MEDICARE FORM

Ocrevus® (ocrelizumab) Medication Precertification Request

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

For Medicare Advantage Part B:
Phone: 1-866-503-0857 (TTY: 711)
FAX: 1-844-268-7263

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.

Ocrevus is preferred for MA plans.

Please indicate: ☐ Start of treatment, start date: ____/____/____ ☐ Continuation of therapy, date of last treatment: ____/____/____

Precertification Requested By: _____ Phone: _____ Fax: _____

A. PATIENT INFORMATION

First Name:	Last Name:		
Address:	City:	State:	ZIP:
Home Phone:	Work Phone:	Cell Phone:	
DOB:	Allergies:	E-mail:	

Current Weight: _____ lbs or _____ kgs Height: _____ inches or _____ cms

B. INSURANCE INFORMATION

Aetna Member ID #: _____ Does patient have other coverage? ☐ Yes ☐ No
Group #: _____ If yes, provide ID#: _____ Carrier Name: _____
Insured: _____ Insured: _____

Medicare: ☐ Yes ☐ No If yes, provide ID #: **Medicaid:** ☐ Yes ☐ No If yes, provide ID #:

C. PRESCRIBER INFORMATION

First Name:		Last Name: *(Check one):* ☐ M.D. ☐ D.O. ☐ N.P. ☐ P.A.			
Address:		City:		State:	ZIP:
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	UPIN:
Provider E-mail:		Office Contact Name:		Phone:	

Specialty *(Check one):* ☐ **Neurologist** ☐ **Primary Care** **Other**: _____

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration:
☐ Self-administered ☐ Physician's Office
☐ Outpatient Infusion Center Phone: _____
Center Name: _____
☐ Home Infusion Center Phone: _____
Agency Name: _____
☐ Administration code(s) (CPT): _____
Address: _____
City: _____ State: _____ ZIP: _____
Phone: _____ Fax: _____
TIN: _____ PIN: _____
NPI: _____

Dispensing Provider/Pharmacy: *Patient Selected choice*
☐ Physician's Office ☐ Retail Pharmacy
☐ Specialty Pharmacy ☐ Other: _____
Name: _____
Address: _____
City: _____ State: _____ ZIP: _____
Phone: _____ Fax: _____
TIN: _____ PIN: _____
NPI: _____

E. PRODUCT INFORMATION

Request is for Ocrevus (ocrelizumab)
Dose: _____ **Frequency:** _____ **HCPCS Code:** _____

F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other any other where applicable (*).

Primary ICD Code: _____ ☐ **Other ICD Code**: _____

G. CLINICAL INFORMATION - Required clinical information must be completed for ALL precertification requests.

For *Initiation* Requests (clinical documentation required for all requests):

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

☐ 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.

_____

For *All* Requests (clinical documentation required for all requests):

☐ 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
☐ Yes ☐ No 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 completed in its entirety for all precertification requests.

☐ Yes ☐ No 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?

→ Please provide a description of the behavioral issue or impairment: _____

☐ Yes ☐ No Is the patient medically unstable which may include respiratory, cardiovascular, or renal conditions that may limit the member's ability to tolerate a large volume or load or predispose the member to a severe adverse event that cannot be managed in an alternate setting without appropriate medical personnel and equipment?

→ Please provide a description of the condition:

☐ Cardiovascular: _____

☐ Respiratory: _____

☐ Renal: _____

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 Required):*** _____ **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.