

Patient Details

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|--------------|---|---------------|-------------------------------|----------------------|
| Forename | Surname | Protocol | CHOP-RITUXIMAB>65YR (TRUXIMA) | SA (m ²) |
| Address | DOB | Patient NO | Local No. | Height (m) |
| Course Name: | CHOP+ RITUXIMAB (TRUXIMA) + GCSF 21d OVER 65YRS | | | Weight (kg) |
| Consultant | Ward | Type of line | Diagnosis | |
| NHS No | | No. of lumen: | | |

| Monitoring | Acceptable Range | Date Due | Date of Test | Value | Checked |
|---------------------------|------------------|----------|--------------|-------|---------|
| Height (m) | | | | | |
| Weight (kg) | | | | | |
| SA (m ²) | | | | | |
| ANC >1 (5 day expiry) | 1.00 | 15.00 | Day 1 | | |
| BILIRUBIN 1.5ULN | 0.00 | 31.50 | Day 1 | | |
| CREATININE(max 130) | 0.00 | 130.00 | Day 1 | | |
| Platelets >75 (5 day exp) | 75.00 | 600.00 | Day 1 | | |
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Additional Prescribing Notes

Please prescribe supportive/preventative care either on Chemocare or on a separate paper prescription, as per local trust policy and guidance:

- 1) Consider stress ulcer PPI prophylaxis i.e. lansoprazole
- 2) Consider allopurinol 300mg OD (100mg OD if CrCl <20mls/min) for first 4 weeks of treatment only.

Administration of rituximab infusions: Refer to and follow Trust guidelines.

| 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 | PREDNISOLONE (40mg/m ²) | mg | None | PO | | | | / | / | Should be given 30-60 minutes prior to rituxumab infusion from take home supply or ward stock. |
| 1 | T=hrs | HYDROCORTISONE (100mg) | 100 mg | None | IV | | Slow Bolus | | / | / | Can be administered in addition to oral prednisolone if required. |
| 1 | T=hrs | PARACETAMOL (1000mg) | 1000 mg | None | PO | | | | / | / | Should be given 30-60 minutes prior to rituximab infusion. |
| 1 | T=hrs | CHLORPHENAMINE (10mg) | 10 mg | | IV | | Slow Bolus | | / | / | Should be given 30-60 minutes prior to rituximab infusion. |

| | | | | | |
|---------------|---------------|----------------|--------------------------|-------------|---|
| Allocated by: | Confirmed by: | Authorised by: | Checked by: (Pharmacist) | | |
| Date: | Date: | Date: | Date: | Parenteral | 2 |
| | | | | Intrathecal | 0 |
| | | | | Oral | 2 |

Patient Details

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Address _____

DOB _____ Patient NO _____ Local No. _____ Course Name: _____

Ward _____

NHS No _____

CHOP-RITUXIMAB>65YR (TRUXIMA)

CHOP+ RITUXIMAB (TRUXIMA) + GCSF 21d OVER 65YRS

SA (m²) _____

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 | RITUXIMAB (TRUXIMA) (375mg/m ²) | mg | SODIUM CHLORIDE 0.9% 500 ml | IV | | | | / | / | Truxima brand. Variable infusion rate - see additional prescribing notes. |
| 1 | T=hrs | FREE FLOWING INFUSION (500ml) | 500 ml | SODIUM CHLORIDE 0.9% | IV | | | | / | / | |
| 1 | T=hrs | ONDANSETRON (8mg) | 8 mg | None | PO | | | | / | / | |
| 1 | T=:hrs | CYCLOPHOSPHAMIDE (750mg/m ²) | mg | None | IV | | Slow Bolus | | / | / | |
| 1 | T=:hrs | DOXORUBICIN (50mg/m ²) | mg | None | IV | | Slow Bolus | | / | / | |
| 1 | T=:hrs | VINCRIStINE (1.4mg/m ²) | mg | SODIUM CHLORIDE 0.9% 50 ml | IV | | | | / | / | Max dose: 2mg. Infuse over 5-10 minutes. Monitor for signs of extravasation and report any incidents as per trust procedure. |

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| DOB | Patient NO | Local No. | NHS No | Weight (kg) |
| | | | | |
| Consultant | | Ward | Diagnosis | |
| Address | | | | |

Record drug allergies or sensitivities

| | | | | Time | Date | | | | | | | | | | | | | | |
|---------------------------|---|-----------------------|--------|------|------|--|--|--|--|--|--|--|--|--|--|--|--|--|--|
| | | | | | | | | | | | | | | | | | | | |
| Drug & dose | PREDNISOLONE | | | | | | | | | | | | | | | | | | |
| Actual dose | mg | Duration | 5 DAYS | | | | | | | | | | | | | | | | |
| Route | PO | Start Date | | | | | | | | | | | | | | | | | |
| Frequency | OM | Start Day | 1 | | | | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | | | |
| Note | Taken preferably in the morning. First dose to be taken before the rituximab infusion. | | | | | | | | | | | | | | | | | | |
| 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 | Metoclopramide 10mg tablets are prescribed with each cycle, discuss with patient and delete if supply not required. If pre-pack supplied record Batch Number : _____. | | | | | | | | | | | | | | | | | | |

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| Date: | Date: | Date: | Date: | |

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| DOB | Patient NO | Local No. | NHS No | Weight (kg) |
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| | | Ward | | |
| Address | | | | |

Record drug allergies or sensitivities

| | | | Time | Date | | | | | | | | | | | | | | | | |
|---------------------------|---|-----------------------|--------|------|--|--|--|--|--|--|--|--|--|--|--|--|--|--|--|--|
| | | | | | | | | | | | | | | | | | | | | |
| Drug & dose | ONDANSETRON | | | | | | | | | | | | | | | | | | | |
| Actual dose | 8 mg | Duration | 2 DAYS | | | | | | | | | | | | | | | | | |
| Route | PO | Start Date | | | | | | | | | | | | | | | | | | |
| Frequency | BD | Start Day | 1 | | | | | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | | | | |
| Note | If pre-pack supplied record Batch Number : _____. | | | | | | | | | | | | | | | | | | | |
| Drug & dose | FILGRASTIM (G-CSF) | | | | | | | | | | | | | | | | | | | |
| Actual dose | microgram | Duration | 3 DAYS | | | | | | | | | | | | | | | | | |
| Route | SC | Start Date | | | | | | | | | | | | | | | | | | |
| Frequency | OD | Start Day | 7 | | | | | | | | | | | | | | | | | |
| Quantity Dispensed | | Dispensed by | | | | | | | | | | | | | | | | | | |
| | | Accuracy check | | | | | | | | | | | | | | | | | | |
| Note | SUBCUTANEOUS BOLUS To be injected ONCE a day by subcutaneous injection on days 7, 11 and 14. | | | | | | | | | | | | | | | | | | | |

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| Date: | Date: | Date: | Date: |