



CC2330 0104 09 2016

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

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

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

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

**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**  $1.5 \times 10^9/L$  and platelets **greater than or equal to**  $75 \times 10^9/L$ , Creatinine Clearance **greater than** 50 mL/minute, otherwise notify Medical Oncologist.
- LFT's and Bilirubin assessed.

**CHEMOTHERAPY (FOR COMMUNITY PHARMACY):**

- Capecitabine 1250 mg/m<sup>2</sup> X BSA = \_\_\_\_\_ mg PO bid with food on Days 1 to 14**
- Dose modification: **1250 mg/m<sup>2</sup> X BSA - \_\_\_\_\_% = \_\_\_\_\_ mg PO bid with food on Days 1 to 14**
- Mitte: \_\_\_\_\_ x 500 mg tablets and \_\_\_\_\_ x 150 mg tablets ( \_\_\_\_\_ days supply)

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.