

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 and according to acceptable standards of care

PROTOCOL CODE: GICOXB

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DOCTOR'S ORDERS	Ht	cm	Wt	kg BSA	m²	
REMINDER: Please ensure drug allergies and previous bleomycin are documented on the Allergy & Alert Form						
DATE: To b	e given:			Cycle(s) #:		
Date of Previous Cycle:						
□ Delay treatment week(s) □ CBC & Diff day of treatment May proceed with doses as written if within 96 hours ANC greater than or equal to 1.2 x 10 ⁹ /L, platelets greater than or equal to 75 x 10 ⁹ /L, creatinine clearance greater than or equal to 50 mL/minute, BP less than or equal to 160/100. For those patients on warfarin, hold bevacizumab if INR greater than 3						
Dose modification for: Hematology Other Toxicity						
Proceed with treatment based on blood work from						
PREMEDICATIONS: Patient to take own supply. RN/Pharmacist to confirm						
ondansetron 8 mg PO prior to treatment			_			
dexamethasone ☐ 8 mg or ☐ 12 mg (select premedication ordered) ☐ For prior oxaliplatin hypersensitivity reaction	ns (Grade 1 or 2	2):	`		sone IV	
45 minutes prior to oxaliplatin: dexamethasone 20 mg IV in 50 mL NS over 15 minutes 30 minutes prior to oxaliplatin: diphenhydrAMINE 50 mg IV in NS 50 mL over 15 minutes and famotidine 20 mg IV in NS 100 mL over 15 minutes (Y-site compatible)						
NO ice chips						
Other:						
** Have Hypersensitivity Reaction Tray & Protocol Available**						
TREATMENT: Repeat in three weeks oxaliplatin line to be primed with D5W; bevacizumab line to be primed with NS						
oxaliplatin 130 mg/m² x BSA = mg ☐ Dose Modification:mg/m² x IV in 250 to 500 mL D5W over 2 hours. Flus	x BSA =	mg ' pre and p	ost oxali _l	olatin		
For moderate vascular pain during oxaliplatin peripheral administration 250 mL D5W at maximum rate of 125 mL/h concurrently with oxaliplatin prn OR						
bevacizumab 7.5 mg/kg x kg = _ IV in 100 mL NS over 15 minutes. Flush line (Blood pressure measurement pre and post					ubsequent cycles)	
Pharmacy to select bevacizumab brand as per	Provincial Systen	nic Therapy	Policy III-	190		
Drug Brand (Pharmacist to comp	lete. Please prin	t.)	Pharma	cist Initial and Date	,	
bevacizumab						
conscitation 1000 mg/m² or	0/\-		na DO D	ID v 14 days		
capecitabine 1000 mg/m² or x BSA x (%) = mg PO BID x 14 days (refer to Capecitabine Suggested Tablet Combination Table for dose rounding)						
DOCTOR'S SIGNATURE:				SIGI	NATURE:	
				UC:		



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DATE:					
RETURN APPOINTMENT ORDERS					
Return in three weeks for Doctor and Cycle					
Return in <u>six</u> weeks for Doctor and Cycle & Book treatment x 2 cycles					
Last Cycle. Return in week(s)					
CBC & Diff, creatinine, total bilirubin, ALT prior to each cycle					
Dipstick Urine or laboratory urinalysis for protein at the beginning of each even numbered cycle. (If results are 2+ or 3+ or greater than or equal to 1 g/L laboratory urinalysis for protein then □ 24 hr urine for total protein must be done within 3 days prior to next cycle.) If clinically indicated:					
□ CEA □ CA 19-9 □ ECG					
☐ alkaline phosphatase ☐ albumin ☐ GGT ☐ sodium ☐ potassium					
☐ INR weekly ☐ INR prior to each cycle					
☐ Other tests:					
☐ Weekly nursing assessment for (specify concern):					
☐ Consults:					
☐ See general orders sheet for additional requests.					
DOCTOR'S SIGNATURE:	SIGNATURE:				
	UC:				