

Patient Details

Forename

Surname

Protocol

VENETOCLAX

SA (m²)

Height (m)

Weight (kg)

DOB

Patient NO

Local No.

Course Name:

VENETOCLAX cycle 2 onwards

Consultant

Ward

Type of line

No. of lumen:

Diagnosis

Chronic lymphocytic leukaemia/Small lymph

NHS No

Monitoring	Acceptable Range		Date Due	Date of Test	Value	Checked
Height (m)						
Weight (kg)						
SA (m²)						
Cockcroft >30mls/min	30.00	300.00	Day [1]			
NEUTROPHILS > 1.0	1.00	15.00	Day [1]			
PLATELETS > 50	50.00	600.00	Day [1]			

Additional Prescribing Notes

Oral hydration consisting of fluid intake of 1.5 to 2 L per day starting at least 48 hours prior to the start of treatment for all subjects prior to first dose and at all subsequent dose increment steps and continued for at least 24 hours after dosing and all of the chemistries laboratory values remain within ULN.

Patients to be managed as per local/national tumour lysis guidelines.

Refer to Venetoclax SPC for dose modifications.

Allocated by:

Confirmed by:

Authorised by:

Checked by: (Pharmacist)

Parenteral

Intrathecal

Oral

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Date:

Date:

Date:

Date:

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Forename	Surname		Protocol	VENETOCLAX					SA (m²)				
			Course Name	VENETOCLAX cycle 2 onwards					Height (m)				
DOB	Patient NO		Local No.		NHS No							Weight (kg)	
Consultant			Ward		Diagnosis		Chronic lymphocytic leukaemia/Small lymph						
Address													

Record drug allergies or sensitivities

				Time	Date													
Drug & dose	VENETOCLAX																	
Actual dose	400 mg	Duration	28 DAYS															
Route	PO	Start Date																
Frequency	OD	Start Day	1															
Quantity		Dispensed by																
Dispensed		Accuracy check																
Note	Avoid grapefruit products,Seville oranges & starfruit. HIGH RISK OF TUMOUR LYSIS. Biochemistry including U&E, phosphate, calcium MUST be checked 6-8 hours after first increased dose is given.																	

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