

(Patient Sticker)

PHYSICIAN ORDER SET:

BELATACEPT CONVERSION (GREATER THAN 6 MONTHS FROM TRANSPLANT)

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Patier	nt:	DOB:		Gender:	
Patier	nt Phone #:	Height:		Weight:	
Diagn	osis:	ICD-10 Code: _			
Treati	ment Start Date:				
^o rovio	der Facility Name:	Provider Facility	y Address:		
Order	ing Provider:	Date:	:		
Siana	ture:				
Crite	ria to Treat Criteria to Treat		Interval	Defer Until	Duration
	If patient's current weight has changed more than 10% from their weight contact provider to discuss adjusting dose	Day 0 dosing	Every visit		
Com	munication Orders				
			Interval	Defer Until	Duration
	Provider Communication		Every visit		Every visit
	Routine, Until discontinued, Starting when released, Until Specific Risk of Post-Transplant Lymphoproliferative Disorder. Use in EBV seronegative or with serostatus.		s only. Do not use	e in tx recipients w	
ledic					ho are EBV
	ations				ho are EBV
	ations		Interval	Defer Until	ho are EBV Duration
	Provider and Nurse Communication		Interval 1 time per week	Defer Until	



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		Weight Type Used to Calculate Dose (Please Circle One):	1 time p week	er	1 time per week	
		Weight Type: Recorded Weight				
		belatacept (NULOJIX) 10 mg/kg in sodium chloride 0.9% 100 m IVPB 10 mg/kg, Intravenous, Administer over 30 Minutes, Once, Starting at treatment start			1 treatment	
		time, For 1 dose DAY 0. Use 0.2 – 1.2 micron filter when infusing this preparation.				
		belatacept (NULOJIX) 5 mg/kg in sodium chloride 0.9% 100 mL IVPB	Every 2 weeks		2 treatments	
		5 mg/kg, Intravenous, Administer over 30 Minutes, Once, Starting at treatment start time, For 1 dose DAY 14, DAY 28, DAY 42, DAY 56 Use 0.2 – 1.2 micron filter when infusing this preparation.				
		belatacept (NULOJIX) 5 mg/kg in sodium chloride 0.9% 100 mL IVPB	Every 4	at	Until discontinued	
			least 25			
		5 mg/kg, Intravenous, Administer over 30 Minutes, Once, Starting at treatment start time, For 1 dose DAY 84 and every 4 weeks.	days ap	ап		
		Use 0.2 – 1.2 micron filter when infusing this preparation.				
_ab	s					
			Interval	Defer Until	Duration	
П	Dr	ovider and Nurse Communication				
	Ro	utine, As needed, Starting when released, Until Specified e provider may have ordered HLA labs to be drawn. Check Order Review and release	e and draw HLA	labs, if ordered.		
Cat	het	er Management				
			Interval	Defer Until	Duration	
		ne Access	PRN		PRN	
		utine, Once Starting when released, Until Specified needed. Until Specified. Insert peripheral IV, or access peripheral, or central venous and the start of the s	access device,	to provide treatme	nt	
	alt	teplace (CATHFLO) 1mg/mL injection 2 mg	PRN		PRN	
	2 n	ng, Intracatheter, As needed, line care, Starting when released. For central venous cess device requiring clearance. May repeat once per lumen.				
	Fo	r patients greater than 30 kg: 2 mg in 2 mL using a 10 mL syringe. r patient less than or equal to 30 kg: instill a volume equal to 110% of the internal nen of CVAD, not to exceed 2 mg in 2 mL.				
		docaine-prilocaine (EMLA) cream	PRN		PRN	



saturation during Activity/Titration: Yes

Min Sp02 (%): 94

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Topical, As needed, apply prior to the PIV insertion or port access, Starting when released. For TOPICAL Use only. Allow at least 1 hour (mild dermal procedures) or at least 2 hours (major dermal procedures) for optimum therapeutic effect.

