

Vancouver Cancer Centre

PROTOCOL CODE: LYHDMRTEM (Inpatient)

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DOCTOR'S ORDERS Wt kg BSA m² Ht<u>c</u>m ALLERGY/ALERT: Reminder to Physicians: Please ensure drug allergies and previous Bleomycin are documented on the Allergy and Alert Form. Date/Time: Cycle #: Admit to inpatient bed DAT, AAT, VSR, ECOG CBC & diff, creatinine, electrolytes panel, total bilirubin, ALT, alk. Phos, LDH, urine pH, CXR HBsAg, HBsAb and HBcore Ab if not previously done Calculate creatinine clearance (see protocol summary) Daily creatinine, electrolytes panel CBC & diff on day 4 If clinically indicated: HBV viral load ☐ If clinically indicated post methotrexate: daily ALT, total bilirubin, alkaline phosphatase, LDH, GGT At hour 48 (from start of methotrexate infusion) or morning of day 3, then daily q am: methotrexate levels (until level less than 0.1 micromol/L; note date and time of withdrawal as well as start time of infusion on specimen. MD to be notified of all results immediately Daily weights, intake / output Administer Folstein Mini Mental Status Exam (MMSE) at 1st treatment and at final treatment and record results in admission/discharge summary. Ocular slit lamp exam (ophthalmology consultation) START ALKALINISING REGIMEN 4 TO 12 HOURS PRIOR TO METHOTREXATE: Discontinue all other IV hydration before starting alkalinizing regimen. Start IV D5W with potassium chloride 20mEq/L and sodium bicarbonate 150 mEq/L at 125 mL/h for at least 4 hours prior to methotrexate until urine pH is greater than 7. Hydration may be temporarily held during methotrexate infusion (per physician/nursing discretion). Continue hydration post-methotrexate infusion until methotrexate level is less than 0.1 micromol/L. PREMEDICATIONS: ondansetron 8 mg PO/IV immediately before methotrexate infusion. prochlorperazine 10 mg PO once after methotrexate infusion completed. ondansetron 8 mg PO prior to temozolomide PRN: prochlorperazine 10 mg PO q4h prn thereafter. CHEMOTHERAPY: Check urine pH prior to starting methotrexate If urine pH less than 7, continue alkalinising regimen until pH greater than or equal to 7 If urine pH greater than or equal to 7: methotrexate 8 gram/m² x BSA = _____ gram IV in 1000 mL NS over 4 hours. Dose modification (____%)= ____gram/m² x BSA = ____gram IV in 1000 mL NS over 4 hours Measure urine pH q6h. If pH less than 7, notify MD 24 hours after start of methotrexate infusion begin leucovorin 25 mg IV g 6h x 4 doses, then leucovorin 25 mg PO q 6h until methotrexate level less than 0.1 micromol/L leucovorin dose may need to be adjusted upwards depending on methotrexate level. See protocol summary for details. POSTHYDRATION: See Alkalinizing Regimen above Note: One staff Physician signature is required. Orders written by other providers MUST be cosigned. Signatures UC: **Doctor 1 Signature: Doctor 2 Signature:** RN:



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Have Hypersensitivity Reaction Tray and Protocol Available

PREMEDICATIONS:

Date:

For intravenous infusion:

diphenhydrAMINE 50 mg PO prior to riTUXimab and then q4h during the IV infusion, if the infusion exceeds 4 hours **acetaminophen 650 mg -975 mg** PO prior to riTUXimab and then q4h during the IV infusion, if the infusion exceeds 4 hours

For subcutaneous injection: diphenhydrAMINE 50 mg PO prior to riTUXimab subcutaneous acetaminophen 650 mg - 975 mg PO prior to riTUXimab subcutaneous

TREATMENT: (Note: riTUXimab is given for a total of 4 doses)

riTUXimab subcutaneous or IV may be administered either before or after chemotherapy, but within 72 hours after methotrexate

mg

TREATMENT #1:

riTUXimab (first dose) 375 mg/m² x BSA = _____

IV in 250 to 500 mL NS. Start at 50 mg/hour.

After 1 hour, increase rate by 50 mg/hour every 30 minutes until rate = 400 mg/hour unless toxicity occurs. NOTE: If the peripheral blood lymphocyte count is above 30×10^9 /L, the riTUXimab should be omitted from that cycle. Patients are to be under constant visual observation during all dose increases and for 30 minutes after infusion completed. Vital signs are not required, unless symptomatic.

