



**Dose modification for hepatic toxicity**

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|---|-----------------|
| • AST/ALT up to 5 x ULN, Bilirubin < 1.5xULN  | Full dose       |
| • AST/ALT 5.1-20 x ULN, Bilirubin 1.5-3 x ULN | Defer by 1 week |
| • AST/ALT > 20xULN, Bilirubin > 3ULN          | Stop treatment  |

**Dose modification for neurological toxicity**

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|-----------------|--|
| • NCI grade 0-1 | Proceed with full dose   |
| • NCI grade 2   | Defer until recovery, then replace Cisplatin with Carboplatin AUC5 |
| • NCI grade 3+  | Change to less neurotoxic regime if appropriate                    |

**Expected toxicities****Neutropenic sepsis & thrombocytopenia****Nausea & vomiting (severe)****Constipation****Peripheral neuropathy****Alopecia****Mucositis****Tinnitus**

**This protocol has been reviewed by the Lancashire & South Cumbria Lung Oncology Consultants' Group and responsibility for the template lies with the Head of Service.**

**Date: September 2019**

**Next review: September 2021**