

Doctor's Order Sheet  
FEC-D Regimen (FEC arm):

**(Cycles 1-3)  
fluorouracil - EpiRUBicin -  
cyclophosphamide (Part I)**

**ARIA Protocol Name: FEC-D**

Adult Chemotherapy- Medical Oncology

Adjuvant Breast Cancer Therapy



CC1710 0042 10 2019

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**  $1.5 \times 10^9/L$  and platelets **greater than or equal to**  $100 \times 10^9/L$ , otherwise notify Medical Oncologist.
- LFT's and Bilirubin assessed.
- Creatinine clearance assessed.

**PREMEDICATIONS**

- fosaprepitant 150 mg IV in 150 mLs normal saline over 30 minutes  
**OR**  
 aprepitant 125 mg PO followed by 80 mg PO on days 2 and 3  
 ondansetron 16 mg PO  
 dexamethasone 12 mg PO

**HYDRATION/SUPPORTIVE CARE**

- Start IV infusion with normal saline 1000 mL and infuse with fluorouracil and epiRUBicin so that 1000 mL is infused prior to cyclophosphamide.

PLEASE REFER TO CHEMOTHERAPY LETTER WHEN ORDERING SUPPORTIVE MEDICATIONS FOR THIS PATIENT

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.

Doctor's Order Sheet  
FEC-D Regimen (FEC arm):  
**(Cycles 1-3)**  
**fluorouracil - EpiRUBicin -  
cyclophosphamide**  
(Part II)

**ARIA Protocol Name:** FEC-D  
Adult Chemotherapy- Medical Oncology  
Adjuvant Breast Cancer Therapy



CC1710 0042 10 2019

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

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

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

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

**CHEMOTHERAPY:**

- epiRUBicin 100 mg/m<sup>2</sup>** X BSA = \_\_\_\_\_ mg IV push on day 1
- Dose modification: **100 mg/m<sup>2</sup>** X BSA - \_\_\_\_\_ % = \_\_\_\_\_ mg IV push on day 1
  
- fluorouracil 500 mg/m<sup>2</sup>** X BSA = \_\_\_\_\_ mg IV in 100 mLs D5W over 30 minutes on day 1
- Dose modification: **500 mg/m<sup>2</sup>** X BSA - \_\_\_\_\_ % = \_\_\_\_\_ mg IV in 100 mLs D5W over 30 minutes on day 1
  
- cyclophosphamide 500 mg/m<sup>2</sup>** X BSA = \_\_\_\_\_ mg IV in 100 mLs normal saline over 60 minutes on day 1 (doses greater than 1000 mg must be diluted in 250 mLs normal saline)
- Dose modification: **500 mg/m<sup>2</sup>** X BSA - \_\_\_\_\_ % = \_\_\_\_\_ mg IV in 100 mLs normal saline over 60 minutes on day 1

**POST CHEMOTHERAPY**

- filgrastim (Brand: \_\_\_\_\_)** \_\_\_\_\_ mcg subcutaneous daily for 7 days starting 24-48 hours post chemotherapy
- pegfilgrastim (Brand: \_\_\_\_\_)** **6 mg** subcutaneous x one dose 24-48 hours post chemotherapy

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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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