PRIOR AUTHORIZATION FORM

Ocrevus - Commercial



Unless otherwise indicated below, authorization quantities are limited to the manufacturer recommended dosage

P.O. Box 30192 Salt Lake City, UT 84130

Complete online at www.selecthealth.org/pa or fax back to: 801-650-3279			
For questions or clarifications, call: 800-442-3129 Patient Information			
Patient information			
Patient's Name:	Patient's Date of Birth:		
Patient's ID:	Patient's Phone #:		
Diagnosis Code(s):			
Requesting Provider Information			
Name:	Phone #:		
NPI/DEA:	Fax #:		
Address:	Supervising Physician (if requesting provider bills under a different provider) Name:		
	NPI/DEA:		
Servicing Provider Information (if different than requesting provider)			
Name of provider or facility:	Phone number:		
NPI/DEA:	Address:		
Drug Name and Strength:	Directions / SIG:		
Q1. Will Ocrevus be administered at Logan Regional or St. George Regional Hospital?			
Note, this drug is not covered at Logan Regional or St. George Regional Hospital			
☐ Yes	□ No		
Q2. Is this a reauthorization request?			
	□ No		
☐ Yes	□ No		
Q3. What is the patient's diagnosis?			
 ☐ Active forms of secondary progressive multiple sclerosis (aSPMS) ☐ Clinically isolated syndrome (CIS) ☐ Primary progressive multiple sclerosis (PPMS) 			
☐ Relapsing remitting multiple sclerosis (RRM☐ Other	io)		
Q4. If other, please specify:			

Q5. Will Ocrevus be used in combination with any other multiple sclerosis disease-modifying therapies (i.e., dimethyl fumarate, Ocrevus, Tysabri, etc.)?		
☐ Yes] No	
Q6. Has a differential diagnosis of neuroinflammatory (NMO), been excluded?	disease, such as neuromyelitis optica	
□Yes] No	
Q7. Will Ocrevus be used in combination with anothe Otezla?	r biologic medication, JAK inhibitor or	
□Yes] No	
Q8. Was the patient's diagnosis of RRMS supported criteria?	by the 2017 McDonald diagnostic	
Please select the appropriate diagnostic finding below number of lesions, and additional clinical criteria need	,	
☐ Two attacks PLUS two lesions on MRI☐ Two attacks, one lesion, and evidence of dissemir	action in space on MPI	
☐ One attack, two lesions, and evidence of dissemir	•	
☐ One attack, one lesion, and evidence of dissemina	ation in space AND time on MRI, OR	
demonstration of CSF-specific oligoclonal bands ☐ One attack, one lesion, and evidence of dissemination in space on MRI AND demonstration		
of CSF-specific oligoclonal bands		
☐ None of the above		
Q9. For new starts only, have the MRI findings been to confirm the diagnosis and findings?	eviewed and interpreted by a radiologist	
Please note: MRI and radiologist interpretation must be submitted		
□Yes] No	
Q10. Has the patient been compliant on at least one of the following preferred generic MS drugs?		
☐ Dimethyl fumarate		
☐ Fingolimod☐ Glatiramer acetate (Glatopa)		
☐ Teriflunomide		
☐ None of the above		
Q11. If none, does the patient have a highly activ	ve disease?	
☐ Yes	□ No	
Q12. If no, does the patient have elevated risk for progression to highly active multiple sclerosis based one of the following clinical factors?		
☐ Onset or diagnosis after age of 40	☐ Short inter-attack interval of relapse	

	 ☐ Expanded Disability Status Scale (EDSS) more than or equal to 3 within the first year of diagnosis ☐ High T2 lesion load, gadolinium enhancing lesions, or T1 lesions at diagnosis ☐ Infratentorial lesions (cerebellum, brain stem, or spinal cord) at diagnosis ☐ Partial or incomplete recovery or multifocal attack 	 □ Relapse requiring hospitalization or administration of corticosteroids □ Rapid accumulation of disability □ None of the above 	
Q13. Chart Notes are required for the request of this medication. Failure to provide chart notes will result in a delay in decision and/or denial. Did you attach relevant chart notes?			
☐ Yes		□No	
Q14. Add	ditional comments:		
This form is intended for SelectHealth members only. All requests for preauthorization should be sent via fax to 1-801-650-3279. Missing, inaccurate, or incomplete information may cause a delay or denial of authorization.			
	Prescriber Signature	Date	

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