

OCREVUS (ocrelizumab) [non-preferred] PRIOR AUTHORIZATION FORM

Prior authorization guidelines for MS Agents and Quantity Limits are accessible at http://www.dhs.pa.gov/provider/pharmacyservices/index.htm.

PRIOR AUTHORIZATION REQUEST INFORMATION			PRESCRIBER INFORMATION					
□New request □Renewal request total pages:			Prescriber name:					
Name of office contact:			Specialty:					
Contact's phone number:			State license #:					
LTC facility contact/phone:			NPI: MA Provider ID#:					
BENEFICIARY INFORMATION			Street address:					
Beneficiary name:			Suite #:	City/state/zip:				
Beneficiary ID#: DOB:			Phone:	Fax:				
CLINICAL INFORMATION								
Medication requested: Ocrevus (ocrelizumab) Streng			gth:	Quantity: Refills:				
Directions:								
Diagnosis (submit documentation):				Dx code (required):				
Specialty Pharmacy Drug Program: Which specialty pharmacy will be used? Diplomat Specialty Walgreen's Specialty								
 Does the beneficiary have a diagnosis of multiple sclerosis? Yes – Submit documentation of diagnosis and disease pattern. No – Submit documentation supporting the use of Ocrevus for the beneficiary's diagnosis. 								
2. Is the requested medication prescribed by a neurologist? Yes No – specialty:								
3. Does the beneficiary have evidence of active hepatitis B infection or other significant active infection					ion?	☐Yes ☐No Submit o	locumentation.	
4. Will the beneficiary receive any antineoplastic, immunosuppressive, or immune modulating therapies while using Ocrevus? Submit documentation						locumentation.		
5. Will the beneficiary receive any live vaccines while using Ocrevus and after discontinuing Ocrevus until B-cell repletion (as recommended in the FDA-approved package labeling for Ocrevus)? Submit documentation.								
6. For females of childbearing potential, does the beneficiary have documentation of a recent negative pregnancy test?							locumentation.	
7. <u>For females of childbearing potential</u> , will the beneficiary use adequate contraception treatment and for 6 months following the last infusion of Ocrevus?				uring		☐Yes ☐No Submit o	locumentation.	
Initial requests								
1. Is the beneficiary up-to-date on immunizations according to current guidelines?						☐Yes ☐No Submit	documentation.	
2. For beneficiaries being treated for a relapsing form of MS, does the ben have a history of trial and failure of or contraindication or intolerance to the public multiple sclerosis agents? Check all that apply. Aubagio Betaseron Gilenya Tecfide Avonex Copaxone (20 mg) Rebif Tysabri				□Yes	failu	Submit documentation of therapeutic failure and contraindications/intolerances.		
3. <u>For beneficiaries being treated for a relapsing form of MS</u> , is the beneficiaries treatment with Ocrevus?				☐Ye:				
Renewal requests								
Does the beneficiary continue to expe starting Ocrevus?	ondition since	□Ye:	☐ Yes Submit documentation of beneficiary's response to therapy.					
PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS - PHARMACY DIVISION								
Prescriber Signature:)ate:		
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