

Chemotherapy Protocol

GERM CELL

BLEOMYCIN-CISPLATIN-ETOPOSIDE

(BEP 5 Day)

Regimen

• Germ Cell – Bleomycin-Cisplatin-Etoposide (5 day BEP)

Indication

- In patients 40 years and below with;
 - metastatic non-seminomatous germ cell tumours
 - metastatic seminoma where radiotherapy is not appropriate
 - renal impairment or a poor performance status

Toxicity

| Drug | Adverse Effect |
|-----------|--|
| Bleomycin | Pulmonary toxicity, rigors, skin pigmentation, nail changes |
| Cisplatin | Neuropathy, nephrotoxicity, ototoxicity |
| Etoposide | Hypotension on rapid infusion, alopecia, hyperbilirubineamia |

The adverse effects listed are not exhaustive. Please refer to the relevant Summary of Product Characteristics for full details.

Monitoring

Drugs

- FBC, LFTs and U&Es on day one of the cycle
- AFP, HCG prior to day one of the cycle
- Chest x-ray
- Consider pulmonary function tests before starting therapy. These should be repeated
 if respiratory symptoms develop during treatment, particularly a drop in oxygen
 saturation on exercise. Bleomycin should be stopped until the results of such
 investigations are known.

Dose Modifications

The dose modifications listed are for haematological, liver and renal function and drug specific toxicities only. Dose adjustments may be necessary for other toxicities as well.

In principle all dose reductions due to adverse drug reactions should not be re-escalated in subsequent cycles without consultant approval. It is also a general rule for chemotherapy that if a third dose reduction is necessary treatment should be stopped.



Patients are being treated with curative intent therefore dose modifications and delays should be kept to a minimum. Please discuss all dose reductions / delays with the relevant consultant before prescribing. The approach may be different depending on the clinical circumstances.

Haematological

Consider a blood transfusion if the patient is symptomatic of anaemia or has a haemoglobin of less than 8g/dL.

Prior to each cycle the following criteria must be met

| Criteria | Eligible Level | | |
|------------|--|--|--|
| Neutrophil | equal to or more than 0.5x10 ⁹ /L | | |
| Platelets | equal to or more than 100x10 ⁹ /L | | |

This is a curative regimen. All dose reductions and delays should be discussed with the relevant consultant. In general if these levels are not met then treatment should be delayed for three days at a time. Treatment should re-start as soon as these haematological parameters are met. Dose delays rather than dose reductions are recommended.

Hepatic Impairment

| Drug | Bilirubin µmol/L | | AST/ALT units/L | Dose (% of original dose) | |
|-----------|---------------------|----|---------------------|--------------------------------|--|
| Bleomycin | | | | Clinical decision | |
| | | | | | |
| Cisplatin | N/A | | N/A | No dose modification necessary | |
| | | | | | |
| | 26-51 | or | 60-180 | Consider dose reducing to 50% | |
| Etoposide | greater than 51 | or | greater than 180 | Clinical decision | |



Renal Impairment

| Drug | Creatinine Clearance (ml/min) | Dose (% of original dose) | | | | |
|------------|--|----------------------------------|--|--|--|--|
| Bleomycin | 50 or more | 100% | | | | |
| Біеоттусіт | less than 50 | discuss with consultant and omit | | | | |
| | | | | | | |
| | 60 or greater | 100% | | | | |
| Cisplatin | If the creatinine clearance is 59ml/min or below please refer to the responsible consultant for advice | | | | | |
| | | | | | | |
| | greater than 50 | 100% | | | | |
| Etoposide | 15-50 | 75% | | | | |
| | less than 15 | 50% | | | | |

Other

Dose reductions or interruptions in therapy are not necessary for those toxicities that are considered unlikely to be serious or life threatening. For example, alopecia, altered taste or nail changes.

For all other non-haematological NCI-CTC grade 3 and above toxicities delay treatment until the adverse effect has resolved to NCI-CTC grade 1 or below. The dose of the causative agent(s) may then be reduced or discontinued at the discretion of the consultant.

Bleomycin

The risk of bleomycin induced pneumonitis is greater in those individuals who are older than forty years of age, have a history of smoking, those with underlying lung disease, previous mediastinal radiotherapy or poor renal function. If pulmonary symptoms develop stop the bleomycin until they can be investigated fully and a diagnosis made.

Regimen

Good prognosis - 3 cycles

Intermediate / Poor prognosis – 4 cycles (if 4 cycles are required omit the day 8, 15 bleomycin on cycle 4 only)

3 cycles will be set in Aria



21 day cycle for 3 cycles

| Drug | Dose | Days | Administration |
|-----------|----------------------|-----------|--|
| Bleomycin | 30,000 IU | 2, 8, 15 | Intravenous infusion in 100ml sodium chloride |
| Distance | 00,000 10 | | 0.9% over 30 minutes |
| Cisplatin | 20mg/m ² | 1,2,3,4,5 | Intravenous infusion in 1000ml sodium chloride 0.9% with 20mmol potassium chloride over 60 minutes |
| Etoposide | 100mg/m ² | 1,2,3,4,5 | Intravenous infusion in 1000ml sodium chloride 0.9% over 60 minutes |

Dose Information

- Aria is set to dose cap all regimens at 2.4m². This regimen must NOT be capped.
 Please override any doses that are capped.
- Cisplatin will be dose banded according to the agreed bands
- Etoposide will be dose banded according to the agreed bands

Administration Information

Extravasation

- Bleomycin neutral
- Cisplatin exfoliant
- Etoposide irritant

Additional Therapy

Antiemetics

15 – 30 minutes prior to chemotherapy

- aprepitant 125mg once a day on day 1
- aprepitant 80mg once a day on days 2, 3
- dexamethasone 4mg once a day on days 1, 2, 3, 4, 5, 6, 7 oral
- metoclopramide 10mg three times a day when required oral
- ondansetron 8mg twice a day on days 1, 2, 3, 4, 5, 6, 7 oral
- On days of bleomycin administration
 - hydrocortisone 100mg intravenous when required
 - chlorphenamine 10mg intravenous when required
- Cisplatin pre-hydration as follows
 - furosemide 40mg oral or intravenous as required
 - sodium chloride 0.9% 1000ml with 8mmol magnesium sulphate over 60 minutes



- Cisplatin post hydration
 - sodium chloride 0.9% 1000ml over 240 minutes
- Ciprofloxacin 500mg twice a day for 7 days starting on day 8 of the cycle
- Consider growth factor support according to local policy, for example;
 - filgrastim or bioequivalent 30 million units once a day for seven days starting on day seven of the cycle subcutaneous
 - lenograstim or bioequivalent 33.6 million units once a day for seven days starting on day seven of the cycle subcutaneous
 - pegfilgrastim or bioequivalent 6mg once a day for one day on day seven of the cycle
- Mouthwashes according to local or national policy on the treatment of mucositis
- Gastric protection with a proton pump inhibitor or a H₂ antagonist may be considered in patients considered at high risk of GI ulceration or bleed.

Coding (OPCS)

- Procurement X70.3
- Delivery N/A

References

1.de Wit R, Roberts JT, Wilkinson PM et al. Equivalence of 3 cycle BEP versus 4 cycles and of the 5 day schedule versus 3 days per cycle in good-prognosis germ cell cancer: a randomised study of the European Organisation for Research and Treatment of Cancer Genitourinary Tract Cancer Cooperative Group and the Medical Research Council. J Clin Oncol 2001; 19: 1629-1640.

2.de Wit R, Stoter G, et al. Four cycles of BEP versus four cycles of VIP in patients with intermediate-prognosis metastatic testicular non seminoma: A randomised study of the EORTC Genitourinary Tract Cancer Cooperative Group. Br J Cancer 1998; 78(6): 828-832.

3.Nichols C, Catalano P, Crawford E et al. Randomised comparison of cisplatin and etoposide and either bleomycin or ifosfamide in the treatment of advanced disseminated germ cell tumours: An Eastern Cooperative Oncology Group, Southwest Oncology Group and Cancer and Leukemia Group B study. J Clin Oncol 1998; 16: 1287-1293.

5.Fossa SD, Kaye SB, Mead GM, Cullen MH, De Wit R, Borogi J, Van Groeningen C, De Mulder P, Stenning S and De Prijck L. Filgrastim (G-CSF) during combination chemotherapy of patients with poor prognosis metastatic germ cell malignancy (A phase III trial of the EORTC GU group/MRC testicular tumour working party) J Clin Oncol 1998: 16: 716-724



REGIMEN SUMMARY

Bleomycin-Cisplatin-Etoposide (5 day BEP)

Cycle 1, 2, 3, 4

Day 1

- 1. Aprepitant 125mg oral
- 2. Dexamethasone 4mg oral or intravenous
- 3. Metoclopramide 10mg oral or intravenous
- 4. Ondansetron 8mg oral or intravenous bolus
- 5. Furosemide 40mg oral or intravenous when required for the maintenance of diuresis
- 6. Sodium chloride 0.9% 1000ml with magnesium sulphate 8mmol intravenous infusion over 60 minutes
- 7. Cisplatin 20mg/m² in 1000ml sodium chloride 0.9% with 20mmol potassium chloride intravenous infusion over 60 minutes
- 8. Sodium chloride 0.9% 1000ml over 240 minutes
- 9. Etoposide 100mg/m² in 1000ml sodium chloride 0.9% intravenous infusion over 60 minutes

Take Home Medicines

- 10. Dexamethasone 4mg once a day in the morning for 2 days starting on day 6 of the cycle oral
- 11. Metoclopramide 10mg up to three times a day when required for the relief of nausea oral
- 12. Ondansetron 8mg to be taken on the evening of days 1, 2, 3, 4 and 5 of chemotherapy and 8mg twice a day for 2 days starting on day 6 of the cycle oral
- 13. Ciprofloxacin 500mg twice a day for 7 days starting on day 8 of the cycle oral

Day 2

- 14. Aprepitant 80mg oral
- 15. Dexamethasone 4mg oral or intravenous
- 16. Metoclopramide 10mg oral or intravenous
- 17. Ondansetron 8mg oral or intravenous bolus
- 18. Furosemide 40mg oral or intravenous when required for the maintenance of diuresis



- 19. Sodium chloride 0.9% 1000ml with magnesium sulphate 8mmol intravenous infusion over 60 minutes
- 20. Cisplatin 20mg/m² in 1000ml sodium chloride 0.9% with 20mmol potassium chloride intravenous infusion over 60 minutes
- 21. Sodium chloride 0.9% 1000ml over 30 minutes
- 22. Etoposide 100mg/m² in 1000ml sodium chloride 0.9% intravenous infusion over 60 minutes
- 23. Bleomycin 30,000 IU in 100ml sodium chloride 0.9% intravenous infusion over 30 minutes
- 24. Chlorphenamine 10mg intravenous when required for bleomycin reactions
- 25. Hydrocortisone 100mg intravenous when required for bleomycin reactions

Days 3

- 26. Aprepitant 80mg oral
- 27. Dexamethasone 4mg oral or intravenous
- 28. Metoclopramide 10mg oral or intravenous
- 29. Ondansetron 8mg oral or intravenous bolus
- 30. Furosemide 40mg oral or intravenous when required for the maintenance of diuresis
- 31. Sodium chloride 0.9% 1000ml with magnesium sulphate 8mmol intravenous infusion over 60 minutes
- 32. Cisplatin 20mg/m² in 1000ml sodium chloride 0.9% with 20mmol potassium chloride intravenous infusion over 60 minutes
- 33. Sodium chloride 0.9% 1000ml over 240 minutes
- 34. Etoposide 100mg/m² in 1000ml sodium chloride 0.9% intravenous infusion over 60 minutes

Days 4, 5

- 35. Dexamethasone 4mg oral or intravenous
- 36. Metoclopramide 10mg oral or intravenous
- 37. Ondansetron 8mg oral or intravenous bolus
- 38. Furosemide 40mg oral or intravenous when required for the maintenance of diuresis
- 39. Sodium chloride 0.9% 1000ml with magnesium sulphate 8mmol intravenous infusion over 60 minutes



- 40. Cisplatin 20mg/m² in 1000ml sodium chloride 0.9% with 20mmol potassium chloride intravenous infusion over 60 minutes
- 41. Sodium chloride 0.9% 1000ml over 240 minutes
- 42. Etoposide 100mg/m² in 1000ml sodium chloride 0.9% intravenous infusion over 60 minutes

Day 8, 15

- 43. Dexamethasone 8mg oral or intravenous
- 44. Bleomycin 30,000 IU in 100ml sodium chloride 0.9% intravenous infusion over 30 minutes
- 45. Chlorphenamine 10mg intravenous when required
- 46. Hydrocortisone 100mg intravenous when required



DOCUMENT CONTROL

| Ver | sion | Date | Amendment | Written By | Approved By |
|-----|------|-----------|-----------|---------------------------------|---|
| | 1 | July 2017 | None | Dr Deborah Wright Pharmacist | Dr Emma Killick Consultant Medical Oncologist |

This chemotherapy protocol has been developed as part of the chemotherapy electronic prescribing project. This was and remains a collaborative project that originated from the former CSCCN. These documents have been approved on behalf of the following Trusts;

Hampshire Hospitals NHS Foundation Trust NHS Isle of Wight Portsmouth Hospitals NHS Trust Salisbury Hospital NHS Foundation Trust University Hospital Southampton NHS Foundation Trust Western Sussex Hospitals NHS Foundation Trust

All actions have been taken to ensure these protocols are correct. However, no responsibility can be taken for errors which occur as a result of following these guidelines.