	Í					
Member has obtained a	Member does NOT have	Member has been tested		bers with a				
baseline	an active infection,	for antibodies to the	uveiti	s and/or di	abetes			
electrocardiogram (ECG)	including clinically	varicella zoster virus (VZV)		; a baseline				
	important localized	important localized or has completed the opt						
	infections	immunization series for	the fu	uding the				
		VZV prior to beginning	macu	starting				
		therapy	treatr					
Prescriber attests that a CBC wi	ith lymphocyte count, ALT, AST, a	nd total bilirubin have been obtain	tained for the D Yes D					
member in the past 6 months?								
For MS, has the member had a th	nerapeutic failure of one-month tria	al of at least two preferred medicati	ons?	□ Yes	🗆 No			
,					_			
Additional information the pres	cribing provider feels is importan	nt to this review. Please specify b	elow or sub	omit medic	al records			
	51							
Signature affirms that information given on this form is true and accurate and reflects office notes.								
Prescribing Provider's Signatu	re:	Date:						
								

Please note: Incomplete forms or forms without the chart notes will be returned

Office notes, labs, and medical testing relevant to the request that show medical justification are required. Standard turnaround time is 24 hours. You can call 855-300-5528 to check the status of a request.