

# OPTIMA HEALTH COMMUNITY CARE

## PHARMACY/MEDICAL PRIOR AUTHORIZATION REQUEST\*

Directions: *The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-844-348-3720. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. Incomplete form will delay authorization process.*

**Drug Requested:** Ocrevus™ (ocrelizumab) Injection (J-2350/C9494) (Medical) (**Non-Preferred**)

**DRUG INFORMATION:** Complete all information below. Authorization process will be delayed if incomplete.

Drug Form/Strength: \_\_\_\_\_

Dosing Schedule: \_\_\_\_\_ Length of Therapy: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code: \_\_\_\_\_

### **RECOMMENDED DOSAGE AND ADMINISTRATION:**

**INITIAL DOSE:** 300 mg intravenous infusion, followed 2 weeks later by a 2<sup>nd</sup> 300 mg intravenous infusion

**SUBSEQUENT DOSES:** single 600 mg intravenous infusion every 6 months

**Medical notes MUST be submitted with this request to support each line checked.**

**CLINICAL CRITERIA:** All boxes that apply **MUST** be checked to qualify. Incomplete information or medical notes are **not** attached with this form request will delay the authorization process.

**Please check all below for Primary Progressive MS indication. If **NOT** checked, authorization process will be delayed. Medical notes **MUST** be attached with this request.**

- Prescriber is a **Neurologist**
- Patient has a **confirmed** diagnosis of **Primary Progressive MS**

**Please check all below for Relapsing-Remitting MS indication. If **NOT** checked, authorization process will be delayed. Medical notes **MUST** be attached with this request.**

- Prescriber is a **Neurologist**
- Patient has a confirmed diagnosis of relapsing-remitting MS
- Patient has had at least one medically documented clinical relapse within 12 months
- Patient has completed a trial and has failed at least **TWO (2)** of the following agents: (**check each that have been tried**):

<input type="checkbox"/> Aubagio® (teriflunomide)	<input type="checkbox"/> Betaseron® (IFN beta-1a)	<input type="checkbox"/> Extavia® (IFN beta-1a)
<input type="checkbox"/> Avonex® (IFN beta-1b)	<input type="checkbox"/> Copaxone® (glatiramer acetate)	<input type="checkbox"/> Gilenya® (fingolimod)
<input type="checkbox"/> Lemtrada® (alemtuzumab)	<input type="checkbox"/> Rebif® (IFN beta-1a)	<input type="checkbox"/> Plegridy® (pegylated-IFN beta-1a)
<input type="checkbox"/> Tecfidera® (dimethyl fumarate)	<input type="checkbox"/> Tysabri® (natalizumab)	

(signature on next page)

**Medication being provided by (check applicable box below):**

Physician's office

OR

Specialty Pharmacy:

PropriumRx

*\*Use of samples to initiate therapy **does not** meet step-edit/preauthorization criteria.\**

**\*Previous therapies will be verified through pharmacy paid claims or submitted chart notes.\***

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

**DEA OR NPI #:** \_\_\_\_\_

REVISED/UPDATED: 8/1/2017; 12/27/2017.