

## Patient Details

Forename	Surname	Protocol	ESCALATED BEACOPDAC	SA (m <sup>2</sup> )
Address	DOB	Patient NO	Local No.	Course Name: Escalated BEACOPDac SA<2.1m2 cycle 1
Consultant	Ward	Type of line	Diagnosis	Height (m)
NHS No	No. of lumen:	Weight (kg)		

Monitoring	Acceptable Range	Date Due	Date of Test	Value	Checked
Height (m)					
Weight (kg)					
SA (m <sup>2</sup> )					
ALA TRANSAM 1.5ULN	0.00 60.00	Day 1			
BILIRUBIN 1.5ULN	0.00 31.50	Day 1			
COCKCROFT (>60)	60.00 300.00	Day 1			
NEUTROPHILS > 1.0	1.00 15.00	Day 1			
PLATELETS > 80	80.00 600.00	Day 1			
WHITE BLOOD CELL > 2.5	2.50 11.00	Day 1			

**Additional Prescribing Notes**

Doxorubicin and vincristine can cause pain and tissue necrosis if extravasated.

Doses capped at 2.1m/2

Day	Date and Time	Drug and dose (per m2) or dose (per kg)	ACTUAL DOSE	Infusion Fluid and Final Volume	Route	Additives	Time/Infusion Rate	Line	Given/Checked by	Time Start/Stop	Comments
1	T=hrs	FREE FLOWING INFUSION (500ml)	500 ml	SODIUM CHLORIDE 0.9%	IV				/	/	
1	T=hrs	AKYNZEO 300MG/0.5MG (1capsule)	1 capsule		PO				/	/	To be given 60 minutes prior to chemotherapy
1	T=hrs	DOXORUBICIN (35mg/m <sup>2</sup> )	mg	None	IV		Slow Bolus		/	/	
1	T=hrs	CYCLOPHOSPHAMIDE (1250mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 250 ml	IV		Infuse over 30 Mins at a rate 500 ml/hr		/	/	

Allocated by:	Confirmed by:	Authorised by:	Checked by: (Pharmacist)	Parenteral	3
Date:	Date:	Date:	Date:	Intrathecal	0
				Oral	4

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Forename \_\_\_\_\_ Surname \_\_\_\_\_ Protocol \_\_\_\_\_

Address \_\_\_\_\_

DOB \_\_\_\_\_ Patient NO \_\_\_\_\_ Local No. \_\_\_\_\_ Course Name: \_\_\_\_\_

Ward \_\_\_\_\_

NHS No \_\_\_\_\_

ESCALATED BEACOPDAC

Escalated BEACOPDac SA<2.1m2 cycle 1

SA (m<sup>2</sup>) \_\_\_\_\_

Height (m) \_\_\_\_\_

Weight (kg) \_\_\_\_\_

Day	Date and Time	Drug and dose (per m2) or dose (per kg)	ACTUAL DOSE	Infusion Fluid and Final Volume	Route	Additives	Time/Infusion Rate	Line	Given/Checked by	Time Start/Stop	Comments
1	T=hrs	<b>ETOPOSIDE</b> (200mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 1000 ml	IV		Infuse over 2 Hrs at a rate 500 ml/hr		/	/	
2	T=hrs	FREE FLOWING INFUSION (500ml)	500 ml	SODIUM CHLORIDE 0.9%	IV				/	/	
2	T=hrs	<b>DACARBAZINE</b> (250mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 500 ml	IV		Infuse over 30 Mins at a rate 1000 ml/hr		/	/	
2	T=hrs	<b>ETOPOSIDE</b> (200mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 1000 ml	IV		Infuse over 2 Hrs at a rate 500 ml/hr		/	/	
3	T=hrs	FREE FLOWING INFUSION (500ml)	500 ml	SODIUM CHLORIDE 0.9%	IV				/	/	
3	T=hrs	<b>DACARBAZINE</b> (250mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 500 ml	IV		Infuse over 30 Mins at a rate 1000 ml/hr		/	/	
3	T=hrs	<b>ETOPOSIDE</b> (200mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 1000 ml	IV		Infuse over 2 Hrs at a rate 500 ml/hr		/	/	

<b>Allocated by:</b>	<b>Confirmed by:</b>	<b>Authorised by:</b>	<b>Checked by: (Pharmacist)</b>
<b>Date:</b>	<b>Date:</b>	<b>Date:</b>	<b>Date:</b>

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Address \_\_\_\_\_

DOB \_\_\_\_\_ Patient NO \_\_\_\_\_ Local No. \_\_\_\_\_ Course Name: \_\_\_\_\_

