

Doctor's Order Sheet  
CAPOX + Bevacizumab Regimen:  
**Oxaliplatin-Capecitabine  
-Bevacizumab (Part I)**  
Adult Chemotherapy- Medical Oncology  
Metastatic Colorectal Carcinoma



CC1340 0005 06 2017

Name: \_\_\_\_\_

HCN: \_\_\_\_\_

Date of Birth: \_\_\_\_\_

**Allergies:**

☐ No Known

Date: DD/MONTH/YYYY Planned Administration Date: DD/MONTH/YYYY  
Cycle \_\_\_\_\_ of \_\_\_\_\_ **Cycle Duration: 21 days** Date of previous cycle: DD/MONTH/YYYY

**MAY PROCEED WITH DOSES AS WRITTEN IF:**

- ANC **greater than or equal to**  $1.2 \times 10^9/L$  and platelets **greater than or equal to**  $75 \times 10^9/L$ , BP **less than or equal to** 160/100, Creatinine Clearance **greater than** 50 mL/minute, otherwise notify Medical Oncologist.
- LFT's and Bilirubin assessed.
- Dipstick Urine or laboratory urinalysis for protein at the beginning of each odd (1, 3, 5) numbered cycle. If results are 2+ or 3+ or greater than or equal to 1 g/L laboratory urinalysis for protein, collect 24 hour urine for total protein within 3 days before the next cycle. If this result is abnormal, dose reductions are required

**PREMEDICATIONS:**

- ☐ Ondansetron 8 mg PO
- ☐ Dexamethasone 8 mg PO
- ☐ Other: \_\_\_\_\_

Authorized Prescriber: \_\_\_\_\_ Date: DD/MONTH/YYYY Time: \_\_\_\_\_

Authorized Prescriber's Signature: \_\_\_\_\_ ID #: \_\_\_\_\_

Nurse's Name: \_\_\_\_\_ Date: DD/MONTH/YYYY Time: \_\_\_\_\_

Nurse's Signature: \_\_\_\_\_

THIS IS A CONTROLLED DOCUMENT; PLEASE ENSURE THAT YOU ARE READING THE MOST RECENT VERSION. USER WILL BE SOLELY RESPONSIBLE FOR VERIFYING ITS CURRENCY AND ACCURACY.

Doctor's Order Sheet  
CAPOX + Bevacizumab Regimen:  
**Oxaliplatin-Capecitabine  
-Bevacizumab (Part II)**  
Adult Chemotherapy- Medical Oncology  
Metastatic Colorectal Carcinoma



CC1340 0005 06 2017

Name: \_\_\_\_\_

HCN: \_\_\_\_\_

Date of Birth: \_\_\_\_\_

Weight: \_\_\_\_\_ kg Height: \_\_\_\_\_ cm Body Surface Area (BSA) = \_\_\_\_\_

**CHEMOTHERAPY (FOR HOSPITAL PHARMACY):**

- ☐ **Oxaliplatin 130 mg/m<sup>2</sup>** X BSA = \_\_\_\_\_ mg IV in 500 mL D5W over 120 minutes on day 1
- ☐ Dose modification: **130 mg/m<sup>2</sup>** X BSA - \_\_\_\_\_ % = \_\_\_\_\_ mg IV in 500 mL D5W over 120 minutes on day 1
- ☐ **Bevacizumab 7.5 mg/kg** X weight (kg) = \_\_\_\_\_ mg IV in 100 mL normal saline over:
- **90** minutes during **Cycle 1**;
  - If tolerated without reaction- **60** minutes during **Cycle 2**;
  - If tolerated without reaction- **30** minutes during **Cycle 3**;
  - If tolerated without reaction-**15** minutes during **Cycle 4** and all other cycles

(Prior to and post administration, flush lines with normal saline as Bevacizumab is not compatible with D5W; blood pressure measurement pre and post dose for first 3 cycles; and prior to Bevacizumab for subsequent cycles)

**CHEMOTHERAPY (FOR COMMUNITY PHARMACY):**

- ☐ **Capecitabine 1000 mg/m<sup>2</sup>** X BSA = \_\_\_\_\_ mg PO bid with food on days 1 to 14
- ☐ Dose modification: **1000 mg/m<sup>2</sup>** X BSA - \_\_\_\_\_ % = \_\_\_\_\_ mg PO bid with food on days 1 to 14

This prescription is NOT eligible for medication management by a pharmacist.

Authorized Prescriber: \_\_\_\_\_ Date: DD/MONTH/YYYY Time: \_\_\_\_\_

Authorized Prescriber's Signature: \_\_\_\_\_ ID #: \_\_\_\_\_

Nurse's Name: \_\_\_\_\_ Date: DD/MONTH/YYYY Time: \_\_\_\_\_

Nurse's Signature: \_\_\_\_\_

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