

PHYO

CHST RITUXIMAB (RITUXAN or BIOSIMILAR)

	Page 1 of 5
Patient Name:	
D ((D: 1)	
Date of Birth:	

Baseline Patient	Demographic

CMC85948-002NS Rev. 11/2021	INFUSION THERAPY	PLAN (SOTP)				
Baseline Patient Demographic						
To be completed by the ordering provider.						
Diagnosis:	Height:	cm W	eight:	kg	Body Surface Area:	(m ²)
☐ NKDA - No Known Drug Allergies	☐ Allergies:					
Therapy Plan orders extend over time (sev		curring treatm	ent.			
Please specify the following regarding the ent	tire course of therapy:					
Duration of treatment: we	eeks mo	onths	unknow	/n		
Treatment should begin: as soon as	possible (within a week)	withi	n the month			
**Plans must be reviewed / re-ordered at le	east annually. **					
ORDERS TO BE COMPLETED FOR EACH	THERAPY					
ADMIT ORDERS						
✓ Height and weight						
✓ Vital signs						
Hypotension Defined Admit						
☑ Nursing communication						
Prior to starting infusion, please determi	ne the patient's threshold	d for hypotensi	on as defined by	the follo	owing parameters. This i	nformation will be
needed in the event of an infusion reaction	on occurring.		-			
Hypotension is defined as follows: 1 month to 1 year - systolic blood pressu	re (SRP) less than 70					
1 year to 11 years - systolic blood pressu		(2 x age in year	s)			
11 years to 17 years - systolic blood pres	ssure (SBP) less than 90		,			
OR any age - systolic blood pressure (SE Baseline systolic blood pressure (SBP) x						
PREGNANCY TESTS AT DALLAS AND P		as Hypotensi	л.			
Nursing communication						
Only one pregnancy test is necessary, b	ased on facility capabiliti	es. Please utiliz	e the lab that is	available	per facility.	
☐ Patient requires a pregnancy test (based	on organizational policy,	, female patient	s 10 years of age	e or over	require a pregnancy tes	t)
Pregnancy test, urine - POC STAT, ONE TIME, for females ≥ 10 year	rs old. If positive do NOT	infuse and cor	ntact the ordering	n provide	r	
Gonadotropin chorionic (HCG) urine	ola. Il positivo, do Ivo I	inidoc dila coi	naor are ordering	g provide		
STAT, ONE TIME, unit collect, for female	es <u>></u> 10 years old. If positi	ive, do NOT inf	use and contact	ordering	provider.	
Gonadotropin chorionic (HCG) quantit						
STAT, ONE TIME, unit collect, for female	es <u>></u> 10 years old. If positi	ive, do NOT inf	use and contact of	ordering	provider.	
INTRA PROCEDURE						
INTRA-PROCEDURE						
Please select all appropriate therapy						
IV START NURSING ORDERS						
Insert peripheral IV / Access IVAD	vailahle					

☐ lidocaine 1% BUFFERED (J-TIP LIDOCAINE)

0.2 mL, INTRADERMAL, PRN

when procedure will take about 1 minute patient / family preference for procedure when immediate procedure needed

Administration Instructions: NOTE: Do not use this medication in patients with bleeding disorders, platelets ≤ 20,000, or in patients taking anticoagulants, when accessing implanted ports or using a vein that will be utilized for chemotherapy administration, nor for pre-term infants or neonates.

☐ lidocaine - prilocaine (EMLA) cream

TOPICAL, PRN

when more than 60 minutes are available before procedure will take more than 1 hour

patient / family preference for procedure

Administration Instructions: NOTE: In children < 3 months of age, or < 5 kg in weight, maximum application time is 1 hour.



CMC85948-002NS	Rev. 11/2021

NURSING ORDERS, CONTINUED

	Page 2 of 5
Patient Name:	
Date of Birth:	

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INFUSION THERAPY PLAN (SOTP)

