ALL ORDERS MUST BE SIGNED, DATED AND TIMED BY PHYSICIAN				
Allergies/Reactions:	☐ Puyallup Infusion Center - Fax: 253-697	7-5066	☐ Gig Harbor Inf	usion Services - Fax: 253-530-8069
	☐ Allenmore Ambulatory Infusion Service:	s - Fax: 253-864-4052	2 DHEC Infusion	Center - Fax: 509-755-5845
	☐ Auburn Infusion Services - Fax: 253-87	6-8282	☐ North Spokan	e Infusion Center - Fax: 509-232-2531
ORDERS WITH CHECK BOXES When box ne	an order is optional (those with check ext to the order. Orders left unchecked	boxes), physicians will not be initiated.	are responsible f	or indicating a check mark in the
Ubilituximab (Briumvi):				
Patient Name:		Requested [Date of Service:	/
Date of Birth://	Patient Phone Number: ()			🗖 May leave message
		ICD -10 Cod	lo.	
Discourie DM Rate Calcusia		<u>ICD -10 C00</u>	<u>ie</u> :	
Diagnosis: ☐ Multiple Sclerosis		U		
Required: H&P with documentation to support above diagnosis including ICD-10 code and supporting labs and documentation **If required documentation not received with order, scheduling of treatment will be delayed until complete information is available**				
Baseline Labs Required: • Hepatitis B • Quantitative serum immunoglobuli • Pregnancy test	ns			
Maintenance labs required: ☐ Pregnancy testing prior to each dose in patients' who may become pregnant				
☑ IV Access: Access and/or maintain IV site in accordance with MHS IV Therapy P&P: Peripheral IV Device Site Selection, Insertion, Maintenance, and Discontinuation; and Maintenance of Central Venous Catheters-Flushing, Dressing Changes and Removal.				
Premeds: ☐ Methylprednisolone 100 mg IVP ☐ Diphenhydramine 25 mg IVP ☐ Acteaminophen 650 mg PO prn				
Treatment Regimen: Ublituximab (Briumvi) 150 mg IV x 1 on day 1, followed by 450 mg IV once 2 weeks later; subsequent doses of 450 mg IV once every 24 weeks (beginning 24 weeks after the first dose of 150 mg)				
✓ Vital signs: Check vital signs prior to Contact provider if systolic BP>180;	•	HR >110; temp >	>38C (100.4F)	
If Hypersensitivity reaction (fever, chill Consult MultiCare hypersensitivity Notify provider of reaction, assessr	guideline for treatment managem			
Code Status: Please note, patients will be considered FULL Code unless marked otherwise. If the patient has a POLST, advance directive or living will, please include a copy with the orders.				
Was consent obtained: ☐ Yes ☐ No (if yes, please send DOCUMENTA	TION of consent	t with order)	
Provider Signature	Print Name	 -	Date	Time
	pontont more hadio		Oud aug	vnivos in 12 manthatt
Another brand of drug, identical in form and a	content, may be dispensed unless che	ckea 🖵	Orders ex	xpires in 12 months**

Patient Identification - Always Attach Patient Label

Name:

MRN #:

CSN #:

Age / Sex and Gender:

Pre-printed Order

MULTIPLE SCLEROSIS

MultiCare 🕰

