

1 PATIENT INFORMATION Patient Name: _____ Phone: _____ Address: _____ City: _____ State: _____ Zip: _____ MRN #: _____ DOB: _____ Drug Allergies: _____	2 PRESCRIBER INFORMATION Prescriber's Name: _____ DEA#: _____ NPI: _____ Clinic/Facility Name: _____ Address: _____ City: _____ State: _____ ZIP: _____ Phone: _____ Fax: _____
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3 Instructions to Provider
 All orders with ✓ will be placed unless otherwise noted. Please fax completed order form to 206-326-2139. For drug prior authorization, call 1-888-767-4670 or visit <https://wa-provider.kaiserpermanente.org/provider-manual/clinical-review/preservice>. Lab orders are not included on this form. KPWAHI does not provide laboratory monitoring.

4 CLINICAL INFORMATION
 Diagnosis (ICD-10 code): _____ Date of Last Dose: _____

5 ALGLUCOSIDASE ALFA PRESCRIPTION INFORMATION

First Dose: No Yes Weight: _____ kg Date Recorded: _____

Alglucosidase alfa (LUMIZYME) in 0.9 % sodium chloride 500 mL IV infusion:
 Dose: 20mg/kg _____ mg/kg x weight (kg) Route: Intravenous Frequency: Every 2 weeks
 Refills: 11 months Other _____
 Infusion Access: PIV CVAD Other: _____
 Infusion Rates, and Supplies: Per protocol (See below) Special Instructions (Specify below)
 Patient's Current Home Care/Specialty Pharmacy: _____

Infusion Reaction Medications & Supplies:

✓ Hydrocortisone sodium succinate injectable 100 mg IV Sig: Once PRN for hypersensitivity ✓ Epinephrine Auto-Injector <input type="checkbox"/> 0.15mg <input type="checkbox"/> 0.3mg QTY: 2 Sig: Inject into lateral thigh muscle for severe allergic reaction. Seek medical attention after use.	✓ Diphenhydramine injectable 25 mg IV Sig: Once PRN, may repeat x1 for urticaria, pruritis, shortness of breath ✓ Sodium Chloride 0.9% IV 250ml Bag Sig: Once PRN for anaphylaxis
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✓ **Sodium Chloride 0.9% IV Flush:** Flush 10 ml IV before/after medication administration or as needed for line maintenance

Labs /Special Instructions/Pre-Meds: _____

Alglucosidase alfa Infusion Protocol: <ul style="list-style-type: none"> • Infuse per manufacturer guidelines, total ~4 hours <ul style="list-style-type: none"> ○ Initiate at 1mg/kg/hr, if tolerated increase by 2mg/kg/hr every 30 min to a max rate of 7mg/kg/hr • Monitor vital signs (Temp, BP, HR, RR) every 15 minutes x 4; then every 30 minutes x2; then every 60 minutes until completion of infusion • Documentation must include: <ul style="list-style-type: none"> ○ Start and end time of infusion 	<ul style="list-style-type: none"> • Observe patient for signs of infusion rate-related adverse reactions: <ul style="list-style-type: none"> ○ Blood pressure changes, increased pulse rate ○ Fever, chills ○ Headache ○ Chest, back or hip pain ○ Dyspnea ○ Mild erythema
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6 PHYSICIAN SIGNATURE REQUIRED

X _____ SUBSTITUTION PERMITTED (Date)	X _____ DISPENSE AS WRITTEN (Date)
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