



Please Fax Completed Form To: 888-898-9113

Please Send a Copy of The Patient's Insurance Cards (Front & Back)

Address: City, State, Zip:										
Address:	PATIENT INF	ORMATION (Co	x Existing Chart)	PRESCRIBER	INFORMAT	ION				
NPI #:	Name: DOB:				Prescriber Name:					
NPI #:	Address:			<u>.</u>	State License: _					
Email: SS#: City, State, Zip: Phone: Fax: Phone: Phone: Fax: Phone: Phone: Fax: Phone: Fax: Phone: Fax: Phone: Phone					NPI #:	#: Tax ID:				
Gender: M F Weight: (lbs) Ht: Office Contact: Phone: Fax: Allergies: Office Contact: Phone: Phone: Allergies: Office Contact: Phone: All	Phone: Alt. Phone:			 	Address:					
Phone:	Email:		SS#:							
Allergies: Office Contact: Phone:					Phone:		Fax: _			
Secondary Insurance (If Applicable):					Office Contact:	Phone:				
Plan #:										
Plan #:	Primary Insuran	ice:			Secondary Insur	ance (If Appli	cable):			
Group #:										
RX Card (PBM):										
BIN:										
G35 Multiple Sclerosis										
G35 Multiple Sclerosis Other Diagnosis/ICD-10 Code:										
Has Patient Completed the First 2 Loading Doses of Ocrevus*? Yes No Expected Date of First/Next Infusion: Date of Last MRI: Past DMT Therapies: Hepatitis B (HBsAg and anti-HBV) Test Results: Positive Negative Quantitative Serum Immunoglobulins Test Results: Please Check to Confirm Understanding: According to immunization guidelines, live or live-attenuated vaccines should be administered at least 4 weeks prior to initiation of OCREVUS* and, whenever possible, for non-live vaccines at least 2 weeks prior to initiation of OCREVUS*. **Obtain the following labs at prior to start of treatment and at frequency CBC CMP CRP ESR LFTs X-Ray Other: OCREVUS* ORDERS Prescription type: New start Restart Continued therapy Total Doses Received: Date of Last Injection/Infusion: Medication Dose Administration Refills Initial Dose: 600 mg dose administered as 2 separate intravenous infusions 2 weeks apart. Maintenance Dose: 600 mg dose administered once every 6 months; 2 infusion options to choose from: Option 1: Single infusion administered over approximately 3.5 to 4 hours. Option 2: Single infusion administered over approximately 2 hours (for eligible patients who have not experienced a serious infusion reaction with any previous OCREVUS infusion)	CLINICAL INF	ORIVIATION								
Has Patient Completed the First 2 Loading Doses of Ocrevus*? Yes No Expected Date of First/Next Infusion:	☐ G35 Multiple Sclerosis ☐ Other Diagnosis/ICD-10 Code:									
Date of Last MRI:	Has Patient Con									
Hepatitis B (HBsAg and anti-HBV) Test Results:										
Please Check to Confirm Understanding: According to immunization guidelines, live or live-attenuated vaccines should be administered at least 4 weeks prior to initiation of OCREVUS® and, whenever possible, for non-live vaccines at least 2 weeks prior to initiation of OCREVUS®. **Obtain the following labs at prior to start of treatment and at frequency CBC CMP CRP ESR LFTS X-Ray Other: OCREVUS® ORDERS Prescription type: New start Restart Continued therapy Total Doses Received: Date of Last Injection/Infusion: Medication Dose Administration Refills										
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Prescription type: New start Restart Continued therapy Total Doses Received: Date of Last Injection/Infusion: Medication Dose Administration Refills Ocrevus* (Ocrelizumab)	** Obtain the following labs at prior to start of treatment and at frequency \(\subseteq \text{CBC} \subseteq \text{CMP} \subseteq \text{CRP} \subseteq \text{ESR} \subseteq \text{LFTs} \subseteq \text{X-Ray} \subseteq \text{Other:} \(\left_{\text{CMP}} \subseteq \text{CRP}									
Medication Dose Administration Refills Initial Dose: 600 mg dose administered as 2 separate intravenous infusions 2 weeks apart. Maintenance Dose: 600 mg dose administered once every 6 months; 2 infusion options to choose from: Option 1: Single infusion administered over approximately 3.5 to 4 hours. Option 2: Single infusion administered over approximately 2 hours (for eligible patients who have not experienced a serious infusion reaction with any previous OCREVUS infusion) Special Instructions:	OCREVUS® ORDERS									
□ Ocrevus® (ocrelizumab) □ 300 mg/10 mL (30 mg/mL) single-dose vial □ Option 1: Single infusion administered over approximately 3.5 to 4 hours. □ Option 2: Single infusion administered over approximately 2 hours (for eligible patients who have not experienced a serious infusion reaction with any previous OCREVUS infusion) Special Instructions:	Prescription typ	e: 🗆 New start 🗀	Restart 🗆 Co	ontinued therapy Total	Doses Received: _		Date of Last I	Injection/Infusion	on:	
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Pre- Medication Route Dose	Special Instructi	ions:								
	Pre- Medication			Route		Dose				
□ Acetaminophen □ PO □ 500mg □ 650mg □ 1000mg	☐ Acetaminophen			□РО		□ 500mg □ 650mg □ 1000mg				
☐ Methylprednisolone (Solu-Medrol) ☐ IV ☐ 60mg ☐ 100 mg ☐mg	☐ Methylprednisolone (Solu-Medrol)			□IV						
	☐ Diphenhydramine (Benadryl)			□ IV □ PO		□ 25mg □ 50mg				
□ Other:				-		0	0			

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ANAPHYLACTIC REACTION (AR):
☐ EpiPen® Auto-injector 0.3 mg (1:1000) Inject IM -or- SubQ to patients who weigh ≥ 66 lbs (≥ 30 kg); may repeat in 3-5 mins x 1 if necessary
☐ EpiPen Jr® Auto-injector 0.15mg (1:2000) Inject IM -or- SubQ to patients who weigh 33 - 66 lbs (15-30 kg): may repeat in 3-5 mins x 1 if necessary
☐ Diphenhydramine 50mg (1mL) - Administer 50 mg VIA slow IVP, administer IM if no IV access; may repeat x 1 after 10 mins, if necessary
☐ Methylprednisolone 40mg - administer 40 mg IVP -or- IM if no IV access
☐ Sodium Chloride 0.9% 500 mL infuse IV at a rate of up to 999 mL/hr
□ Other:
SIGNATURE
We hereby authorize Valustar to provide all supplies and additional services (nursing/patient training) required to provide and deliver the medicine as prescribed in this referral.
X Date:
Prescriber Signature

To ensure payment by insurance carrier, please include supporting clinical documentation for specified ICD 10 Code, demographic, and insurance information along with faxed order. Initial appointment will be verified upon insurance approval.

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