

CARBOPLATIN desensitisation

INDICATION (ICD10)

1. Carboplatin desensitisation for carboplatin hypersensitivity.
PS 0, 1, 2

REGIMEN

| | | | | |
|-------|--|-----------|-------------|---------------------------------|
| Day 0 | Premedication starting the night prior to infusion: Dexamethasone 8mg twice daily orally Cetirizine 10mg once daily orally H ₂ antagonist once daily orally | | | |
| Day 1 | Premedication 30 minutes prior to infusion: Dexamethasone 8mg IV bolus Chlorphenamine 10mg IV bolus Ondansetron 8mg IV bolus H ₂ antagonist | | | |
| | CARBOPLATIN | AUC 0.005 | IV infusion | 50ml glucose 5% over 30 minutes |
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| | CARBOPLATIN | AUC 4.445 | IV infusion | #ml glucose 5% over 60 minutes |

Dose calculated by EDTA GFR or calculated $(CrCl + 25) \times AUC$.

diluent volume for dose prescribed as per national standardised product specification

CYCLE FREQUENCY AND NUMBER OF CYCLES

Every 21 days for the equivalent number of cycles would have received in treatment reacted to, up to 6 cycles

ANTI-EMETICS

Moderate risk day 1

CONCURRENT MEDICATION REQUIRED

| | |
|-------------|--|
| Carboplatin | Ensure premedication given before carboplatin. Anaphylaxis treatment should be prescribed if the patient has had an anaphylactic episode previously. Dexamethasone 20mg IV bolus Chlorphenamine 10mg IV bolus H ₂ antagonist Carboplatin should be given at a slower rate e.g 2-4 hours. |
|-------------|--|

EXTRAVASATION AND TYPE OF LINE / FILTERS

Carboplatin - irritant

Filter not required

Peripheral or central line

INVESTIGATIONS

Blood results required before SACT administration

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| FBC, U&E including Mg ⁺⁺ (>0.4) and LFTs Neutrophils x 10 ⁹ /L ≥1.0 provided patient is well Platelets ≥100x10 ⁹ /L | baseline and every cycle |
| GFR assessed using EDTA result (BMI <19 or >30 or calculated creatinine clearance at the Consultant's discretion) | baseline and every cycle |
| Patients with hydronephrosis or serum creatinine ≥100micromol/L | baseline and every cycle |
| CA125 | baseline and day 1 every cycle |
| Virology | before cycle 1 if not previously checked |
| Weight | baseline and every cycle |

MAIN TOXICITIES AND ADVERSE REACTIONS

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|-------------|--|
| Carboplatin | Ototoxicity – monitor Neurotoxicity - monitor |
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DOSE MODIFICATIONS

Hepatic impairment

Carboplatin

No need for dose adjustment is expected

Renal impairment

Carboplatin

| | |
|---|-----------------|
| GFR/ calculated CrCl ≤20ml/min or ≤30ml/min with pre-existing severe renal impairment | contraindicated |
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REFERENCES

- Royal Marsden desensitisation regimen

Assessments

| | Pre | Cycle 1 | Cycle 2 | Cycle 3 | Cycle 4 | Ongoing |
|-------------------------------------|-----|---------|-----------|---------|-----------|--|
| Clinical assessment | X | | Pre cycle | | Pre cycle | Every cycle |
| SACT assessment (PS and toxicities) | X | X | X | X | X | Every cycle |
| FBC | X | X | X | X | X | Every cycle |
| U&E, calcium, magnesium & LFT | X | X | X | X | X | Every cycle |
| CrCl | X | X | X | X | X | Every cycle |
| CT scan | X | | | | | At cycle 6, Inform consultant team if not booked |
| Informed consent | X | | | | | Verbal each cycle |
| Height | X | | | | | |
| Weight recorded | X | X | X | X | X | Every cycle |