



Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient?  Yes  No

**What is your patient's diagnosis?**

- A relapsing form of multiple sclerosis – Please Note: Examples include clinically isolated syndrome, relapsing remitting disease, and active secondary progressive disease
- Primary Progressive multiple sclerosis
- Other diagnoses or indications

**Clinical Information:**

(if Relapsing forms of MS or PPMS) Is the requested medication being prescribed by (or in consultation with) a neurologist or a physician who specializes in the treatment of multiple sclerosis?  Yes  No

(if Relapsing forms of MS or PPMS) Is the requested medication to be used in combination with other disease-modifying agents for multiple sclerosis? Please Note: Examples of disease modifying agents for MS include Avonex, Betaseron, Briumvi, Rebif, Plegridy, Ponvory, Copaxone, Glatopa, glatiramer acetate subcutaneous injection, Gilenya, fingolimod capsules, Aubagio, Tecfidera, dimethyl fumarate delayed-release capsules, Tysabri, Mayzent, Mavenclad, Vumerity, Bafiertam, Zeposia, Kesimpta, teriflunomide tablets, Tyruko, Tascenso ODT, Ocrevus Zunovo, and Lemtrada.  Yes  No

(if Relapsing forms of MS) Is the patient currently receiving Ocrevus?  Yes  No

(if currently receiving) Has the patient already received at least 1 year of therapy with Ocrevus? Please Note: Answer No if the patient has received less than 1 year of therapy or if the patient is restarting therapy with Ocrevus.  Yes  No

(if Currently receiving Ocrevus for at least 1 Year) Has the patient experienced a beneficial clinical response when assessed by at least one objective measure? Please Note: Examples include stabilization or reduced worsening in disease activity as evaluated by magnetic resonance imaging (MRI) [absence or a decrease in gadolinium enhancing lesions, decrease in the number of new or enlarging T2 lesions]; stabilization or reduced worsening on the Expanded Disability Status Scale (EDSS) score; achievement in criteria for No Evidence of Disease Activity-3 (NEDA-3) or NEDA-4; improvement on the fatigue symptom and impact questionnaire-relapsing multiple sclerosis (FSIQ-RMS) scale; reduction or absence of relapses; improvement or maintenance on a the six-minute walk test or 12-Item MS Walking Scale; improvement on the Multiple Sclerosis Functional Composite (MSFC) score; and/or attenuation of brain volume loss.  Yes  No

(if no) Has the patient experienced stabilization, slowed progression, or improvement in at least one symptom such as motor function, fatigue, vision, bowel/bladder function, spasticity, walking/gait, or pain/numbness/tingling sensation?  Yes  No

*(Please note: there are different preferred products depending on your patient's plan. Please refer to the applicable Cigna health care professional resource [e.g. [cignaforhcp.com](http://cignaforhcp.com)] to determine benefit availability and the terms and conditions of coverage)*

**Additional Information:**

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

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

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*NDC number is required on the medical claims to confirm claim is payable for the drug Betaseron. The NDC number can be found on the drug packaging. In addition you may refer to the Crosswalk of HCPCS Codes Requiring NDC on Claims at the Cigna for Health Care Professionals website ([CignaforHCP.com](http://CignaforHCP.com) > Resources > Clinical Reimbursement Policies and Payment Policies >."*

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