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# Physician's Orders RITUXIMAB OR BIOSIMILAR WEEKLY TIMES 4 DOSES - ADULT, OUTPATIENT, INFUSION CENTER

Patient Name
DOB
MRN
Physician
CSN

Anticipated Infusion Date	ICD 10 Code with D	escription	
Height(cm)	Weight(kg) Allergies	S	<del> </del>
Site of Service			
☐ CH Gerber	☐ CH Lemmen Holton (GR)	□ CH Pennock	□ CH Greenville
☐ CH Helen DeVos (GR)	☐ CH Ludington	□ CH Reed City	☐ CH Zeeland
☐ CH Blodgett (GR)			
Provider Specialty			
☐ Allergy/Immunology	☐ Infectious Disease	□ OB/GYN	☐ Rheumatology
□ Cardiology	☐ Internal Med/Family Practice	□ Other	☐ Surgery
☐ Gastroenterology	□ Nephrology	☐ Otolaryngology	☐ Urology
☐ Genetics	☐ Neurology	□ Pulmonary	☐ Wound Care
Treatment Intent			
□ Conditioning	☐ Curative	☐ Mobilization	☐ Supportive
☐ Control	☐ Maintenance	□ Palliative	

#### Cycle 1

One dose every 7 days x 4 doses on day 1, 8, 15, 22

Provider Reminder

**ONC PROVIDER REMINDER 2** 

Confirm that the appropriate informed consents have been signed and are in the medical record.

Appointment Requests

ONCBCN CALCULATED LENGTH INFUSION APPOINTMENT REQUEST 1

Labs

COMPLETE BLOOD COUNT (CBC) W/DIFFERENTIAL

**■** Pregnancy tests recommended for Females aged 12 to 60 with Uterus intact.

Please order as appropriate for clinical presentation

Interval: Once Selection conditions: Patient could become pregnant

**HCG, QUANTITATIVE** 

**■ ONC PROVIDER REMINDER** 

Review prior hepatitis labs and consider repeat screening as indicated.

**HEPATITIS PANEL** 

Selection conditions: Hides NCCN Hepatitis Screening orders in background if the patient has had a completed hepatitis lab within the last 365 days.

**■ HEP B CORE TOTAL AB** 

Selection conditions: Hides NCCN Hepatitis Screening orders in background if the patient has had a completed hepatitis lab within the last 365 days.

Vitals

VITAL SIGNS

RITUXIMAB - Dose 1 & 2: Vital signs every 15 minutes x 2 hours, then hourly throughout infusion. Subsequent doses: Vital signs with rate changes.

**Nursing Orders** 

- HYPERSENSITIVITY REACTION ADULT ONCOLOGY PROTOCOL
- ONC NURSING COMMUNICATION 100

May Initiate IV Catheter Patency Adult Protocol





## **RITUXIMAB OR BIOSIMILAR WEEKLY TIMES 4 DOSES -**

Patient Name DOB MRN Physician CSN

ADULT, OUTPATIENT, INFUSION CENTER	(CONTINUED)
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Treat	ment Parameters						
	ONC MONITORING AND HOLD PARAMETERS 3						
	May proceed with treatment if hepatitis B core antibody and surface antigen labs have been						
	resulted prior to the first dose, and the results are negative.						
Pre-N	-Medications						
	acetaminophen (Tylenol) tablet 650 mg						
	Dose: 650 mg Route: Oral Once Instructions: Administer 30 minutes prior to rituximab.						
	oliphenhydrAMINE (Benadryl) capsule 50 mg						
	Dose: 50 mg Route: Oral Once Instructions: Administer 30 minutes prior to rituximab.						
Pre-N	Medications						
	ONC PROVIDER REMINDER 7						
	HISTORY OF INFUSION REACTION: consider adding CORTICOSTEROID pre-medication prior						
	to riTUXimab or biosimilar						
	OdexAMETHasone (Decadron) injection 10 mg						
	Dose: 10 mg Route: Intravenous Once over 5 Minutes for 1 dose						
	Instructions:						
	For patient with a history of reaction to riTUXimab or biosimilar. Administer 30 minutes prior to start of riTUXimab or biosimilar.						
	_						
	methylPREDNISolone sodium succinate (SOLU-Medrol) injection 125 mg						
	Dose: 125 mg Route: Intravenous Once for 1 dose Instructions:						
	For patient with a history of reaction to riTUXimab or biosimilar. Administer 30 minutes prior to						
	start of riTUXimab or biosimilar.						
	hydrocortisone sodium succinate (PF) (Solu-CORTEF) injection 100 mg						
	Dose: 100 mg Route: Intravenous Once for 1 dose						
	Instructions:						
	For patient with a history of reaction to riTUXimab or biosimilar. Administer 30 minutes prior to start of riTUXimab or biosimilar.						
Mono	start of ITTOXIMAD of biosimilar.						
VIOLIC							
	riTUXimab-pvvr (RUXIENCE) 2 mg/mL infusion 375 mg/m2						
	Dose: 375 mg/m2 Route: Intravenous Titrate @ 25-200 mL/hr for 1 dose Instructions:						
	Before start of riTUXimab-pvvr, infusion, prime 0.9% sodium chloride 250 mL with primary						
	tubing and have available at bedside in case of hypersensitivity or infusion reaction. Do NOT						
	connect this saline with any other intravenous drugs, fluids, or blood products.						
	Start infusion at 50 mg/hr (25 mL/hr). For patients tolerating the infusion well, the rate of						
	infusion may be increased in increments of 50 mg/hr (25 mL/hr) at intervals of 30 minutes to a						
	maximum rate of 400 mg/hr (200 mL/hr).						
	SUBSEQUENT INFUSIONS: If patient tolerated initial infusion, start at 100 mg/hour (50 mL/hr); if there is no infusion related reaction, ingresses the rate by 100 mg/hour (50 mL/hr) ingresses the rate by 100 mL/hr) ingresses the rate by 100 mL/hr) ingresses the rate by 100 mL/hr) ingresses t						
	if there is no infusion-related reaction, increase the rate by 100 mg/hour (50 mL/hr) increments every 30 minutes, to a maximum rate of 400 mg/hour (200 mL/hr)						
	riTUXimab-abbs (TRUXIMA) 2 mg/mL infusion 375 mg/m2						
	Dose: 375 mg/m2 Route: Intravenous Titrate @ 25-200 mL/hr for 1 dose						
	Instructions:						
	Before start of riTUXimab-abbs, infusion, prime 0.9% sodium chloride 250 mL with primary						
	tubing and have available at bedside in case of hypersensitivity or infusion reaction. Do NOT						
	connect this saline with any other intravenous drugs, fluids, or blood products.						
	Start infusion at 50 mg/hr (25 mL/hr). For patients tolerating the infusion well, the rate of						





