Lemtrada (Alemtuzumab)

Provider Order Form rev. 4/10/2022

PATIENT INFORMATION	Referral	Status (check one):	🗆 New Referral	Updated Order	r 🗌 Order Renewal	
Patient Name:		DOB:				
NKDA 🗌 Allergies:		Weight	Please specify	r: □ lbs □ kg	Height:	
Patient Status (check one): New to Therapy Continuing Therapy		Last Treatment Date:		Next Due Date:		
ICD-10 code (required): ICD-10 c	descriptio	on:				
REQUIRED: Demographics & Most Recent: H&P, clini past tried and/or failed therapies, intolerand					ny	
PRESCRIPTION						
SUPPORTING DOCUMENTATION Ensure baseline labs have been drawn & provide results: Ensure patient has taken & prescribed an anti-viral: Acyclovir 400mg Home medications: Zyrtec 10mg / □ Hydroxyzine 50mg / Zantac 150mg / Pepcid 20mg (Staff to verify patient has taken.) NURSING		PRE-MEDICATION ORDERS (ADDITIONAL) Ibuprofen (Advil) 400mg PO (If indicated, acetaminophen will be held) cetirizine (Zyrtec) 10mg PO loratadine (Claritin) 10mg PO ranitidine (Zantac) 150mg PO Methylprednisolone (Solu-Medrol) mg IV mixed in ml NS over 1 hour on days: dexamethasone 80mg IV mixed in 100ml NS over 1 hour on days:				
 Provide nursing care per AdaptIV Infusion Nursing Procedures, inclureaction management and post-procedure observation Verify patient & provider are enrolled/authorized in REMS. Ensure REMS authorization call has occurred prior to infusion. Provide patient with What You Need to Know about Lemtrada Treat 	_			Route:		
and Infusion Reactions: A Patient Guide Complete & submit LEMTRADA REMS Infusion Checklist upon completion of each treatment cycle. LABORATORY ORDERS CBC with differential on days: CMP		 Dose & Rd Frequence Mix in 100 (protect find) Flush with (infuse at) 	(Lemtrada) year o oute: 12mg intraver y: daily for 5 days ml 0.9% sodium ch rom light) n 0.9% sodium chlo same rate as Lemtr	nous infusion nloride, infuse over fo ride at the completio	on of infusion	
 PRE-MEDICATION ORDERS (REQUIRED) acetaminophen (Tylenol) 1000mg PO each day diphenhydramine (Benadryl) 50mg PO each day methylprednisolone (Solu-Medrol) 1000mg IV mixed in 100ml 0.9% over 1 hour on days 1, 2, 3 of each cycle treatment *Unless contraindicated, the above will be given with each treatment cycle. 	6 NS	 Dose Frequ Mix in (prot Flush (infus) 	& Route: 12mg intr uency: daily for 3 da n 100ml 0.9% sodiu ect from light) with 0.9% sodium se at same rate as L	ays m chloride, infuse ov chloride at the comp	ver four hours pletion of infusion	

Ordering Provider: Initial here _____ and proceed to the next page.



PRN MEDICATIONS (GIVEN BASED ON PATIENT ASSESSMENT)

- □ acetaminophen (Tylenol) 650mg PO every 6 hours for **mild** pain or fever (alternate with ibuprofen)
- □ ibuprofen (Advil) 400mg PO every 4 hours for **mild** pain or fever (alternate with acetaminophen)
- ketorolac [Toradol] 30mg SIVP x 1 for moderate to severe pain/ headache [Do not give with elevated creatinine. If pain/headache not relieved 15-20 minutes after administration notify provider. Consider stopping infusion and transfer to an acute care setting.]
- □ diphenhydramine [Benadryl] 25-50mg PO every 4 hours for **mild** itching or hives
- □ hydroxyzine 50mg PO every 12 hours for **mild** itching or hives (consider if diphenhydramine already given)
- □ diphenhydramine 25-50mg SIVP, for **severe** itching, rash, or shortness of breath. May repeat 25-50mg SIVP × 1
- ondansetron (Zofran) 4mg SIVP every 4-6 hours for nausea/vomiting, may repeat 4mg SIVP x1 for a max dose of 8mg

HYPERTENSION MANAGEMENT

SBP > 30mmhg above baseline or SBP > or = 160

□ clonidine 0.1mg PO x 1

SBP > 40mmhg above baseline or BP > or = 170/100 Notify provider and repeat VS q 5 minutes

□ hydralazine 10mg SIVP over 2-3 minutes, may repeat dose x 1 in 20 minutes (Do not give if heart rate >100 BPM)

SPECIAL INSTRUCTIONS

INFUSION/MONITORING PARAMETERS

- □ If any of the following below are present, stop infusion, monitor vital signs every 5 minutes and notify provider.
- □ If blood pressure remains >40mmhg above baseline or \ge 170/100 after administration of PRN medications.
- □ If chest pain, pressure or tightness that is not relieved with PRN medication administration.
- □ If heart rate < 50 or > 110 and patient symptomatic; dizziness, shortness of breath, chest pain, pressure or discomfort.
- □ If SPO2 < 92% with or without supplemental oxygen.
- \square Any sudden onset or change in neurological symptoms.
- * Premedicate patients with high dose corticosteroids (1,000 mg methylprednisolone or equivalent) immediately prior to LEMTRADA infusion and for the first 3 days of each treatment course.
- * Administer anti-viral prophylaxis for herpetic viral infections starting on the first day of each treatment course and continue for a minimum of two months following treatment with LEMTRADA or until the CD4+ lymphocytecount is at least 200 cells per microliter, whichever occurs later.
- * Observe patients for infusion reactions during and for at least 2 hours after each LEMTRADA infusion.
- * Conduct the following laboratory tests at baseline and at periodic intervals until 48 months after the last treatment course of LEMTRADA in order to monitor for early signs of potentially serious adverse effects:
 - Complete blood count (CBC) with differential (prior to treatmentinitiation and at monthly intervals thereafter)
 - Serum creatinine levels (prior to treatment initiation and at monthly intervals thereafter)
 - Urinalysis with urine cell counts (prior to treatment initiation and at monthly intervals thereafter)
 - A test of thyroid function, such as thyroid stimulating hormone(TSH) level (prior to treatment initiation and every 3 months thereafter)
 - Serum transaminases (alanine aminotransferase [ALT] and aspartate aminotransferase [AST]) and total bilirubin levels (prior to treatment initiation and periodically thereafter)
- * Providers choosing to refer patients for Lemtrada infusions must complete this order set. Outside order sets will not be accepted. Please direct any questions or comments regarding the use of this order set to Matt Munden, RN Director of Nursing or Andrew Lasher, MD Chief Medical Officer.

PROVIDER INFORMATION

Referral Coordinator Name:		Referral Coordinator Email:				
Ordering Provider:	Provider NPI:					
Referring Practice Name:		Phone:	Fax:			
Practice Address:		City:	State:	Zip Code:		
Provider Name (Print)	Provider Signature			Date		

REQUIRED: PLEASE INCLUDE ALL REQUIRED LABS AND A COPY OF PATIENT'S INSURANCE CARD - FRONT AND BACK