



Provincial Health Services Authority

Information on this form is a guide only. User will be solely responsible for verifying its currency and accuracy with the corresponding BC Cancer treatment protocols located at www.bccancer.bc.ca/terms-of-use and according to acceptable standards of care.

PROTOCOL CODE: BRAVTR

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DOCTOR'S ORDERS								
Ht _____ cm	Wt _____ kg	BSA _____ m ²						
REMINDER: Please ensure drug allergies and previous bleomycin are documented on the Allergy & Alert Form								
DATE:	To be given:	Cycle # of Trastuzumab:						
Date of Previous Cycle:								
Indicate the number of trastuzumab doses patient has received together with chemotherapy (not as single-agent) to date: _____								
Have Hypersensitivity Reaction Tray and Protocol Available								
TREATMENT:								
<input type="checkbox"/> Cycle 1 (NEW patients ONLY – Omit for patients continuing single-agent trastuzumab following a trastuzumab-containing chemotherapy regimen): trastuzumab 8 mg / kg x _____ kg = _____ mg IV in 250 mL NS over 1 hour 30 minutes. Observe for 1 hour post infusion*. Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190								
<table border="1" style="width: 100%; border-collapse: collapse;"><tr><th style="width: 20%;">Drug</th><th style="width: 50%;">Brand (Pharmacist to complete. Please print.)</th><th style="width: 30%;">Pharmacist Initial and Date</th></tr><tr><td>trastuzumab</td><td></td><td></td></tr></table>	Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date	trastuzumab				
Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date						
trastuzumab								
OR								
<input type="checkbox"/> Cycle 2 trastuzumab 6 mg/kg x _____ kg = _____ mg IV in 250 mL NS over 1 hour. Observe for 30 minutes post-infusion*. Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190								
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trastuzumab								
<input type="checkbox"/> Cycle 3 and Subsequent: (For patients who have just completed a trastuzumab-containing chemotherapy regimen) trastuzumab 6 mg/kg x _____ kg = _____ mg IV in 250 mL NS over 30 minutes** every three weeks x _____ cycle(s). Observe for 30 minutes post-infusion*. Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190								
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trastuzumab								
<small>* Observation period not required after 3 treatments with no reaction ** 30 minute infusion time for Cycle 3 and all subsequent cycles, if no previous adverse reactions.</small>								
acetaminophen 325 to 650 mg PO PRN for headache and rigors								
Proceed with treatment based on blood work from _____								
DOCTOR'S SIGNATURE:		SIGNATURE:						
		UC:						



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DOCTOR'S ORDERS		Ht _____ cm	Wt _____ kg	BSA _____ m ²
DATE: _____				
RETURN APPOINTMENT ORDERS				
<input type="checkbox"/> Return in three weeks for Doctor and Cycle _____.				
<input type="checkbox"/> Return in _____ weeks for Doctor and Cycle(s) _____.				
CBC & Diff, platelets prior to Cycle #2				
<input type="checkbox"/> CBC & Diff, platelets every 12 weeks				
If clinically indicated x _____ weeks:				
<input type="checkbox"/> ECG <input type="checkbox"/> Echocardiogram <input type="checkbox"/> MUGA Scan <input type="checkbox"/> CA15-3				
<input type="checkbox"/> Tot. Prot <input type="checkbox"/> Albumin <input type="checkbox"/> Bilirubin <input type="checkbox"/> GGT <input type="checkbox"/> Alk Phos.				
<input type="checkbox"/> LDH <input type="checkbox"/> ALT <input type="checkbox"/> BUN <input type="checkbox"/> Creatinine				
<input type="checkbox"/> Other tests:				
<input type="checkbox"/> Consults:				
<input type="checkbox"/> See general orders sheet for additional requests.				
DOCTOR'S SIGNATURE: _____			SIGNATURE: _____	
			UC: _____	