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infusion may be increased in increments of 50 mg/hr (25 mL/hr) at intervals of 30 minutes to a maximum rate of 400 mg/hr (200 mL/hr).

SUBSEQUENT INFUSIONS: If patient tolerated initial infusion, start at 100 mg/hour (50 mL/hr); if there is no infusion-related reaction, increase the rate by 100 mg/hour (50 mL/hr) increments every 30 minutes, to a maximum rate of 400 mg/hour (200 mL/hr)

#### riTUXimab (RITUXAN) 2 mg/mL infusion 375 mg/m2

Dose: 375 mg/m2 Route: Intravenous Titrate @ 25-200 mL/hr for 1 dose Instructions:

Before start of riTUXimab, infusion, prime 0.9% sodium chloride 250 mL with primary tubing and have available at bedside in case of hypersensitivity or infusion reaction. Do NOT connect this saline with any other intravenous drugs, fluids, or blood products.

Start infusion at 50 mg/hr (25 mL/hr). For patients tolerating the infusion well, the rate of infusion may be increased in increments of 50 mg/hr (25 mL/hr) at intervals of 30 minutes to a maximum rate of 400 mg/hr (200 mL/hr).

SUBSEQUENT INFUSIONS: If patient tolerated initial infusion, start at 100 mg/hour (50 mL/hr); if there is no infusion-related reaction, increase the rate by 100 mg/hour (50 mL/hr) increments every 30 minutes, to a maximum rate of 400 mg/hour (200 mL/hr)

#### niTUXimab-arrx (RIABNI) 2 mg/mL infusion 375 mg/m2

Dose: 375 mg/m2 Route: Intravenous Titrate @ 25-200 mL/hr for 1 dose Instructions:

Before start of riTUXimab-arrx infusion, prime 0.9% sodium chloride 250 mL with primary tubing and have available at bedside in case of hypersensitivity or infusion reaction. Do NOT connect this saline with any other intravenous drugs, fluids, or blood products.

Start infusion at 50 mg/hr (25 mL/hr). For patients tolerating the infusion well, the rate of infusion may be increased in increments of 50 mg/hr (25 mL/hr) at intervals of 30 minutes to a maximum rate of 400 mg/hr (200 mL/hr).

SUBSEQUENT INFUSIONS: If patient tolerated initial infusion, start at 100 mg/hour (50 mL/hr); if there is no infusion-related reaction, increase the rate by 100 mg/hour (50 mL/hr) increments every 30 minutes, to a maximum rate of 400 mg/hour (200 mL/hr)

#### Supportive Care

#### prochlorperazine (COMPAZINE) tablet 10 mg

Dose: 10 mg Route: Oral Every 6 hours PRN

Instructions: Use 1st line for nausea/vomiting for patients able to tolerate oral medications.

#### prochlorperazine (COMPAZINE) injection 10 mg

Dose: 10 mg Route: Intravenous Every 6 hours PRN over 5 Minutes Instructions: Use 1st line for nausea/vomiting for patients unable to tolerate oral medications.

#### Ondansetron (ZOFRAN-ODT) disintegrating tablet 8 mg

Dose: 8 mg Route: Oral Every 8 hours PRN

Instructions: Use 2nd line for Nausea/Vomiting. May give IV if unable to tolerate PO.

#### Ondansetron (ZOFRAN) injection 8 mg

Route: Intravenous Every 8 hours PRN over 5 Minutes Instructions: Use 2nd line for Nausea/Vomiting. May give IV if unable to tolerate PO.

Telephone order/Verbal order documented and read-back completed. Practitioner's initials.

NOTE: Unless Order is written DAW (dispense as written), medication may be supplied which is a generic equivalent by nonproprietary name.

TRANSCRIBE	:D:	VALIDATED	1	ORDERED:			
TIME	DATE	TIME	DATE	TIME	DATE	Pager#	
			R.N.		Phys	ician	Physician
	Sign		Sign			Print	Sign

EPIC VERSION DATE: 07/16/20