ALL ORDERS MUST BE SIGNED, DATED AND TIMED BY PHYSICIAN			
Allergies/Reactions:	Fax all infusions to: 833-380-8800 Please mark the appropriate infusion center:	☐ Allenmore Infusion Center☐ Auburn Infusion Center☐ Gig Harbor Infusion Services☐ Puyallup Infusion Center☐	☐ DHEC Infusion Center☐ North Spokane Infusion Center☐ North Star Lodge Infusion Center☐
ORDERS WITH CHECK BOXES When an order is optional (those with check boxes), physicians are responsible for indicating a check mark in the box next to the order. Orders left unchecked will not be initiated.			
Efgartigimod alfa (Vyvgart):			
Patient Name:Requested Date of Service://			
Diagnosis: ☐ Myasthenia Gravis		ICD -10 Code:	
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: CBC * Patients should be up to date with all immunizations before initiating therapy. Avoid the use of live vaccines in patients undergoing efgartigimod treatment			
☑ 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.			
Treatment Regimen: once weekly for 4 weeks *Subsequent cycles may be administered based on clinical evaluation and no sooner than 50 days from start of the previous treatment cycle □ Efgartigimod alfa (Vyvgart) 10 mg/kg IV infusion over 1 hour for patients less than 120 kg □ Efgartigimod alfa (Vyvgart) 1200 mg IV infusion over 1 hour for patients >/= 120 kg			
✓ Vital signs: Check vital signs prior to and at completion of infusion. Contact provider if systolic BP>180; diastolic BP>100; systolic BP<90; HR >110; temp >38C (100.4F)			
If Hypersensitivity reaction (fever, chills, hypotension, rigors, itching, rash, etc.) • Consult MultiCare hypersensitivity guideline for treatment management • Notify provider of reaction, assessment and need for further orders			
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 DOCUMENTATION of consent with order)			
Provider Signature	Print Name	 Date	Time
Another brand of drug, identical in form and content,			xpires in 12 months**
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Patient Identification - Always Attach Patient Label

Name:

MRN #:

CSN #:

Age / Sex and Gender:

Pre-printed Order

MYASTHENIA GRAVIS

MultiCare 🕰

