

PROTOCOL CODE: GIAVTTB

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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 #: _____
Date of Previous Cycle: _____		
<input type="checkbox"/> Delay treatment _____ week(s) <input type="checkbox"/> CBC & Diff day of treatment		
If within 96 hours of Day 1: ANC greater than or equal to 1.5 x 10⁹/L, platelets greater than or equal to 75 x 10⁹/L, BP less than or equal to 160/100 mmHg. For patients on warfarin, hold bevacizumab if INR greater than 3. And if ordered, previous cycle Day 15 ANC greater than or equal to 0.5 x 10⁹/L, platelets greater than or equal to 25 x 10⁹/L		
May proceed with doses as written if on Day 15: BP less than or equal to 160/100 mmHg. For patients on warfarin, hold bevacizumab if INR greater than 3.		
Dose modification for: <input type="checkbox"/> Hematology <input type="checkbox"/> Other Toxicity _____ Proceed with treatment based on blood work from _____		
PREMEDICATIONS: Patient to take own supply. RN/Pharmacist to confirm _____.		
Antiemetics per protocol		
<input type="checkbox"/> Other: _____		
TREATMENT: <input type="checkbox"/> Repeat in 4 weeks		
bevacizumab 5 mg/kg x _____ kg = _____ mg on Days 1 and 15. IV in 100 mL NS over 15 minutes. (Blood pressure measurement pre and post dose for first 3 doses and prior to bevacizumab for subsequent cycles.) Pharmacy to select bevacizumab brand as per Provincial Systemic Therapy Policy III-190		
Drug	Brand (Pharmacist to complete. Please print.)	Pharmacist Initial and Date
bevacizumab		
trifluridine-tipiracil 35 mg/m² x BSA = _____ mg PO (Maximum dose = 80 mg/dose; based on trifluridine component) <i>twice</i> daily on Days 1-5 and 8-12 of each 28 days cycle. Round dose to nearest 5 mg. Dose modification: (Maximum dose = 80 mg/dose; based on trifluridine component, Round dose to nearest 5 mg)		
<input type="checkbox"/> trifluridine-tipiracil 30 mg/m² x BSA = _____ mg PO <i>twice</i> daily on Days 1-5 and 8-12 of each 28 days cycle (dose level -1) Supply for: _____ days.		
<input type="checkbox"/> trifluridine-tipiracil 25 mg/m² x BSA = _____ mg PO <i>twice</i> daily on Days 1-5 and 8-12 of each 28 days cycle (dose level -2) Supply for: _____ days.		
<input type="checkbox"/> trifluridine-tipiracil 20 mg/m² x BSA = _____ mg PO <i>twice</i> daily on Days 1-5 and 8-12 of each 28 days cycle (dose level -3) Supply for: _____ days.		
<input type="checkbox"/> trifluridine-tipiracil _____ mg/m² x BSA = _____ mg PO <i>twice</i> daily on Days 1-5 and 8-12 of each 28 days cycle. Supply for: _____ days.		
DOCTOR'S SIGNATURE:		SIGNATURE:
		UC:

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Date:	
RETURN APPOINTMENT ORDERS	
<input type="checkbox"/> Return in four weeks for Doctor and Cycle _____. Book chemo on Days 1 and 15 <input type="checkbox"/> Return in eight weeks for Doctor and Cycle _____. Book chemo x 2 cycles on Days 1 and 15. <input type="checkbox"/> Return in _____ weeks for Doctor and Cycle _____. <input type="checkbox"/> Last Cycle. Return in _____ week(s).	
<p>CBC & Diff, creatinine, total bilirubin, ALT, dipstick urine OR laboratory urinalysis for protein prior to each cycle</p> <p>CBC & Diff on Day 15 of Cycle 1</p> <p>If clinically indicated:</p> <input type="checkbox"/> CBC & Diff on Day 15 (for Cycle 2 onwards) <input type="checkbox"/> CEA <input type="checkbox"/> CA 19-9 <input type="checkbox"/> ECG <input type="checkbox"/> alkaline phosphatase <input type="checkbox"/> albumin <input type="checkbox"/> GGT <input type="checkbox"/> sodium <input type="checkbox"/> potassium <input type="checkbox"/> 24 hr urine for total protein (must be done 3 days prior to next cycle, if urine protein 2+ or 3+ or greater than or equal to 1 g/L laboratory urinalysis for protein) <input type="checkbox"/> INR weekly <input type="checkbox"/> INR prior to each cycle <input type="checkbox"/> Other tests: <input type="checkbox"/> Consults: <input type="checkbox"/> See general orders sheet for additional requests.	
DOCTOR'S SIGNATURE:	SIGNATURE:
	UC: