

# **Chemotherapy protocol**

## **DRUG REGIMEN**

EC100

## **Indications for use**

Neoadjuvant and adjuvant treatment for breast cancer

## Regimen

**DRUG** 

Epirubicin 100mg/m<sup>2</sup> IV bolus Cyclophosphamide 500mg/m<sup>2</sup> IV bolus

Regimen given every 3 weeks as part of the following programmes:

EC100-T: 3 cycles of EC100 followed by 3 cycles of docetaxel

EC100-Paclitaxel: 3 cycles of EC100 followed by 9 doses of weekly paclitaxel

EC100-TPH: 3 cycles of EC100 followed by 3 cycles of docetaxel, trastuzumab and pertuzumab

#### Investigation prior to initiating treatment

**FBC** 

Routine Biochemistry CXR at clinician's discretion

Echocardiogram / MUGA scan (see protocol)

#### **Cautions**

Pre-existing cardiac morbidity
LVEF < 50% (consider alternative therapy)
Altered LFT

### Investigations and consultations prior to each cycle

To be seen by clinician before every cycle.

**FBC** 

U&Es

**LFTs** 

The U&Es and LFTs may be retrospectively looked at (i.e. after the chemotherapy treatment) <u>unless</u> they are known to be abnormal then they need to be repeated the day before so that the results are available prechemotherapy

#### Side Effects

Nausea and vomiting, alopecia, mucositis, possible diarrhoea, myelosuppression, cardiac side effects, vein pain

# Acceptable levels for treatment to proceed (if outside these defer one week or contact consultant)

Neutrophil > 1.5 and plts > 100

If Neutrophils 1.2-1.5 contact consultant

## **Dose Modification Criteria**

None specific

#### Specific Information on Administration

Epirubicin is a vesicant and should be administered first via the side port of a fast running infusion. Primary GCSF prophylaxis should be given

THIS PROTOCOL HAS BEEN DIRECTED BY  $\underline{\mathsf{DR}}\ \mathsf{HOGG}, \mathsf{DESIGNATED}\ \mathsf{LEAD}\ \mathsf{CLINICIAN}\ \mathsf{FOR}\ \mathsf{BREAST}\ \mathsf{CANCER}$ 

# RESPONSIBILITY FOR THIS PROTOCOL LIES WITH THE HEAD OF SERVICE

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VERSION 1