

## Patient Details

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

Protocol		BENDAMUSTINE-R (NHL) (TRUXIMA)		SA (m <sup>2</sup> )	
Course Name:		Bendamustine+ Rituximab (Truxima) 375mg/m2		Height (m)	
DOB	Patient NO	Local No.	Diagnosis	Weight (kg)	
Consultant	Ward	Type of line	SINGLE LINE		
NHS No	No. of lumen:				

Monitoring	Acceptable Range	Date Due	Date of Test	Value	Checked
Height (m)					
Weight (kg)					
SA (m <sup>2</sup> )					
ALA TRANSAM2.5ULN	0.00 100.00	Day [1]			
BILIRUBIN 1.5ULN	0.00 31.50	Day [1]			
Cockcroft >40ml/min	40.00 300.00	Day [1]			
NEUTROPHILS > 1.0	1.00 15.00	Day [1]			
PLATELETS> 75	75.00 600.00	Day [1]			

**Additional Prescribing Notes**

Administration of rituximab infusions: Refer to and follow Trust guidelines

Encourage patient to take day Day 2 ondansetron on morning of or at least one hour prior to bendamustine treatment.

Patients should only receive irradiated blood products.

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	PARACETAMOL (1000mg)	1000 mg	None	PO				/	/	Should be given 30-60 minutes prior to rituximab infusion.
1	T=hrs	HYDROCORTISONE (100mg)	100 mg	None	IV		Slow Bolus		/	/	Should be given 30-60 minutes prior to rituximab infusion.
1	T=00Hhrs	CHLORPHENAMINE (10mg)	10 mg		IV		Slow Bolus		/	/	Should be given 30-60 minutes prior to rituximab infusion.
1	T=:hrs	RITUXIMAB (TRUXIMA) (375mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 500 ml	IV				/	/	Truxima brand. Variable infusion rate - see additional prescribing notes.
1	T=hrs	ONDANSETRON (8mg)	8 mg	None	PO				/	/	

<b>Allocated by:</b>		<b>Confirmed by:</b>		<b>Authorised by:</b>		<b>Checked by: (Pharmacist)</b>		<table border="1" style="width:100%; border-collapse: collapse;"> <tr> <td>Parenteral</td> <td style="text-align: right;">2</td> </tr> <tr> <td>Intrathecal</td> <td style="text-align: right;">0</td> </tr> <tr> <td>Oral</td> <td style="text-align: right;">2</td> </tr> </table>		Parenteral	2	Intrathecal	0	Oral	2
Parenteral	2														
Intrathecal	0														
Oral	2														
<b>Date:</b>		<b>Date:</b>		<b>Date:</b>		<b>Date:</b>									

Trust location: \_\_\_\_\_

# Parenteral Cytotoxic Chart

## Patient Details

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

Protocol

BENDAMUSTINE-R (NHL) (TRUXIMA)

SA (m<sup>2</sup>)  
Height (m)  
Weight (kg)

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

Bendamustine+ Rituximab (Truxima) 375mg/m2

Ward \_\_\_\_\_

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

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	<b>BENDAMUSTINE</b> (90mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 500 ml	IV		Infuse over 1 Hrs at a rate 500 ml/hr		/	/	
2	T=hrs	ONDANSETRON (8mg)	8 mg	None	PO				/	/	
2	T=hrs	FREE FLOWING INFUSION (500ml)	500 ml	SODIUM CHLORIDE 0.9%	IV				/	/	
2	T=:hrs	<b>BENDAMUSTINE</b> (90mg/m <sup>2</sup> )	mg	SODIUM CHLORIDE 0.9% 500 ml	IV		Infuse over 1 Hrs at a rate 500 ml/hr		/	/	Ensure patient has taken ondansetron

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

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		Course Name	Bendamustine+ Rituximab (Truxima) 375mg/m2										Height (m)	
DOB	Patient NO	Local No.	NHS No											Weight (kg)
Consultant		Ward	Diagnosis											
Address														

Record drug allergies or sensitivities

			Time	Date												
Drug & dose	ALLOPURINOL															
Actual dose	300 mg	Duration	28 DAYS													
Route	PO	Start Date														
Frequency	OD	Start Day	1													
Quantity Dispensed		Dispensed by														
		Accuracy check														
Note	For cycle 1 only, unless required. If pre-pack supplied record Batch Number : _____.															
Drug & dose	METOCLOPRAMIDE															
Actual dose	10 mg	Duration	PRN													
Route	PO	Start Date														
Frequency	TDS	Start Day	1													
Quantity Dispensed		Dispensed by														
		Accuracy check														
Note	If pre-pack supplied record Batch Number : _____.															

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

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		<b>Course Name</b>	Bendamustine+ Rituximab (Truxima) 375mg/m2										<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>	28 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>	If pre-pack supplied record Batch Number : _____.																	
<b>Drug &amp; dose</b>	CO-TRIMOXAZOLE																	
<b>Actual dose</b>	960 mg	<b>Duration</b>	28 DAYS															
<b>Route</b>	PO	<b>Start Date</b>																
<b>Frequency</b>	OD M,W,F	<b>Start Day</b>	1															
<b>Quantity Dispensed</b>		<b>Dispensed by</b>																
		<b>Accuracy check</b>																
<b>Note</b>	960mg of Co-trimoxazole to be taken on Mondays, Wednesdays and Fridays throughout treatment																	

<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>