	lidocaine - tetracaine (SYNERA) patch TOPICAL, PRN ☐ when 20 - 30 minutes are available before pro ☐ when anticipated pain is less than 5 mm from lidocaine with transparent dressing 4% kit TOPICAL, PRN ☐ when 20 - 30 minutes are available before pro ☐ patient / family preference for procedure Heparin flush	n skin surface	dure will take more than 1 hour / family preference for procedure dure will take more than 1 hour	
	heparin flush 10 - 50 units, INTRAVENOUS, PRN, IV line flus used with all central lines including IVADs, with t		• •	s. This heparin flush should be
	heparin flush 100 - 300 units, INTRAVENOUS, PRN, IV line de-accessing IVADs.	e flush. Per protocol, hepa	arin should not be used to flush perip	oheral IVs. For use only when
	Sodium chloride flush			
	Sodium chloride flush 0.9% injection 1 - 20 mL, INTRAVENOUS, PRN, IV line flush			
	Sodium chloride - preservative free 0.9% inje 1 - 30 mL, INTRAVENOUS, PRN, IV line flush	ction		
PR	E-PROCEDURE LABS			
V	Completed Blood Count with differential Unit collect	INTERVAL: Every visit		DURATION: Until discontinued
√	Renal Function Panel Unit collect	INTERVAL: Every visit		DURATION: Until discontinued
V	Hepatic Function Panel Unit collect	INTERVAL: Every visit		DURATION: Until discontinued
	Gamma Glutamyl Transferase Unit collect	INTERVAL: Every visit		DURATION: Until discontinued
V	Magnesium Unit collect	INTERVAL: Every visit		DURATION: Until discontinued
V	Phosphorus Unit collect	INTERVAL: Every visit		DURATION: Until discontinued
	CMV Quantitative PCR Unit collect	INTERVAL: Once	DEFER UNTIL:	DURATION: Until discontinued
	Epstein Barr Virus Quantitative PCR Unit collect	INTERVAL: Once	DEFER UNTIL:	DURATION: Until discontinued
	BK Virus DNA PCR Quantitative Unit collect	INTERVAL: Once	DEFER UNTIL:	DURATION: Until discontinued
	Tacrolimus Unit collect needs to be drawn PRIOR to morning dose	INTERVAL: Once	DEFER UNTIL:	DURATION: Until discontinued
	Cyclosporine Random Unit collect needs to be drawn PRIOR to morning dose	INTERVAL: Once	DEFER UNTIL:	DURATION: Until discontinued
	Sirolimus Unit collect needs to be drawn PRIOR to morning dose	INTERVAL: Once	DEFER UNTIL:	DURATION: Until discontinued



PHYO CMC85948-002NS Rev. 11/2021

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Patient Name:	
Date of Birth:	

Maximum rate: 100 mg / hr

Maximum rate: 400 mg / hr

CHST RITUXIMAB (RITUXAN or BIOSIMILAR) INFUSION THERAPY PLAN (SÓTP)

RDFRS TO	RE COMPI	ETED FOR	EACH	THEDADY

PRE-MEDICATION	ons							
Nursing com	Acetaminophen pre-medication 30 minutes prior (15 mg / kg, maximum 650 mg) Nursing communication Administer only one of the acetaminophen orders, suspension or tablets, do not give both.							
15 mg / kg, O	acetaminophen suspension 15 mg / kg, ORAL, for 1 dose pre-medication, give 30 minutes prior to infusion Dose:							
acetaminoph 15 mg / kg Of Dose:	RAL, for 1 dose pre-medication, give	e 30 minutes prior to infusion						
Nursing com	nmunication only one of the diphenhydrAMINE pro	e-medication orders, liquid, capsule or injection, do no	ot give more than one of the orders as	;				
diphenhydr <i>l</i> 1 mg / kg, OR Dose:	RAL, for 1 dose pre-medication, give	e 30 minutes prior to infusion						
1 mg / kg OR	diphenhydrAMINE capsule 1 mg / kg ORAL, for 1 dose pre-medication, give 30 minutes prior to infusion Dose:							
1 mg / kg, IN	MINE injection TRAVENOUS, 1 dose pre-medication	on, give 30 minutes prior to infusion						
methylPREDNISolone RTA infusion 2 mg / kg INTRAVENOUS, for 1 dose. Give 30 minutes prior to riTUXimab. Ready to administer by infusion. Doses > 15 mg / kg should be given over a minimum of 1 hour. (see protocol for monitoring parameters.) Dose:								
INTRA-PROCEDURE								
Nursing com Adverse reac if these condi	tions may include fever, chills, rigor	rs, hypotension and severe allergic reactions (anaphy	laxis). Please contact the ordering pro	ovider				
Vital signs Basline vitals after infusion		then monitor vitals every 15 minutes during after initia	ation of the infusion and for 30 minutes	s				
In the event of		usion related symptoms, the infusion should be stoppe	ed. Please contact the ordering provid	ler.				
Initial infusion	First hour infusion rate	0.5 mg / kg / hr	Maximum rate: 50 mg / hr					
	If no infusion related events observed after 60 minutes	Increase rate by 1 mg / kg / hr	Maximum rate: 100 mg / hr					
	If no infusion related events observed after 30 minutes	Continue to increase rate by 0.5 mg / kg / hr every 30 minutes until maximum rate has been reached	Maximum rate: 400 mg / hr					
Subsequent	First hour infusion rate	1 mg/kg/hr	Maximum rate: 100 mg / hr					

Increase rate by 1 mg / kg / hr every 30 minutes

until maximum rate has been reached

1 mg / kg / hr

First hour infusion rate

If no infusion related events

are observed after 30 minutes

infusion



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PHYO CMC85948-002NS Rev. 11/2021

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ORDERS TO BE COMPLETED FOR EACH THERAPY	

