



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

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;"> <thead> <tr> <th style="width: 15%;">Drug</th> <th style="width: 45%;">Brand (Pharmacist to complete. Please print.)</th> <th style="width: 40%;">Pharmacist Initial and Date</th> </tr> </thead> <tbody> <tr> <td>trastuzumab</td> <td></td> <td></td> </tr> </tbody> </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											
<table border="1" style="width:100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 15%;">Drug</th> <th style="width: 45%;">Brand (Pharmacist to complete. Please print.)</th> <th style="width: 40%;">Pharmacist Initial and Date</th> </tr> </thead> <tbody> <tr> <td>trastuzumab</td> <td></td> <td></td> </tr> </tbody> </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											
<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 <input type="checkbox"/> three or <input type="checkbox"/> four weeks (select one) x _____ cycle(s). Observe for 30 minutes post-infusion*.											
Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190											
<table border="1" style="width:100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 15%;">Drug</th> <th style="width: 45%;">Brand (Pharmacist to complete. Please print.)</th> <th style="width: 40%;">Pharmacist Initial and Date</th> </tr> </thead> <tbody> <tr> <td>trastuzumab</td> <td></td> <td></td> </tr> </tbody> </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											
* 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.											
acetaminophen 325 to 650 mg PO PRN for headache and rigors											
Proceed with treatment based on blood work from _____											
DOCTOR'S SIGNATURE:					SIGNATURE:						
					UC:						



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

DOCTOR'S ORDERS		Ht _____ cm Wt _____ kg BSA _____ m ²
DATE: _____		
RETURN APPOINTMENT ORDERS		
<input type="checkbox"/> Return in three or four 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:	