

Trust location: _____

Parenteral Cytotoxic Chart

Patient Details

Forename

Surname

Protocol

IDELALISIB + RITUXIMAB (TRUXIMA)

DOB

Patient NO

Local No.

Course Name:

Idelalisib 150mg BD+Ritux (Truxima) 500mg/m2 C5-8

Consultant

VALIDATION

Ward

Type of line

No. of lumen:

Diagnosis

Chronic lymphocytic leukaemia/Small lymph

NHS No

SA (m²)

Height (m)

Weight (kg)

| Monitoring | Acceptable Range | | Date Due | Date of Test | Value | Checked | <div>Additional Prescribing Notes</div> <div>Administration of rituximab infusions: Refer to and follow Trust guidelines.</div> |
|-------------------|------------------|--------|----------|--------------|-------|---------|---|
| Height (m) | | | | | | | |
| Weight (kg) | | | | | | | |
| SA (m²) | | | | | | | |
| ALA TRANSAM 3ULN | 0.00 | 120.00 | Day [1] | | | | |
| NEUTROPHILS > 1.0 | 1.00 | 15.00 | Day [1] | | | | |
| PLATELETS> 75 | 75.00 | 600.00 | Day [1] | | | | |
| | | | | | | | |
| | | | | | | | |
| | | | | | | | |

| 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 | | | | <div></div> <div>Batch No.</div> | <div></div> | Should be given 30-60 minutes prior to rituximab infusion. |
| 1 | T=hrs | HYDROCORTISONE (100mg) | 100 mg | None | IV | | Slow Bolus | | <div></div> <div>Batch No.</div> | <div></div> | Should be given 30-60 minutes prior to rituximab infusion. |
| 1 | T=00Hhrs | CHLORPHENAMINE (10mg) | 10 mg | | IV | | Slow Bolus | | <div></div> <div>Batch No.</div> | <div></div> | Should be given 30-60 minutes prior to rituximab infusion. |
| 1 | T=:hrs | RITUXIMAB (TRUXIMA) (500mg/m²) | mg | SODIUM CHLORIDE 0.9% 500 ml | IV | | | | <div></div> <div>Batch No.</div> | <div></div> | Variable infusion rate - follow Trust guidelines. Ensure patient has received chlorphenamine and paracetamol. |

Allocated by:

Confirmed by:

Authorised by:

Checked by: (Pharmacist)

Date:

Date:

Date:

Date:

Chart Id.:

Parenteral

Intrathecal

Oral

1

0

2

Patient Details

| | | | | | |
|------------|------------|-----------|-------------|---|-------------|
| Forename | Surname | | Protocol | IDELALISIB + RITUXIMAB (TRUXIMA) | SA (m²) |
| | | | Course Name | Idelalisib 150mg BD+Ritux (Truxima) 500mg/m2 C5-8 | 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 | IDELALISIB | | | | | | | | | | | | | | | | | |
| Actual dose | 150 mg | | Duration | 28 DAYS | | | | | | | | | | | | | | |
| Route | PO | | Start Date | | | | | | | | | | | | | | | |
| Frequency | BD | | Start Day | 1 | | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | | |
| Note | Supply 1 original container of 60 tablets. Swallow whole with or without food. | | | | | | | | | | | | | | | | | |
| Drug & dose | CO-TRIMOXAZOLE | | | | | | | | | | | | | | | | | |
| Actual dose | 960 mg | | Duration | 28 DAYS | | | | | | | | | | | | | | |
| Route | PO | | Start Date | | | | | | | | | | | | | | | |
| Frequency | OD M,W,F | | Start Day | 1 | | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | | |
| Note | 960mg of Co-Trimoxazole to be taken on Mondays, Wednesdays and Fridays throughout treatment. | | | | | | | | | | | | | | | | | |

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|---------------|---------------|----------------|--------------------------|------------|
| Allocated by: | Confirmed by: | Authorised by: | Checked by: (Pharmacist) | Chart Id.: |
| Date: | Date: | Date: | Date: | |
| / / | / / | / / | / / | |

| | | | | | | | | | | | | | | | | | |
|--|--|----------------|-------------|---|--------|--|--|--|--|--|--|--|-------------|------------|--|--|--|
| Patient Details | | | | | | | | | | | | | | | | | |
| Forename | Surname | | Protocol | IDELALISIB + RITUXIMAB (TRUXIMA) | | | | | | | | | | SA (m²) | | | |
| | | | Course Name | Idelalisib 150mg BD+Ritux (Truxima) 500mg/m2 C5-8 | | | | | | | | | | Height (m) | | | |
| DOB | Patient NO | | Local No. | | NHS No | | | | | | | | Weight (kg) | | | | |
| | | | | | | | | | | | | | | | | | |
| | | | Ward | | | | | | | | | | | | | | |
| Address | | | | | | | | | | | | | | | | | |
| Record drug allergies or sensitivities | | | | | | | | | | | | | | | | | |
| | | | | | | | | | | | | | | | | | |
| | | | | Time | Date | | | | | | | | | | | | |
| | | | | | | | | | | | | | | | | | |
| Drug & dose | ACICLOVIR | | | | | | | | | | | | | | | | |
| Actual dose | 200 mg | | Duration | 28 DAYS | | | | | | | | | | | | | |
| Route | PO | | Start Date | | | | | | | | | | | | | | |
| Frequency | TDS | | Start Day | 1 | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | |
| Note | If pre-pack supplied record Batch Number : _____. | | | | | | | | | | | | | | | | |
| Drug & dose | LOPERAMIDE | | | | | | | | | | | | | | | | |
| Actual dose | 2 mg | | Duration | SEE NOTE | | | | | | | | | | | | | |
| Route | PO | | Start Date | | | | | | | | | | | | | | |
| Frequency | SEE NOTE | | Start Day | 1 | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | |
| Note | Take 4mg after first loose stool then 2mg after each loose stool thereafter upto maximum of 16mg in 24 hours. If pre-pack supplied, record Batch Number: _____ | | | | | | | | | | | | | | | | |

| | | | | |
|---------------|---------------|----------------|--------------------------|------------|
| Allocated by: | Confirmed by: | Authorised by: | Checked by: (Pharmacist) | Chart Id.: |
| Date: | Date: | Date: | Date: | |
| / / | / / | / / | / / | |