	52.K0 10 52 00	22 : 23 : 61(2)(61: 1112:01: 1							
INT	RA-PROCEDUR	E, CONTINUED							
		riTUXimab dose, if clinically accept	table, to nearest 100 mg to minimize waste. Dose reg treatment). Please enter the dose of riTUXimab in						
(ima	ab (RITUXAN or I	biosimilar) - 1 time a week (Sele	ct one product below):						
		e, starting 0.5 hours after treatment on should be 1 mg / mL	INTERVAL: 1 time a week start time	DURATION: For 4 treatments					
☐ Sodium Chloride 0.9% ☐ Dextrose (Dilute to a final concentration of 1 - 4 mg / mL). Dose: mL.									
	Initial infusion	First hour infusion rate	0.5 mg / kg / hr	Maximum rate: 50 mg / hr					
		If no infusion related events observed after 60 minutes	Increase rate by 1 mg / kg / hr	Maximum rate: 100 mg / hr					
		If no infusion related events observed after 30 minutes	Continue to increase rate by 0.5 mg / kg / hr every 30 minutes until maximum rate has been reached	Maximum rate: 400 mg / hr					
	Subsequent	First hour infusion rate	1 mg / kg / hr	Maximum rate: 100 mg / hr					
	infusion	If no infusion related events are observed after 30 minutes	Increase rate by 1 mg / kg / hr every 30 minutes until maximum rate has been reached	Maximum rate: 400 mg / hr					
	Intravenous, once Final concentration Dose:		INTERVAL: 1 time a week start time te to a final concentration of 1 - 4 mg / mL). Dos	DURATION: For 4 treatments e: mL.					
	Initial infusion	First hour infusion rate	0.5 mg / kg / hr	Maximum rate: 50 mg / hr					
		If no infusion related events observed after 60 minutes	Increase rate by 1 mg / kg / hr	Maximum rate: 100 mg / hr					
		If no infusion related events observed after 30 minutes	Continue to increase rate by 0.5 mg / kg / hr every 30 minutes until maximum rate has been reached	Maximum rate: 400 mg / hr					
	Subsequent	First hour infusion rate	1 mg / kg / hr	Maximum rate: 100 mg / hr					
	infusion	If no infusion related events are observed after 30 minutes	Increase rate by 1 mg / kg / hr every 30 minutes until maximum rate has been reached	Maximum rate: 400 mg / hr					
d	Expires in 365 da	epartment for the therapy appoin	_	☐ Dallas Neurology					
ЕМ	ERGENCY MEDI	CATIONS							
	Nursing commu								
	1 Hives or cu	itaneous reaction only - no other sy	stem involvement						

PATIENT IS HAVING A DRUG REACTION:

- a. Stop the infusion
- **b.** Give diphenhydramine as ordered
- c. Check heart rate, respiratory rate and blood pressure every 5 minutes until further orders from provider.
- d. Connect patient to monitor (cardiac / apnea, blood pressure and oxygen saturation) if not already on one
- e. Notify provider for further orders



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ORDERS TO BE	COMPLETED FOR EACH	I THERAPY	•					
EMERGENCY MI	EDICATIONS							
✓ Nursing com	munication							
respirat	or cutaneous reaction pl cory distress, mouth / tongo NT IS HAVING ANAPHYL		ominal cramping, v	omiting, hypotension	n, altered mental status,			
b. C; c. G; d. N; e. C; f. C; g. G; h. M; te i. M; se H; 1 1	onnect patient to monitor (ve diphenhydramine once ay repeat epinephrine eve am arrives. ay give albuterol as order tutration. ypotension is Defined as month to 1 year – systolic year to 11 years – systolic l years to 17 years – systolic R any age – systolic blood	rate and blood pressure every 5 cardiac / apnea, blood pressure a as needed for hives ery 5 minutes x 2 doses for pers ed for wheezing with oxygen sa	and oxygen saturation sistent hypotension turation stable while 70 70 + (2 x age in year in 90 30% from baseline.	on), if not already on and respiratory disti e waiting for code te	ress with desaturation until code			
(AMPULE / E 0.01 mg / kg, distress with o	lesaturation until the code	0.01 mg / kg RY 5 MINUTES PRN, for anaph			ent hypotension and respiratory			
Rationale for Clir Clir Rec	nically significant cardiac a cent acute life-threatening explained or acutely abnor ficial airway (stent, trache ute, fluctuating or consisten	event mal vital signs ostomy, oral airway)	turation □ Respir	atory rate				
diphenhydrA 1 mg / kg, INT Dose:	-	for hives or cutaneous reaction,	for 1 dose. Maximur	m dose = 50 mg per	dose, 300 mg per day.			
Albuterol for 0.25 mg / kg. saturation for Dose:	, INHALATION ONCE PF 1 dose	RN, for wheezing, but oxygen sa	turations stable whil	e waiting for code te	eam, continue to monitor oxyger	1		
POST- PROCED	URE							
Nursing communication Flush PIV or IVAD with 20 mL 0.9% sodium chloride (250 mL bag) at the completion of the infusion. Flush IVAD with saline and heparin flush per protocol prior to de-accessing IVAD. Discontinue PIV prior to discharge.								
INTRAVENOL	ride 0.9% infusion JS at 0 - 25 mL / hr, ONCE	E, for 1 dose.						
Dose:	mL		(circle one):					
Signature of Prov	ider		<u>MD</u> <u>DÓ</u> Credentials	Date	 Time			
-								