

Ocrelizumab (Ocrevus™)

Goal(s):

- Restrict use of ocrelizumab in patients with relapsing-remitting multiple sclerosis (RRMS) to those who have failed multiple drugs for the treatment of RRMS.
- Ensure appropriate baseline monitoring to minimize patient harm.

Length of Authorization:

- 6 to 12 months

Requires PA:

- Ocrevus™ (ocrelizumab) pharmacy or physician administered claims

Covered Alternatives:

- Current PMPDP preferred drug list per OAR 410-121-0030 at www.orpdl.org
- Searchable site for Oregon FFS Drug Class listed at www.orpdl.org/drugs/

Approval Criteria		
1. What diagnosis is being treated?	Record ICD10 code.	
2. Is the medication FDA-approved or compendia-supported for the requested indication?	Yes: Go to #3	No: Pass to RPh. Deny; medical appropriateness
3. Is the drug being used to treat an OHP-funded condition?	Yes: Go to #4	No: Pass to RPh. Deny; not funded by the OHP.
4. Is this a request for continuation of therapy?	Yes: Go to Renewal Criteria	No: Go to #5
5. Is the patient an adult (age ≥18 years) diagnosed with relapsing multiple sclerosis?	Yes: Go to #6	No: Go to #7
6. Has the patient failed trials for at least 2 drugs indicated for the treatment of relapsing multiple sclerosis?	Yes: Document drug and dates trialed: 1. _____ (dates) 2. _____ (dates) Go to #7	No: Pass to RPh. Deny; medical appropriateness
7. Has the patient been screened for an active Hepatitis B infection?	Yes: Go to #8	No: Pass to RPh. Deny; medical appropriateness

Approval Criteria

8. Is the drug prescribed by or in consultation with a neurologist who regularly treats multiple sclerosis?

Yes: Approve ocrelizumab 300 mg every 2 weeks x 2 doses followed by 600mg IV every 6 months for 12 months

No: Pass to RPh. Deny; medical appropriateness

Renewal Criteria

1. Has the patient's condition improved as assessed by the prescribing physician and physician attests to patient's improvement.

Yes: Approve for 12 months.

Document baseline assessment and physician attestation received.

No: Pass to RPh; Deny; medical appropriateness.

P&T/DUR Review: 6/21(DM); 6/20; 11/17 (DM); 1/17
Implementation: 7/1/20; 1/1/18; 4/1/17