

Weekly cisplatin with concurrent radiotherapy for head and neck cancer

Indication

Locally advanced head and neck cancer

Regimen details

DRUG	FLUID	TIME
20mmol Potassium chloride + 10mmol magnesium sulphate	1 litre 0.9% sodium chloride	2 hours
Cisplatin 40 mg/m ²	1000ml 0.9% sodium chloride	1 hour
	500ml 0.9% sodium chloride	30 minutes

(Cap body surface area at 2m²)

Cycle frequency

Repeat weekly concurrently with radiotherapy

Number of cycles

6 weeks

Administration

Radiotherapy should be given during the post-hydration period

Ideally chemotherapy should be administered on Monday, Tuesday or Wednesday

Emetogenicity

Highly emetogenic

Investigations – pre first cycle

Investigation	Validity period
FBC	14 days
U+E (including creatinine)	14 days
LFT (including AST)	14 days
Magnesium	14 days

Investigations –pre subsequent cycles

FBC, U+E (including creatinine), LFT (including AST), Magnesium

Standard limits for administration to go ahead

If blood results not within range, authorisation to administer **must** be given by prescriber/ consultant.

At Consultant's direction

Defer treatment 1 week until neutrophils ≥ 1.5 and platelets ≥ 100

If neutrophils 1.2-1.5 contact consultant

Withhold treatment if calculated Creatinine Clearance is < 50 ml/min. Repeat before further treatments.

Dose modifications

Consider discontinuing concurrent cisplatin if acute kidney injury

Consider changing to carboplatin AUC 2 if persistent tinnitus or deafness or if calculated creatinine clearance is <50

Discuss with consultant before modifying dose

Adverse effects

Nausea and vomiting

Renal impairment, tinnitus, hearing loss, neuropathy

THIS PROTOCOL HAS BEEN DIRECTED BY DR MIRZA, CLINICIAN FOR HEAD AND NECK CANCER

RESPONSIBILITY FOR THIS PROTOCOL LIES WITH THE HEAD OF SERVICE

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