

## Regimen Reference Order

### BRST – pembrolizumab + paclitaxel-protein bound weekly

ARIA: BRST - [pembro + wkly pacl (prot)]

**Planned Course:** pembrolizumab (every 21 days) + paclitaxel-protein bound weekly until disease progression or unacceptable toxicity, up to a maximum of 2 years of therapy (1 cycle = 21 days)

**Indication for Use:** Breast Cancer; Triple Negative; Metastatic

**Drug Alert:** Immune Checkpoint Inhibitor (pembrolizumab)

**CVAD:** At Provider's Discretion

#### Proceed with treatment if:

##### **Day 1**

- ANC equal to or greater than  $1.5 \times 10^9/L$  AND Platelets equal to or greater than  $100 \times 10^9/L$
- AST/ALT equal to or less than 3 times the upper limit of normal
- Total bilirubin equal to or less than 1.5 times the upper limit of normal
- Creatinine clearance is equal to or greater than 30 mL/minute

##### **Days 8 and 15**

- ANC equal to or greater than  $1.5 \times 10^9/L$  AND Platelets equal to or greater than  $100 \times 10^9/L$
- ❖ Contact Physician if parameters not met

## SEQUENCE OF MEDICATION ADMINISTRATION

### Pre-treatment Requirements

Drug	Dose	CCMB Administration Guideline
Not Applicable		

### Treatment Regimen – BRST – pembrolizumab + paclitaxel-protein bound weekly

Establish primary solution 500 mL of: normal saline

Drug	Dose	CCMB Administration Guideline
<b>Day 1</b>		
pembrolizumab	2 mg/kg	IV in normal saline 50 mL over 30 minutes Use 0.2 or 0.22 micron filter
metoclopramide	20 mg	Orally 30 minutes prior to chemotherapy
paclitaxel-protein bound (nab-PACLitaxel)	100 mg/m <sup>2</sup>	IV over 30 minutes (administered undiluted)
<b>Days 8 and 15</b>		
metoclopramide	20 mg	Orally 30 minutes prior to chemotherapy

paclitaxel-protein bound (nab-PACLitaxel)	100 mg/m <sup>2</sup>	IV over 30 minutes (administered undiluted) <i>Use non-DEHP bags and non-DEHP administration sets</i>
<b>Maximum pembrolizumab dose is 200 mg</b> All doses will be automatically rounded that fall within CCMB Approved Dose Bands. See Dose Banding document for more information		

**In the event of an infusion-related hypersensitivity reaction, refer to the ‘Hypersensitivity Reaction Standing Order’**

## REQUIRED MONITORING

### All Cycles

#### Day 1

- CBC, serum creatinine, urea, electrolytes, liver enzymes, total bilirubin, albumin, glucose and TSH as per Physician Orders
- Medical Oncologist or designate (i.e. family practitioner in oncology) must assess patient for immune-mediated adverse reactions prior to each dose of pembrolizumab
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O<sub>2</sub> saturation) at baseline and as clinically indicated
- No observation period is required after pembrolizumab or paclitaxel-protein bound administration. Patient can be discharged from treatment room if stable whether they had a reaction or not

#### Days 8 and 15

- CBC
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O<sub>2</sub> saturation) at baseline and as clinically indicated
- No observation period is required after paclitaxel-protein bound administration. Patient can be discharged from treatment room if stable whether they had a reaction or not

## Recommended Support Medications

Drug	Dose	CCMB Administration Guideline
metoclopramide	10 – 20 mg	Orally every 4 hours as needed for nausea and vomiting

## DISCHARGE INSTRUCTIONS

- Patients should be instructed to contact their cancer team immediately if symptoms of hypersensitivity reactions occur after discharge
- Confirm that patient has received the CCMB Immune Checkpoint Inhibitor Medical Alert wallet card
- Reinforce to patient the immune-mediated adverse reactions and importance of reporting immediately
  - For severe symptoms, the patient should be instructed to go to the nearest emergency room. Oncologist on call should be contacted
- Instruct patient to continue taking anti-emetic(s) at home
- Reinforce applicable safe handling precautions of medications, blood and body fluids for 48 hours after completion of chemotherapy

## ADDITIONAL INFORMATION

- pembrolizumab is an Immune Checkpoint Inhibitor. Consult with medical oncologist for immune-mediated adverse reactions; corticosteroids are often indicated
- paclitaxel-protein bound is also called ABRAXANE®, paclitaxel albumin bound, nab-PACLitaxel or PACLitaxel nanoparticle albumin bound