

**PROTOCOL CODE: HLHETCSPA**  
(Week 3 to 8) (Page 1 of 3)

<b>DOCTOR'S ORDERS</b>		Ht _____ cm	Wt _____ kg	BSA _____ m <sup>2</sup>
<b>REMINDER: Please ensure drug allergies are documented on the Allergy &amp; Alert Form</b>				
<b>DATE:</b>	<b>To be given:</b>	<b>Cycle #:</b>		
Date of Previous Cycle:				
<input type="checkbox"/> Delay treatment _____ week(s) <input type="checkbox"/> <b>CBC &amp; Diff and platelets, bilirubin, creatinine</b> day 1 of treatment				
Dose modification for: <input type="checkbox"/> Hematology <input type="checkbox"/> Other Toxicity _____				
Proceed with treatment based on blood work from _____				
<b>PREMEDICATIONS:</b> Patient to take own supply. RN/Pharmacist to confirm _____.				
<input type="checkbox"/> Other				
<b>** Have Hypersensitivity Reaction Tray and Protocol Available**</b>				
<b>TREATMENT: Week 3 to 8 ONLY:</b>				
<input type="checkbox"/> <b>cycloSPORINE 3 mg/kg</b> x Wt = _____ mg PO BID (round to the nearest 25 mg) Mitte: _____ capsules				
<input type="checkbox"/> week 3 and 4: <b>dexamethasone 5 mg/m<sup>2</sup></b> x BSA = _____ mg PO daily (round to the nearest 2 mg) Mitte: _____ tablets OR				
<input type="checkbox"/> week 5 and 6: <b>dexamethasone 2.5 mg/m<sup>2</sup></b> x BSA = _____ mg PO daily (round to the nearest 0.5 mg) Mitte: _____ tablets OR				
<input type="checkbox"/> week 7 and 8: <b>dexamethasone 1.25 mg/m<sup>2</sup></b> x BSA = _____ mg PO daily (round to the nearest 0.5 mg) Mitte: _____ tablets				
<b>etoposide 150 mg/m<sup>2</sup></b> x BSA = _____ mg <input type="checkbox"/> Dose Modification: _____ % = _____ mg/m <sup>2</sup> x BSA = _____ mg IV in 500 to 1000 mL (non-DEHP bag) NS over 45 minutes to 1 hour 30 minutes on <b>Days 15, 22, 29, 36, 43, 50</b> (Use non-DEHP tubing with 0.2 micron in-line filter)				
<b>EMERGENCY DRUGS FOR MANAGEMENT OF ETOPOSIDE TOXICITY:</b>				
hydrocortisone 100 mg IV prn / diphenhydrAMINE 50 mg IV prn				
See page 2				
<b>DOCTOR'S SIGNATURE</b>				<b>SIGNATURE</b>
				<b>UC:</b>



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<b>DOCTOR'S ORDERS</b>		Ht _____ cm	Wt _____ kg	BSA _____ m <sup>2</sup>
<b>REMINDER: Please ensure drug allergies and previous bleomycin are documented on the Allergy &amp; Alert Form</b>				
<b>DATE:</b>	<b>To be given:</b>	<b>Cycle #:</b>		
Date of Previous Cycle: _____				
<input type="checkbox"/> Delay treatment _____ week(s) <input type="checkbox"/> <b>CBC &amp; Diff and Platelets</b> prior to day 1, 8 and 15 of treatment May proceed with doses as written if within 24 hours <b>ANC greater than or equal to 0.5 x 10<sup>9</sup>/L, Platelets greater than or equal to 40 x 10<sup>9</sup>/L</b>				
Dose modification for: <input type="checkbox"/> Hematology <input type="checkbox"/> Other Toxicity _____				
Proceed with treatment based on blood work from _____				
<b>INTRATHECAL (IT) CHEMOTHERAPY:</b>				
methotrexate _____ mg (standard dose 12 mg) and hydrocortisone _____ mg (standard dose 50 mg) qs to 6 mL with preservative-free NS intrathecally on week _____.				
<b>RETURN APPOINTMENT ORDERS</b>				
<input type="checkbox"/> Return in 1 week for Doctor and week _____. <input type="checkbox"/> Last Cycle. Return in _____ week(s).				
<input type="checkbox"/> <b>Other tests:</b> <input type="checkbox"/> <b>Consults:</b> <input type="checkbox"/> See general orders sheet for additional requests.				
<b>DOCTOR'S SIGNATURE:</b>			<b>SIGNATURE:</b>	
			<b>UC:</b>	
<b>MEDICATION VERIFICATION CHECKS</b>				
Full Signatures Required				
<b>Date (dd/mm/yy)</b>				
methotrexate _____ mg and hydrocortisone _____ mg IT		(RN)		
		(MD)		