

## *Ocrevus® (Ocrelizumab), Ocrevus Zunovo™ (ocrelizumab and hyaluronidase-ocsq) Prior Authorization (PA) Form*

Virginia | HealthKeepers, Inc. | Anthem HealthKeepers Plus Medicaid products

If the following information is not complete, correct, or legible, the PA process can be delayed. Please use one form per member.

### Member information

Last name:

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First name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Medicaid ID number:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Date of birth:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Weight in kilograms: \_\_\_\_\_

### Prescriber information

Last name:

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First name:

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NPI number:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Phone number:

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Fax number:

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### Drug information

Drug name:

\_\_\_\_\_

Drug form:

\_\_\_\_\_

Drug strength:

\_\_\_\_\_

Dosing frequency:

\_\_\_\_\_

Length of therapy:

\_\_\_\_\_

Quantity:

\_\_\_\_\_

(Form continued next page)

Member's first name:

[illegible]

For an initial request, complete the following questions to receive a six-month approval:

1. Is the member at least 18 years of age?  
☐ Yes      ☐ No
2. Has the member been screened for the presence of Hepatitis B virus (HBV) prior to initiating treatment AND does not have active disease (for example, positive HBsAg and anti-HBV tests)? AND  
☐ Yes      ☐ No
3. Has the member had baseline serum immunoglobulin assessed? AND  
☐ Yes      ☐ No
4. Will the member not receive live or live attenuated vaccines while on therapy or within four weeks prior to the initiation of treatment? AND  
☐ Yes      ☐ No
5. Is the member free of an active infection? AND  
☐ Yes      ☐ No
6. Will Ocrevus/Ocrevus Zunovo be used as a single therapy? AND  
☐ Yes      ☐ No
7. Has the member not received a dose of ocrelizumab or ublituximab within the past five months? AND  
☐ Yes      ☐ No
8. Does the member have a confirmed diagnosis of multiple sclerosis (MS) as documented by laboratory report (for example, MRI)? AND
  - a. Does the member have a diagnosis of a relapsing form of MS for example, relapsing-remitting MS (RRMS)\*, active secondary progressive disease (SPMS)\*\*, or clinically isolated syndrome (CIS)\*\*\*? OR
  - b. Does the member have a diagnosis of primary progressive MS (PPMS)\*\*\*\*? AND
    - i. Is the member less than 65 years of age? AND
    - ii. Does the member have an expanded disability status scale (EDSS) score of  $\leq 6.5$ ? AND☐ Yes      ☐ No

(Form continued on next page.)

Member's first name:

[illegible]

1. Does the member continue to meet the relevant criteria identified in the initial criteria? AND  
☐ Yes ☐ No
2. Does the member have an absence of unacceptable toxicity from the drug? AND  
☐ Yes ☐ No
3. Is the member being continuously monitored for response to therapy that indicates a beneficial response?  
☐ Yes ☐ No

<p>(Definitive diagnosis of MS with a relapsing-remitting course is based upon both dissemination in time and space. Unless contraindicated, MRI should be obtained (even if criteria are met).</p>	
<p>Dissemination in time (Development/appearance of new CNS lesions over time)</p>	<p>Dissemination in space (Development of lesions in distinct anatomical locations within the CNS - multifocal)</p>
<ul style="list-style-type: none"> <li>• ≥ Two clinical attacks; OR</li> <li>• One clinical attack and one of the following:                             <ul style="list-style-type: none"> <li>○ MRI indicating simultaneous presence of gadolinium-enhancing and non-enhancing lesions at any time or by a new T2-hyperintense or gadolinium-enhancing lesion on follow-up MRI compared to baseline scan</li> <li>○ CSF-specific oligoclonal bands</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>• ≥ Two lesions;</li> <li>• One lesion and one of the following:                             <ul style="list-style-type: none"> <li>○ Clear-cut historical evidence of a previous attack involving a lesion in a distinct anatomical location</li> <li>○ MRI indicating ≥ one T2-hyperintense lesions characteristic of MS in ≥ two of four areas of the CNS (periventricular, cortical or juxtacortical, infratentorial, or spinal cord)</li> </ul> </li> </ul>
<p><b>**Active secondary progressive MS (SPMS) is defined as the following:</b></p> <ul style="list-style-type: none"> <li>• Expanded Disability Status Scale (EDSS) score ≥ 3.0; and</li> <li>• Disease is progressive ≥ three months following an initial relapsing-remitting course (for example., EDSS score increase by 1.0 in members with EDSS ≤5.5 or increase by 0.5 in members with EDSS ≥6); AND                             <ul style="list-style-type: none"> <li>○ ≥ One relapse within the previous two years; OR</li> <li>○ Member has gadolinium-enhancing activity OR new or unequivocally enlarging T2 contrast-enhancing lesions as evidenced by MRI</li> </ul> </li> </ul>	

(Form continued on next page.)

Member's first name:

[illegible]

- A monophasic clinical episode with member-reported symptoms and objective findings reflecting a focal or multifocal inflammatory demyelinating event in the CNS
- Neurologic symptom duration of at least 24 hours, with or without recovery
- Absence of fever or infection
- Member is not known to have multiple sclerosis

- One year of disability progression independent of clinical relapse; **AND**
- **Two** of the following:
  - ≥ One T2-hyperintense lesion characteristic of MS in one or more of the following regions of the CNS: periventricular, cortical or juxtacortical, or infratentorial
  - ≥ Two T2-hyperintense lesions in the spinal cord
  - Presence of CSF-specific oligoclonal bands

The completed form may be **faxed to 844-512-7020**.