

## **MEDICARE FORM**

## Ocrevus® (ocrelizumab) Medication Precertification Request

Page 1 of 2

(All fields must be completed and return all pages for precertification review.)

For New Jersey HMO D-SNP: FAX: 1-833-322-0034 PHONE: 1-844-362-0934

For other lines of business: Please use other form.

Note: Ocrevus is non-preferred for relapsing forms of multiple sclerosis for MAPD plans. The preferred product is Kesimpta.

Please indicate: Start of treatment, start date:/ Co					Continuation of therapy, date of last treatment://						
Precertification Requested	Ву:				Phone	e:		Fax	:		
A. PATIENT INFORMATION											
First Name:			Last Name:								
Address:			City:					State:	ZIP:		
Home Phone:		Work F	Phone:			Ce	ell Phone:		l.		
DOB:	Allergies:							E-mail:			
Current Weight:	_	 S	Height:		inches or		cms				
B. INSURANCE INFORMATION			<u> </u>								
			Does patient have oth	ner c	overage?	ПУ	i □ No				
Aetna Member ID #:			If yes, provide ID#:								
			Insured:								
Medicare: ☐ Yes ☐ No If y	es provide ID #	————L		Medi	caid: Yes	□ No I	f ves provide	: ID #·			
C. PRESCRIBER INFORMAT					odiai 🗀 100		r you, provide	, 15 II.			
First Name:			Last Name:			(Ch	eck one): [	¬мр Г	lno F	NP [	ПРА
Address:						(0		State:	ZIP:	1.4	<u> </u>
	T_		City:	1	N. //		DEA #	State.			
Phone:	Fax:		St Lic #:		ય #:		DEA #:	1	UPIN:		
Provider E-mail:			Office Contact Name:					Phone:			
Specialty (Check one): Neurologist Primary Care Other:											
D. DISPENSING PROVIDER/	ADMINISTRATION IN	FORM/	ATION								
Place of Administration:  ☐ Self-administered ☐ Physician's Office ☐ Outpatient Infusion Center Phone:  ☐ Center Name:					Dispensing Provider/Pharmacy: Patient Selected choice  ☐ Physician's Office ☐ Retail Pharmacy ☐ Specialty Pharmacy ☐ Other:  Name:						
☐ Home Infusion Center Phone:					Address:						
Agency Name:			City:			State:	7IP·				
Administration code(s) (CP					Phone:						
Address:		7	'ID·	_	TIN:						
Phone:				_	NPI:						
TIN:	PIN:				· · · · ·						
NPI:											
E. PRODUCT INFORMATION											
Request is for Ocrevus (c	crelizumab) Dose:			F	Frequency: _						
F. DIAGNOSIS INFORMATIO	N - Please indicate pri	imary IC	D code and specify a	ny o	ther any other v	vhere ap	plicable (*).				
Primary ICD Code:				ther	· ICD Code:						
G. CLINICAL INFORMATION	- Required clinical info	ormatior	n must be completed f	or Al	LL precertificati	on reque	ests.				
For All Requests (clinical d	ocumentation requi	ired for	r all requests):								
Note: Ocrevus is non-preferred for relapsing forms of multiple sclerosis for MAPD plans. The preferred product is Kesimpta.    Yes   No   Has the patient had prior therapy with Ocrevus (ocrelizumab) within the last 365 days?   Yes   No   Has the patient had a trial and failure, intolerance, or contraindication to Kesimpta (ofatumumab)?   Please explain if there are any medical reason(s) that the patient cannot use Kesimpta (ofatumumab) when indicated for the patient's diagnosis.											
Yes No Is this infusion request in an outpatient hospital setting?  Yes No Is this request to continue previously established treatment with the requested medication?  Please explain: This is a new therapy request (patient has not received requested medication in the last 6 months)  This is a continuation of an existing treatment  Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (e.g., acetaminophen, steroids, diphenhydramine, fluids, other pre-medications or slowing of infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion?  Yes No Does the patient have severe venous access issues that require the use of special interventions only available in the											
	outpatient hospital	setting?	•								



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB					
G. CLINICAL INFORMATION (continued) -	Required clinical information must be com	pleted in its entirety for all precer	tification requests.					
<ul> <li>Yes ☐ No</li> <li>Does the patient have significant behavioral issues and/or physical or cognitive impairment that would impact the safety of the infusion therapy AND the patient does not have access to a caregiver?</li> <li>→ Please provide a description of the behavioral issue or impairment:</li></ul>								
Cardiovascular:								
Respiratory:								
Please indicate the type of multiple sclerosis the patient has been diagnosed with:								
Relapsing form of multiple sclerosis (relapsing-remitting and secondary progressive disease for those who continue to experience relapses)								
☐ Primary-progressive MS (PPMS) ☐ Clinically isolated syndrome ☐ Other (please explain):								
Yes No Is the patient taking the requested medication with any other medication used for the treatment of multiple sclerosis other than Ampyra?								
For Continuation requests (Clinical documentation required for all requests):								
☐ Yes ☐ No Is the patient experiencing disease stability or improvement while receiving the requested medication?								
H. ACKNOWLEDGEMENT								
Request Completed By (Signature Require	red):		Date://					
Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.								

The plan may request additional information or clarification, if needed, to evaluate requests.