Pharmacy to select riTUXimab brand as per Provincial Systemic Therapy Policy III-190

Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date
riTUXimab		

FOR ALL SUBSEQUENT TREATMENTS:

□ Patient tolerated a full dose of IV riTUXimab (no severe reactions requiring early termination) and can proceed to subcutaneous riTUXimab:

riTUXimab subcut (RITUXAN SC) 1400 mg (fixed dose in 11.7 mL) subcutaneously into abdomen over 5 minutes. Observation for 15 minutes following administration.

NB. During treatment with subcutaneous riTUXimab, administer other subcutaneous drugs at alternative injection sites whenever possible.

Note: One staff Physician signature is required.	Orders written by other providers MUST be co-signed.	Signatures
		UC:
Doctor 1 Signature:	Doctor 2 Signature:	RN



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Date:

Treatment continued:

FOR ALL SUBSEQUENT TREATMENTS:

Patient did not tolerate a full dose of IV riTUXimab (experienced severe reactions requiring early termination) in the previous treatment and will continue with IV riTUXimab for this cycle:

riTUXimab (subsequent dose) 375 mg/m² x BSA = ____ mg

IV in 250 to 500 mL NS. Infuse 50 mL (or 100 mL of 500 mL bag) of the dose over 30 minutes, then infuse the remaining 200 mL (or 400 mL of 500 mL bag) over 1 hour (total infusion time = 1 hour, 30 minutes).

For all subsequent doses, constant visual observation is not required.

If flushing, dyspnea, rigors, rash, new pruritus, vomiting, chest pain or any other new acute discomfort occurs, stop infusion and page physician.

Pharmacy to select riTUXimab brand as per Provincial Systemic Therapy Policy III-190

Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date
riTUXimab		

ON ALTERNATE CYCLES:

May proceed with day 7 if ANC <u>greater than or equal to</u> 1.0 x 10 ⁹ /L, Platelets <u>greater than or equal to</u> 50 within 72 hours	x 10 ⁹ /L
temozolomide 150 mg/m ² or 100 mg/m ² (<i>select one</i>) x BSA =mg PO daily at bedtime days 7 to 11. (refer to <u>Temozolomide Suggested Capsule Combination Table</u> for dose rounding)	e x 5 days on
SUPPORTIVE MEDICATIONS: REMINDER: Write orders for Dexamethasone, Famotidine and Cotrimoxazole for inpatient use, if applicable.	
Check one: Readmit in 2wks for cycle	
Final Treatment: RTC in 2 months (reminder: administer MMSE at FU visits)	
TESTS:	
At cycle # 3, prebook for repeat 🗌 CT Brain or 🗌 MRI Brain immediately prior to cycle # 5.	
Note: One staff Physician signature is required. Orders written by other providers MUST be co-signed.	Signatures UC: RN:
Doctor 1 Signature: Doctor 2 Signature:	



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DOCTOR'S ORDERS	
REMINDER: Please ensure drug allergies and previous bleomycin are documented on the Allergy & Alert Form	
Date:To be given:Cycle #:	
Date of LYHDMRTEM Chemotherapy	
TREATMENT: leucovorin 25 mg PO q6h xdoses NOTE: leucovorin to be continued until methotrexate level is less than 0.1 micromoL/L.	
RN or Pharmacist to instruct patient on exact dosing times.	
leucovorin is a BC Cancer Benefit Drug – this prescription should be filled at a BC Cancer Outpatient Pharmacy.	
TESTS: Physician to order methotrexate Level in a.m. daily if needed	
Doctor's Signature:	Signature UC:



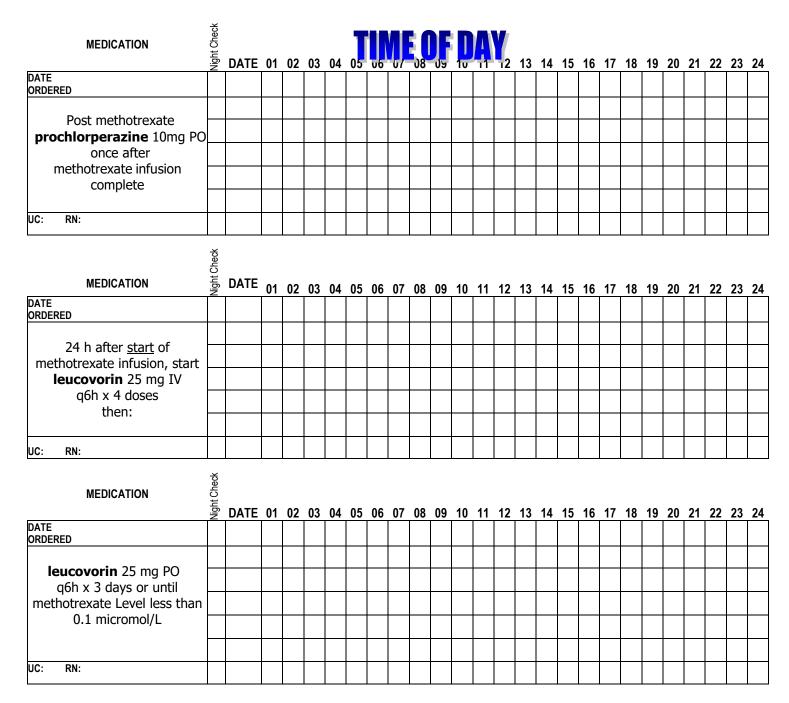
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UC: RN:																										
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UC: RN:																										
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methotrexate g IV in 1000 mL NS over 4 h																										
UC: RN:																										



