

**PROTOCOL CODE: UBRAVTTCAP**

A BC Cancer "Compassionate Access Program" request form must be completed and approved prior to treatment.

<b>DOCTOR'S ORDERS</b>		Ht _____ cm	Wt _____ kg	BSA _____ m <sup>2</sup>
<b>REMINDER:</b> Please ensure drug allergies and previous bleomycin are documented on the Allergy & Alert Form				
<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, Platelets</b> day of treatment				
May proceed with doses as written if within 96 hours <b>ANC greater than or equal to 1.5 x 10<sup>9</sup>/L, Platelets greater than or equal to 75 x 10<sup>9</sup>/L, Creatinine Clearance greater than 50 mL/min, bilirubin less than or equal to 1.5 x ULN, ALT less than or equal to 5 x ULN</b>				
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:</b>				
<input type="checkbox"/> <b>CYCLE #1:</b>				
trastuzumab 8 mg/kg x _____ kg = _____ mg IV in 250 mL NS over 1 hour 30 minutes on Day 1.				
Observe for 1 hour post infusion.				
Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190				
Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date		
trastuzumab				
tucatinib* 300 mg PO BID on days 1 to 21 continuously				
Dose modification if required:				
<input type="checkbox"/> tucatinib* 250 mg PO BID on days 1 to 21 continuously <input type="checkbox"/> tucatinib* 200 mg PO BID on days 1 to 21 continuously <input type="checkbox"/> tucatinib* 150 mg PO BID on days 1 to 21 continuously				
* Dispense in original container				
capecitabine 1000 mg/m <sup>2</sup> x BSA x ( _____ %) = _____ mg PO BID x 14 days on days 1 to 14.				
(refer to <a href="#">Capecitabine Suggested Tablet Combination Table</a> for dose rounding)				
<b>*** SEE PAGE 2 FOR TREATMENT CYCLE 2 ONWARDS***</b>				
<b>DOCTOR'S SIGNATURE:</b>				<b>SIGNATURE:</b>
				<b>UC:</b>



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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](http://www.bccancer.bc.ca/terms-of-use) and according to acceptable standards of care.

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<b>DOCTOR'S ORDERS</b>	
DATE:	
<b>TREATMENT: (Continued)</b>	
<b>*** SEE PAGE 1 FOR TREATMENT CYCLE 1 ***</b>	
<u>OR</u>	
<input type="checkbox"/> <b>CYCLE #2</b>	
trastuzumab 6 mg/kg x _____ kg = _____ mg IV in NS 250 mL over NS over 1 hour.	
Observe for 30 minutes post infusion.	
Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190	
<b>Drug</b>	<b>Brand (Pharmacist to complete. Please print.)</b>
trastuzumab	
<b>tucatinib* 300 mg</b> PO BID on days 1 to 21 continuously. Dose modification if required: <input type="checkbox"/> <b>tucatinib* 250 mg</b> PO BID on days 1 to 21 continuously <input type="checkbox"/> <b>tucatinib* 200 mg</b> PO BID on days 1 to 21 continuously <input type="checkbox"/> <b>tucatinib* 150 mg</b> PO BID on days 1 to 21 continuously * Dispense in original container	
<b>capecitabine 1000 mg/m<sup>2</sup></b> x BSA x ( _____ %) = _____ mg PO BID x 14 days on days 1 to 14. (refer to <a href="#">Capecitabine Suggested Tablet Combination Table</a> for dose rounding)	
<u>OR</u>	
<input type="checkbox"/> <b>CYCLE 3 and subsequent</b>	
trastuzumab 6 mg/kg x _____ kg = _____ mg IV in 250 mL NS over 30 minutes.	
Observe for 30 minutes post infusion. Observation period not required after 3 treatments with no reaction.	
Pharmacy to select trastuzumab brand as per Provincial Systemic Therapy Policy III-190	
<b>Drug</b>	<b>Brand (Pharmacist to complete. Please print.)</b>
trastuzumab	
<b>tucatinib* 300 mg</b> PO BID on days 1 to 21 continuously. Dose modification if required: <input type="checkbox"/> <b>tucatinib* 250 mg</b> PO BID on days 1 to 21 continuously <input type="checkbox"/> <b>tucatinib* 200 mg</b> PO BID on days 1 to 21 continuously <input type="checkbox"/> <b>tucatinib* 150 mg</b> PO BID on days 1 to 21 continuously * Dispense in original container	
<b>capecitabine 1000 mg/m<sup>2</sup></b> x BSA x ( _____ %) = _____ mg PO BID x 14 days on days 1 to 14. (refer to <a href="#">Capecitabine Suggested Tablet Combination Table</a> for dose rounding)	
<b>acetaminophen 325 mg to 650 mg</b> PO PRN for headache and rigors	
<b>DOCTOR'S SIGNATURE:</b>	<b>SIGNATURE:</b>
	<b>UC:</b>



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<b>DOCTOR'S ORDERS</b>	
DATE:	
<b>RETURN APPOINTMENT ORDERS</b>	
<input type="checkbox"/> Return in <b>three</b> weeks for Doctor and Cycle _____. <input type="checkbox"/> Last Cycle. Return in _____ weeks.	
<b>CBC &amp; Diff, Platelets, creatinine, bilirubin, ALT</b> prior to each cycle <b>Cycle 1: weekly nursing assessment</b> <input type="checkbox"/> INR Weekly <input type="checkbox"/> INR prior to each cycle If clinically indicated: <input type="checkbox"/> Tot. Prot <input type="checkbox"/> Albumin <input type="checkbox"/> GGT <input type="checkbox"/> Alk Phos <input type="checkbox"/> LDH <input type="checkbox"/> BUN <input type="checkbox"/> CA 15-3 <input type="checkbox"/> Other tests: <input type="checkbox"/> ECG <input type="checkbox"/> Echocardiogram <input type="checkbox"/> MUGA Scan <input type="checkbox"/> Weekly nursing assessment <input type="checkbox"/> Consults: <input type="checkbox"/> See general orders sheet for additional requests.	
<b>DOCTOR'S SIGNATURE:</b>	<b>SIGNATURE:</b> <b>UC:</b>