Regimen Reference Order

GYNE – bevacizumab + PACLitaxel + CARBOplatin (ovarian)

ARIA: GYNE - [bev + PACL + CARBO (Ovarian)]

Planned Course: Cycle 1: PACLitaxel + CARBOplatin, then Cycles 2 to 6: bevacizumab + PACLitaxel + CARBOplatin, then Cycles 7 to 18: bevacizumab (1 cycle= 21 days)

Indication for Use: Ovarian Cancer

CVAD: At Provider's Discretion

Proceed with treatment if:		
Cycle 1		
• ANC equal to or greater than 1.5 x 10 ⁹ /L	AND	Platelets equal to or greater than 100 x 10 ⁹ /L
Cycles 2 to 6		
• ANC equal to or greater than 1.2 x 10 ⁹ /L	AND	Platelets equal to or greater than 75 x 10 ⁹ /L
Cycles 7 to 18		
• ANC equal to or greater than 1.2 x 10 ⁹ /L	AND	Platelets equal to or greater than 75 x 10 ⁹ /L
 Contact Physician if parameters not met 		

SEQUENCE OF MEDICATION ADMINISTRATION

Pre-treatment Requirements		
Drug	Dose	CCMB Administration Guideline
	Ν	ot Applicable

Establish primary solution 500 mL of: normal saline		
Drug	Dose	CCMB Administration Guideline
Cycle 1		
cetirizine	20 mg	Orally 1 hour prior to PACLitaxel
aprepitant	125 mg	Orally 1 hour pre-chemotherapy
ondansetron	16 mg	Orally 30 minutes pre-chemotherapy
dexamethasone 20 mg	IV in normal saline 50 mL over 15 minutes <u>1 hour</u> prior to PACLitaxel	
	*Nursing Alert: PACLitaxel starts 1 hour after completion of dexamethasone infusion	



PACLitaxel	175 mg/m ²	 IV in normal saline 500 mL over 3 hours, following the administration rates below: Administer at 100 mL/hour for 15 minutes, then Administer remaining volume over 2 hours and 45 minutes Use non-DEHP bags and non-DEHP administration sets with 0.2 or 0.22 micron filter
		*Nursing Alert: Gently invert bag 8 to 10 times immediately prior to administration of PACLitaxel to evenly distribute the drug
CARBOplatin	AUC 6 mg/mL.min; maximum dose 900 mg (see table below)	IV in D5W 250 mL over 30 minutes
Cycles 2 to 6		
bevacizumab (brand name specific)	7.5 mg/kg	IV in normal saline 100 mL over 15 minutes *Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order
cetirizine	20 mg	Orally 1 hour prior to PACLitaxel
aprepitant	125 mg	Orally 1 hour pre-chemotherapy
ondansetron	16 mg	Orally 30 minutes pre-chemotherapy
dexamethasone	20 mg	IV in normal saline 50 mL over 15 minutes <u>1 hour</u> prior to PACLitaxel *Nursing Alert: PACLitaxel starts 1 hour after completion of dexamethasone infusion
Wait 1 hour after comp	letion of IV pre-medicati	on(s) before starting PACLitaxel
PACLitaxel	175 mg/m ²	IV in normal saline 500 mL over 3 hours, following the administration rates below:
		 Administer at 100 mL/hour for 15 minutes, then Administer remaining volume over 2 hours and 45 minutes Use non-DEHP bags and non-DEHP administration sets with 0.2
		or 0.22 micron filter *Nursing Alert: Gently invert bag 8 to 10 times immediately prior to administration of PACLitaxel to evenly distribute the drug
CARBOplatin	AUC 6 mg/mL.min; maximum dose 900 mg (see table below)	IV in D5W 250 mL over 30 minutes
Cycles 7 to 18		
bevacizumab (brand name specific)	7.5 mg/kg	IV in normal saline 100 mL over 15 minutes *Alert: Ensure brand name on prescription label (indicated in brackets on prescription label) matches prescribed order



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

Cycle 1

- CBC, serum creatinine and liver enzymes as per Physician Orders
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O₂ saturation) at baseline and as clinically indicated
- No observation period is required after PACLitaxel administration. Patient can be discharged from treatment room if stable whether they had a reaction or not

Cycles 2 to 6

- CBC, serum creatinine, liver enzymes, urine protein and blood pressure as per Physician Orders
 - Urinalysis for protein: Where urinalysis is not possible, use dipstick. If lab urinalysis for protein is greater than or equal to 1 g/L or dipstick proteinuria shows 2+ or 3+, notify prescriber
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O₂ saturation) at baseline and as clinically indicated
- No observation period is required after bevacizumab or PACLitaxel administration. Patient can be discharged from treatment room if stable whether they had a reaction or not

Cycles 7 to 18

- CBC as per Physician Orders
- Urine protein and blood pressure as per Physician Orders
 - Urinalysis for protein: Where urinalysis is not possible, use dipstick. If lab urinalysis for protein is greater than or equal to 1 g/L or dipstick proteinuria shows 2+ or 3+, notify prescriber
- Full vital signs (temperature, heart rate, respiratory rate, blood pressure and O₂ saturation) at baseline and as clinically indicated
- No observation period is required after bevacizumab administration. Patient can be discharged from treatment room if stable whether they had a reaction or not

Recommended Support Medications			
Drug	Dose	Dose CCMB Administration Guideline	
Cycles 1 to 6 ONLY			
aprepitant	80 mg	Orally once daily on Days 2 and 3	
dexamethasone	8 mg	Orally once daily on Days 2 and 3	
metoclopramide	10 – 20 mg	Orally every 4 hours as needed for nausea and vomiting	

DISCHARGE INSTRUCTIONS

All Cycles

• Patients should be instructed to contact their cancer team immediately if symptoms of hypersensitivity reactions occur after discharge

Cycles 1 to 6

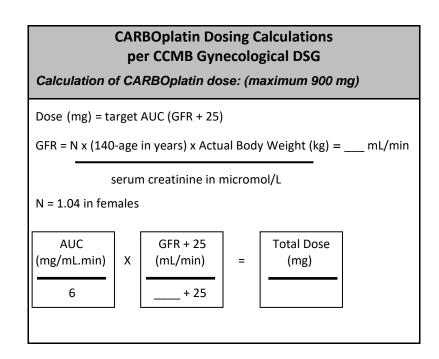
• Instruct patient to continue taking anti-emetic(s) at home



• Reinforce applicable safe handling precautions of medications, blood and body fluids for 48 hours after completion of chemotherapy

ADDITIONAL INFORMATION

- bevacizumab can cause increased risk of hypertension, post-operative bleeding, would healing complications and thromboembolic events
- bevacizumab is available from more than one manufacturer and uses several different brand names. Brand name
 will be indicated in brackets after bevacizumab. Ensure prescription label matches the brand name on prescribed
 order
- PACLitaxel may cause progressive, irreversible neuropathy
- CARBOplatin dose considerations:
 - o CCMB Gynecological DSG uses actual body weight to calculate GFR
 - o CCMB Gynecological DSG uses a maximum CARBOplatin dose of 900 mg for this regimen
 - If calculated CARBOplatin dose differs more than 10% from prescribed CARBOplatin dose, contact the prescriber



<u>AUC = Area Under Curve</u>

The estimated creatinine clearance is based on limited evidence. Sound clinical judgment and interpretation of the estimation are required, because the equation may not be appropriate for some patient populations (for example, acute renal failure).

