



Doctor's Order Sheet

**Perjeta® (PERTuzumab) 420 mg -  
Herceptin® (trastuzumab) 6 mg/kg  
- Abraxane® (NAB-PACLitaxel)**

**100 Regimen: Post Cycle 1  
(Part I)**

**ARIA Protocol Name: PERT/Herceptin(trastuzumab)21 ABRAX100**

Adult Chemotherapy - Medical Oncology

Metastatic Breast Cancer Therapy

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

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

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



CC1850 0056 06 2020

<b>Allergies:</b>	<input type="checkbox"/> <b>No Known</b>
Date: <u>DD/MONTH/YYYY</u> Cycle _____ of _____ <b>Cycle Duration: 21 days</b>	
Planned Administration Date: <u>DD/MONTH/YYYY</u> Date of previous cycle: <u>DD/MONTH/YYYY</u>	
<b>MAY PROCEED WITH DOSES AS WRITTEN IF:</b>	
<ul style="list-style-type: none"><li>• ANC <b>greater than or equal to</b> 1.5 X 10<sup>9</sup>/L and platelets <b>greater than or equal to</b> 100 X 10<sup>9</sup>/L, otherwise notify Medical Oncologist.</li><li>• LFTs and Bilirubin assessed.</li><li>• Creatinine clearance assessed.</li></ul>	
<b>PREMEDICATIONS:</b>	
<input type="checkbox"/> Other: _____	

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.

**Perjeta®(PERTuzumab) 420 mg -  
Herceptin®(trastuzumab) 6 mg/kg  
- Abraxane®(NAB-PACLitaxel)  
100 Regimen: Post Cycle 1  
(Part II)**

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

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

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

**ARIA Protocol Name:** PERT/Herceptin(trastuzumab)21 ABRAX100

Adult Chemotherapy - Medical Oncology

Metastatic Breast Cancer Therapy



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

**CHEMOTHERAPY:**

**Perjeta® (PERTuzumab) 420 mg**

**IV** in 250 mL normal saline over 60 minutes on day 1

Cycle 3 and beyond can be administered over 30 minutes if no previous reaction.

**Herceptin® (trastuzumab) 6 mg/kg X Weight (kg) = \_\_\_\_\_ mg**

**IV** in 250 mL normal saline over 60 minutes on day 1

Cycle 3 and beyond can be administered over 30 minutes if no previous adverse reaction.

**Abraxane® (NAB-PACLitaxel) 100 mg/m<sup>2</sup> X BSA = \_\_\_\_\_ mg**

Dose modification: **Abraxane® (NAB-PACLitaxel) 100 mg/m<sup>2</sup> X BSA - \_\_\_\_\_ % = \_\_\_\_\_ mg**

**IV** in Viaflex bag over 30 minutes on days 1 and 8

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: \_\_\_\_\_

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