MEDICATION ADMINISTRATION RECORD Protocol: LYHDMRTEM

DRUG ALLERGIES:

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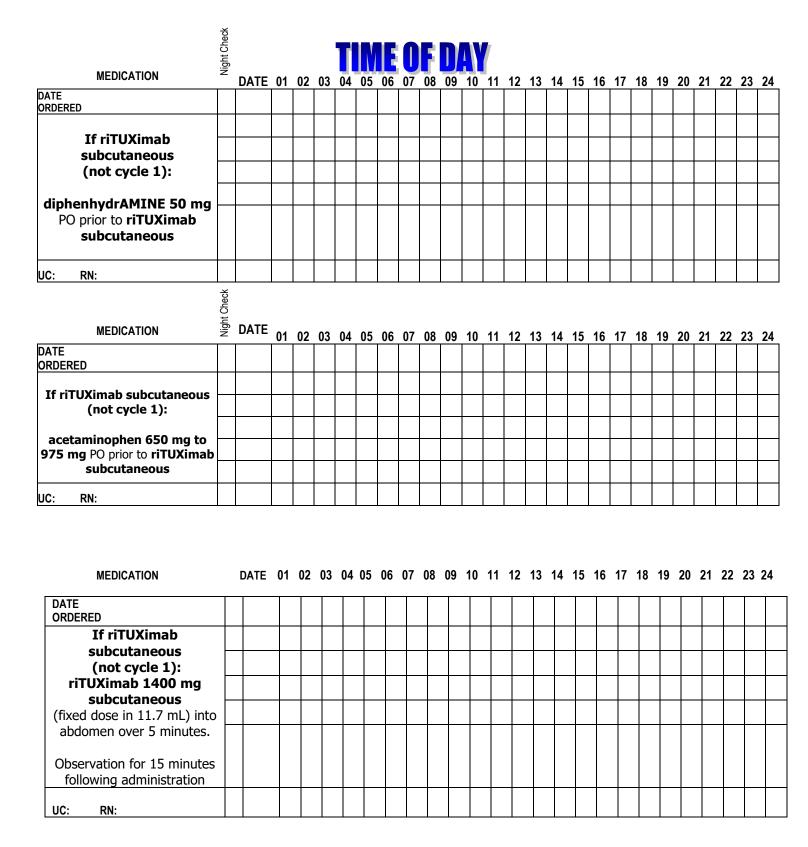


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If IV rituximab: diphenhydrAMINE 50 mg PO prior to riTUXimab, then q4h during the IV																										
infusion, if the infusion exceeds 4 hours																										
UC: RN:																										
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DATE ORDERED																										
If IV rituximab:																										
riTUXimab mg IV in																									┢──┦	
250-500 mL NS within 72																										
hours of methotrexate																										
See orders for rate of infusion																										
UC: RN:																										
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MEDICATION ADMINISTRATION RECORD Protocol: LYHDMRTEM DRUG ALLERGIES:

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MEDICATION ADMINISTRATION RECORD Protocol: LYHDMRTEM DRUG ALLERGIES:

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TIME OF DAY

MEDICATION	DA	ATE	01	02	03	04	05	06	07	08	09	10	11 1	2 1	13 ⁻	14	15	16	17	18	19	20	21	22	23	24
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30 minutes prior to temozolomide on days 7 to																										
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UC: RN:																										

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PO at HS on days 7 to 11																					-		_					
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DATE ORDERED																													
prochlorperazine 10 mg PO q4h PRN																													
10 mg PO q4h PRN																												-	
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H:\Everyone\Systemic\Chemo\Orders\VCC\Lymphoma&Myeloma\LYHDMRTEM_ippo_VCC_18Jan2016 Created: 1 July 2007 (as LYHDMRP) Revised: 1 Dec 2024 (Tests updated)