

<b>Indication</b>	Head and neck: Malignant salivary gland tumours
<b>Treatment Intent</b>	Palliative
<b>Frequency and number of cycles</b>	Every 21 days for 6 cycles
<b>Monitoring Parameters pre-treatment</b>	<ul style="list-style-type: none"> <li>• ECG must be checked prior to cycle 1.</li> <li>• C+G should be used to measure CrCl prior to cycle 1</li> <li>• If CrCl &lt;60ml/min then obtain EDTA result.</li> <li>• If GFR 30-59ml/min consider dose reduction of cisplatin or substitute with carboplatin, if GFR &lt;30ml/min stop platinum.</li> <li>• Monitor LFT's and serum creatinine at each cycle.</li> <li>• <u>Day 1</u> If neuts 1.0-1.4 and Plts <math>\geq</math>100 d/w consultant. If neuts &lt;1.0 or Plts &lt;100 delay epirubicin and cisplatin one week</li> <li>• <u>Day 8 + 15</u> continue 5FU provided neuts <math>\geq</math>0.5 and PLT <math>\geq</math>75.</li> <li>• Dose reduction should be considered if grade 3 or 4 non-haematological toxicity or repeat appearance of grade 2 (except N&amp;V and alopecia). Delay until resolution of toxicity to <math>\leq</math> grade 1</li> <li>• Consider audiology test for hearing impaired patients and monitor all patients for ototoxicity throughout treatment.</li> <li>• <b>DPD testing:</b> DPD testing must be undertaken in all patients before starting treatment; the result must be checked before treatment is started.</li> <li>• <b>Cardiotoxicity:</b> Caution in patients with prior history of coronary heart disease, arrhythmias and angina pectoris.</li> <li>• Maximum recommended cumulative dose epirubicin 900mg/m<sup>2</sup>.</li> <li>• <b>Drug interactions (for comprehensive list refer to BNF/SPC):</b></li> <li>• Caution, ciclosporin increases concentration of epirubicin.</li> <li>• In patients receiving cisplatin and phenytoin, the serum level of phenytoin might be reduced, monitor phenytoin levels with concomitant use.</li> </ul>
<b>References</b>	KMCC protocol MULTI-011 v1 SPC epirubicin accessed online 17.06.21 SPC cisplatin accessed online 17.06.21

NB For funding information, refer to the CDF and NICE Drugs Funding List

Protocol No	MULTI-011	Kent and Medway SACT Protocol Disclaimer: No responsibility will be accepted for the accuracy of this information when used elsewhere.	
Version	V2	Written by	M.Archer
Supersedes version	V1	Checked by	B Willis / C Waters
Date	23.08.21	Authorising consultant (usually NOG Chair)	K Nathan

## Repeat every 21 days

Day	Drug	Dose	Route	Infusion Duration	Administration
Day 1	Sodium chloride 0.9%	1000ml	IV	2 hours	+ 20mmol KCL + 10mmol Mg <sup>2+</sup>
	Mannitol 10%	200ml	IV	15 min	
	Ondansetron	<75yrs 16mg ≥75yrs 8mg	IV	15 min	Sodium Chloride 0.9% 50ml
	Dexamethasone	8mg	PO		
	<b>EPIRUBICIN</b>	<b>50mg/m<sup>2</sup></b>	IV	3 min	through the side of a fast running 0.9% Sodium chloride intravenous infusion
	<b>CISPLATIN</b>	<b>60mg/m<sup>2</sup></b>	IV	2 hours	In 1000ml Sodium chloride 0.9%
	Furosemide	40mg	IV/PO	bolus	Only if urine output <100ml/hour or weight gain >1kg
	Sodium Chloride 0.9%	1000ml	IV	2 hours	+ 20mmol KCL + 10mmol Mg <sup>2+</sup>
	Sodium Chloride 0.9%	500ml	IV	1 hour	or 500ml water, orally
	*(Furosemide)	40mg	IV/PO	<b>*only if required</b>	If patient remains in a 2L positive balance
	<b>5-FLUOROURACIL</b> prescribe for a total of 7 days	<b>200mg/m<sup>2</sup>/ day</b> <b>ie 1400mg/m<sup>2</sup>/7 days</b>	IV	7 days	Continuous infusion pump
Day 8	<b>5-FLUOROURACIL</b> prescribe for a total of 7 days	<b>200mg/m<sup>2</sup>/ day</b> <b>ie 1400mg/m<sup>2</sup>/7 days</b>	IV	7 days	Continuous infusion pump
Day 15	<b>5-FLUOROURACIL</b> prescribe for a total of 7 days	<b>200mg/m<sup>2</sup>/ day</b> <b>ie 1400mg/m<sup>2</sup>/7 days</b>	IV	7 days	Continuous infusion pump
TTO	Drug	Dose	Route	Directions	
Day 1	Dexamethasone	6mg	PO	OM for 3 days	
	Metoclopramide	10mg	PO	3 times a day for 3 days, then 10mg up to 3 times a day as required. Do not take for more than 5 days continuously.	

Protocol No	MULTI-011	Kent and Medway SACT Protocol Disclaimer: No responsibility will be accepted for the accuracy of this information when used elsewhere.		
Version	V2	Written by		M.Archer
Supersedes version	V1	Checked by		B Willis / C Waters
Date	23.08.21	Authorising consultant (usually NOG Chair)		K Nathan