	heparin (PF) 100 unit/mL flush 5 mL 5 mL, Intravenous, As needed, line care, Starting when released	PRN		PRN
		551		
	heparin (PF) 10 unit/mL flush 3 mL 3 mL, Intravenous, As needed, line care, Starting when released	PRN		PRN
		551		2011
	heparin (PF) 10 unit/mL flush 5 mL	PRN		PRN
	5 mL, Intravenous, As needed, line care, Starting when released			
	heparin (PF) 1000 unit/mL catheter injection 2 mL	PRN		PRN
	2 mL, Intracatheter, As needed, APHERESIS LINE CARE ONLY. HEPARIN MUST BE			
	WITHRAWN FROM EACH LUMEN PRIOR TO FLUSHING OR INFUSING THROUGH THE APHERESIS CATHETER, Starting when released			
	sodium chloride (NS) 0.9% syringe flush 3 mL	PRN		PRN
_	3 mL, Intravenous, As needed, line care, Starting when released.			
	sodium chloride (NS) 0.9% syringe flush 10 mL	PRN		PRN
	10 mL, Intravenous, As needed, line care, Starting when released.	LKIN		LLIN
		DDM		DDM
	sodium chloride (NS) 0.9% syringe flush 20 mL	PRN		PRN
	20 mL, Intravenous, As needed, line care, Starting when released.			
	sodium chloride 0.9% infusion	PRN		PRN
	20 mL/hr, Intravenous, Continuous PRN, Keep vein open to provide treatment, Starting	l		
	when released, For 24 hours	DDN		DDN
	D5W infusion	PRN		PRN
	20 mL/hr, Intravenous, Continuous PRN, Keep vein open to provide treatment Starting when released, For 24 hours			
Eme	ergency Medications/Anaphylaxis			
		Interval	Defer Until	Duration
	Provider and Nurse Communication	PRN		PRN
	Routine, Once, Starting when released.			
	Treatment of SEVERE reaction (ANAPHYLAXIS): hypotension, throat swelling, wheezing			
	decreased oxygen saturation. Stop the infusion and treat with epinephrine FIRST. Notify personnel, administer oxygen as needed, monitor vital signs and proceed with administer		emergency	
	HYPERSENSITIVITY medications as clinically indicated.			
	EPINEPHrine (ADRENALIN) injection 0.3 mg	PRN		PRN
	0.3 mg, Intramuscular, As needed, anaphylaxis, Administer FIRST for anaphylaxis.			
	May repeat times 1 dose, Starting when released For 2 doses. Pharmacy's Suggested Dose Instructions; Epinephrine 1:1000 equivalent to 1 mg/mL			
	sodium chloride 0.9% bolus 1,000 mL	PRN		PRN
ш	1,000 mL, Intravenous, Once as needed, Hypotension, Starting when released, For 1	1 1314		1 1314
	dose			
	Oxygen Therapy - Non-Rebreather	PRN		PRN
	Routine			
	Select a Mode of Therapy: Non-Rebreather Titrate Ovugen and use the most appropriate device to maintain Target Ovugen			
	LITERTE LIXVIDED RIDITUSE THE MOST REPORTATE DEVICE TO MRIDIAIN LARGET DAVIGED			



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Hyp	persensitivity			
		Interval	Defer Until	Duration
	Provider and Nurse Communication	PRN		PRN
	Routine, Until discontinued, Starting when released, Until Specified Treatment for mild-moderate infusion reaction: Stop the infusion, notify provider and en monitor vital signs and proceed with administering medication as clinically indicated. If Medications section.			
	albuterol (ACCUNEB) 1.25 mg/3 mL nebulizer solution 2.5 mg	PRN		PRN
	2.5 mg, Nebulization, Once as needed, wheezing, shortness of breath, Starting when released, For 1 dose			
	acetaminophen (TYLENOL) tablet 975 mg	PRN		PRN
	975 mg, Oral, Once as needed, fever, Starting at treatment start time, For 1 dose			
	diphenhydrAMINE (BENADRYL) injection 25 mg	PRN		PRN
	25 mg, Intravenous, As needed, itching hives or adjunct treatment for mild-moderate, or SEVERE reaction, Starting when released, For 2 doses Begin with 25 mg. If patient has continued reaction, administer additional 25 mg.			
	methylprednisolone sodium succinate (PF) (SOLU-Medrol)	PRN		PRN
	 injection 40 mg 40 mg, Intravenous, Once as needed, Adjunct treatment for mild-moderate, or SEVERE reaction, Starting when released, For 1 dose. To be administered along with H1 antihistamine and methylprednisolone. 			
	famotidine (PEPCID) 20 mg in sodium chloride 0.9% 100 mL IVPB	PRN		PRN
	20 mg, Intravenous, Administer over 15 Minutes, at 400 mL/hr, Once as needed, Adjunct treatment for mild-moderate, or SEVERE reaction, Starting when released, For 1 dose. To be administered along with H1 antihistamine and methylprednisolone. HOLD IF: given as premed.			
	ondansetron (PF) (ZOFRAN) injection 4 mg	PRN		PRN
	4 mg, Intravenous, As needed, nausea, vomiting, may repeat x 1 dose, Starting when released, For 2 doses			
	cetirizine (ZyrTEC) tablet 10 mg	PRN		PRN
	10 mg, Oral, Once as needed, Adjunct treatment for mild-moderate, or SEVERE reaction, Starting at treatment start time, For 1 dose. If patient unable to tolerate cetirizine, administer fexofenadine if available. HOLD IF: given fexofenadine.			
	fexofenadine (ALLEGRA) tablet 180 mg	PRN		PRN
	180 mg, Oral, Once as needed, Adjunct treatment for mild-moderate, or SEVERE			

tolerate cetirizine. HOLD IF: given cetirizine.