

Preauthorization Request Form

[Multiple Sclerosis (MS) therapies only]

Please complete each section of this form. Incomplete forms may be returned to sender for additional information. NOTE: For your patient to receive the lowest out-of-pocket costs, use in-network providers unless preauthorization is obtained. Decisions are based on eligibility, benefit determination and medical necessity.

Member name:		Date of Birth:	
Member ID Number:		Group Number:	
NPI number of administering facility:		Requested start date:	
Drug (pharmacy benefit preferred drugs listed	d)		
Oral drugs Aubagio (teriflunomide) Mavenclad (cladribine) Bafiertam (monomethyl fumarate)	 Gilenya (fingolimod) Dimethyl fumarate 	Mayzent (simponimod)Zeposia (ozanimod)	
Self-injectable drugs Avonex (interferon beta-1a) Extavia (interferon beta-1b) Plegridy (peginterferon beta-1a) Other:		 Copaxone (glatiramer) Glatopa (glatiramer) Kesimpta (ofatumumab) 	
Dose & Schedule Requested:			
HCPCS code(s), please list all that apply (if applicable):			
Drug (medical benefit drugs listed)			
 Lemtrada (alemtuzumab) Other: 	Ocrevus (ocrelizumab)	🔲 Tysabri (natalizumab)	
Dose & Schedule Requested:			
HCPCS code(s), please list all that apply (if applicable):			
Indication			
ICD code(s), please list all that apply:			
 Multiple Sclerosis Relapse Remitting Primary Progressive Secondary Progressive Clinically Isolated Syndrome Other: 			
Clinical information			
Initial therapy Continuation of therapy (see	e below)		
For initial therapy, please complete questions below.			
1. Is the member greater than or equal to 18 years of	of age? Yes 🗌 No 🗌		
2. Prior to initiating therapy, has patient received a b	oaseline MRI? Yes 🗌 No 🗌		

3.	Will the medication be used as monotherapy and avoid combination with any other disease modifying therapy (DMT) medication? Yes 🗌 No 🗌		
	If "No" please provide clinical rationale as to why multiple DMT therapies are clinically necessary:		
4.	Is the medication prescribed by, or in consultation with, a neurologist? Yes 🗌 No 🗌		
5.	Does the member have a documented contraindication (per individual product FDA label) that would result in member not being eligible for the requested medication? Yes 🗌 No 🗌		
For	Drug-specific clinical contraindications Aubagio (teriflunomide): severe hepatic impairment Gilenya (fingolimod): Class III or IV heart failure Concomitant Class la or Class III anti-arrhythmic drugs Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Myocardial infarction, decompensated heart failure requiring hospitalization, stroke, TIA, or unstable angina within the last 6 months QTc interval at baseline 500 milliseconds or greater Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Mayzent (simponimod): Class III or IV heart failure Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Second-degree or third-degree atrioventricular block (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the patient has a functional pacemaker Sick-sinus syndrome (history or current), unless the p		
NOTE: Clinical effectiveness established based on the use of drug samples that bypasses policy requirements is not considered a prerequisite for continued coverage. Additionally, prior coverage of a drug under a previous insurance carrier is not a prerequisite for continued coverage under the Health Plans.			
Pro	vider Name: Office/Facility Name:		
Per	son completing the form: Form Completion Date:		
Per	son reviewing the form: Form Review Date:		
Phone Number: () Fax Number: ()			
Det	ermination of medical necessity requires the submission of clinical documentation.		
	Clinical documentation is available in the Avera electronic medical record for review. Please list date(s) of pertinent records:		
D pre	Clinical documentation is not available in the Avera electronic medical record for review. Pertinent clinical records for the vious 12 months are attached for review.		

Final determination will be faxed to the prescriber. Final determination will be mailed to the member.

IMPORTANT NOTICE: This determination does not guarantee benefits or payment of services. Payment of services is subject to patient eligibility at the time of treatment, benefit plan limitations and the other terms of the benefit plan. Payment of benefits is only made for services deemed medically necessary and appropriate. The final payment decision will be made upon submission of a claim. If you have questions about your benefits, please contact Avera Health Plans Customer Care team at 605-322-4545 or toll-free at 1-888-322-2115. This form is not all-inclusive of services requiring preauthorization. Refer to patient's Certificate of Coverage, Master Contract or Summary Plan Document for more information.

Fax this completed form to Avera Health Plans at 1-800-269-8561 or send a secure email to Pharmacy@AveraHealthPlans.com.