Ward \_\_\_\_\_

NHS No \_\_\_\_\_

ESCALATED BEACOPDAC

Escalated BEACOPDac SA<2.1m2 cycle 1

SA (m<sup>2</sup>) \_\_\_\_\_

Height (m) \_\_\_\_\_

Weight (kg) \_\_\_\_\_

Day	Date and Time	Drug and dose (per m2) or dose (per kg)	ACTUAL DOSE	Infusion Fluid and Final Volume	Route	Additives	Time/Infusion Rate	Line	Given/Checked by	Time Start/Stop	Comments
8	T=hrs	ONDANSETRON (8mg)	8 mg	None	PO				/	/	
8	T=hrs	VINCRIStINE (1.4mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 50 ml	IV				/	/	Max dose: 2mg. Infuse over 5-10 minutes.
8	T=hrs	FREE FLOWING INFUSION (100ml)	100 ml	SODIUM CHLORIDE 0.9%	IV				/	/	
8	T=hrs	HYDROCORTISONE (100mg)	100 mg	None	IV		Slow Bolus		/	/	
8	T=hrs	BLEOMYCIN (10000unit/m <sup>2</sup> )	unit	None	IV		Slow Bolus		/	/	

Allocated by:	Confirmed by:	Authorised by:	Checked by: (Pharmacist)
Date:	Date:	Date:	Date:



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		<b>Course Name</b>	Escalated BEACOPDac SA<2.1m2 cycle 1	<b>Height (m)</b>
<b>DOB</b>	<b>Patient NO</b>	<b>Local No.</b>	<b>NHS No</b>	<b>Weight (kg)</b>
		<b>Ward</b>		
<b>Address</b>				

Record drug allergies or sensitivities

			Time	Date															
<b>Drug &amp; dose</b>	METOCLOPRAMIDE																		
<b>Actual dose</b>	10 mg	<b>Duration</b>	PRN																
<b>Route</b>	PO	<b>Start Date</b>																	
<b>Frequency</b>	TDS	<b>Start Day</b>	1																
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																	
		<b>Accuracy check</b>																	
<b>Note</b>	If pre-pack supplied record Batch Number : _____.																		
<b>Drug &amp; dose</b>	CO-TRIMOXAZOLE																		
<b>Actual dose</b>	960 mg	<b>Duration</b>	21 DAYS																
<b>Route</b>	PO	<b>Start Date</b>																	
<b>Frequency</b>	OD MWF	<b>Start Day</b>	1																
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																	
		<b>Accuracy check</b>																	
<b>Note</b>	To be taken DAILY on Mondays, Wednesdays and Fridays. This is continuous treatment. If pre-pack supplied record Batch Number : _____.																		

<b>Allocated by:</b>	<b>Confirmed by:</b>	<b>Authorised by:</b>	<b>Checked by: (Pharmacist)</b>	
<b>Date:</b>	<b>Date:</b>	<b>Date:</b>	<b>Date:</b>	

**Patient Details**

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		<b>Course Name</b>	Escalated BEACOPDac SA<2.1m2 cycle 1	<b>Height (m)</b>
<b>DOB</b>	<b>Patient NO</b>	<b>Local No.</b>	<b>NHS No</b>	<b>Weight (kg)</b>
		<b>Ward</b>		
<b>Address</b>				

Record drug allergies or sensitivities

			Time	Date														
<b>Drug &amp; dose</b>	ACICLOVIR																	
<b>Actual dose</b>	200 mg	<b>Duration</b>	21 DAYS															
<b>Route</b>	PO	<b>Start Date</b>																
<b>Frequency</b>	TDS	<b>Start Day</b>	1															
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																
		<b>Accuracy check</b>																
<b>Note</b>	This is continuous treatment supply original packs. If pre-pack supplied record Batch Number : _____.																	
<b>Drug &amp; dose</b>	RANITIDINE																	
<b>Actual dose</b>	150 mg	<b>Duration</b>	21 DAYS															
<b>Route</b>	PO	<b>Start Date</b>																
<b>Frequency</b>	BD	<b>Start Day</b>	1															
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																
		<b>Accuracy check</b>																
<b>Note</b>	This is continuous treatment supply original packs. If pre-pack supplied record Batch Number : _____.																	

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<b>Date:</b>	<b>Date:</b>	<b>Date:</b>	<b>Date:</b>

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		<b>Course Name</b>	Escalated BEACOPDac SA<2.1m2 cycle 1	<b>Height (m)</b>
<b>DOB</b>	<b>Patient NO</b>	<b>Local No.</b>	<b>NHS No</b>	<b>Weight (kg)</b>
		<b>Ward</b>		
<b>Address</b>				

Record drug allergies or sensitivities

			Time	Date														
<b>Drug &amp; dose</b>	METOCLOPRAMIDE																	
<b>Actual dose</b>	10 mg	<b>Duration</b>	PRN															
<b>Route</b>	PO	<b>Start Date</b>																
<b>Frequency</b>	TDS	<b>Start Day</b>	8															
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																
		<b>Accuracy check</b>																
<b>Note</b>	Discuss with patient and delete if supply not required. If pre-pack supplied record Batch Number : _____.																	
<b>Drug &amp; dose</b>	FILGRASTIM (G-CSF)																	
<b>Actual dose</b>	microgram	<b>Duration</b>	5 DAYS															
<b>Route</b>	SC	<b>Start Date</b>																
<b>Frequency</b>	OD	<b>Start Day</b>	9															
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																
		<b>Accuracy check</b>																
<b>Note</b>	SUBCUTANEOUS BOLUS Discontinue when, after reaching nadir the WBC remains over 1.0 on 3 successive days																	

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<b>Date:</b>	<b>Date:</b>	<b>Date:</b>	<b>Date:</b>