Indication	Metastatic prostate cancer (hormone naïve) NB Within this indication, docetaxel is commissioned for use as follows: in men either commencing, or who have commenced within 12 weeks, long-term ADT for metastatic disease for the first time; and men of sufficient performance status to be treated with 6 cycles of docetaxel chemotherapy.				
Treatment Intent	Palliative				
Frequency and number of cycles	Repeat every 21 days for up to 6 cycles				
Monitoring Parameters pre-treatment	 Monitor FBC, U&Es and LFT's at each cycle. If neuts ≥ 1.5 and PLT ≥100 continue with treatment. If neuts 1.0-1.4 and Plts ≥100 d/w consultant. If neuts <1.0 or Plts <100 delay one week and consider dose reduction. Hepatic Impairment: Consider dose reduction in liver impairment. Not recommended in severe liver impairment. Renal Impairment: no dose adjustment required. Dose Modification: Dose reduction should be considered if grade 3 or 4 non-haematological toxicity or repeat appearance of grade 2 (except N&V and alopecia). Delay until resolution of toxicity to ≤ grade 1 Prednisolone: Dose may be omitted on the days of dexamethasone pre-medication. Dose may be adjusted at clinician's discretion. Common drug interactions: (for comprehensive list refer to BNF/SPC) Concomitant use with medicines which induce, inhibit or are metabolised by cyto-chrome P450-3A (eg ciclosporin, ketoconazole and erythromycin) may affect levels of docetaxel use with caution. Avoid concomitant use with strong CYP3A4 inhibitors (eg ketoconazole, itraconazole, clarithromycin and ritonavir), if treatment cannot be avoided consider dose reduction of docetaxel and monitor patient closely for signs of toxicity. Ensure dexamethasone pre-medication (8mg PO BD for 3 days, starting the morning of the day prior to the next cycle of docetaxel) is prescribed and given to the patient 				
References	at new patient chatKMCC proforma URO-026 v3 SPC accessed online27/02/20 BNF accessed online 27/02/20				

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

Protocol No	URO-026	Kent and Medway SACT Protocol Disclaimer: No responsibility will be accepted for the accuracy of this information when used elsewhere.		
Version	V4	Written by	M.Archer	
Supersedes	V3	Checked by	C.Waters	
version			M.Capomir	
Date	29.06.01	Authorising consultant (usually NOG Chair)	A.Edwards	

Repeat every 21 days

Day	Drug	Dose	Route	Infusion Duration	Administration	
Day 1	Metoclopramide	20mg	IV			
	DOCETAXEL	75mg/m²	IV	1 hour	Sodium Chloride 0.9% 250ml	
TTO	Drug	Dose	Route	Direction	Directions	
	PREDNSIOLONE	5mg	PO	BD continuously (dispense 3 weeks supply) Take with or just after food, or a meal. On the final cycle of treatment, the patient should commence a reducing prednisolone dose: 5mg BD for 1 week and then 5mg OD for 2 weeks. BD for 3 days, starting the morning of the day prior to the next cycle of docetaxel. Take with or just after food, or a meal. Do not dispense on last cycle.		
	Dexamethasone	8mg	РО			
	Metoclopramide	10mg	PO	Up to 3 times a day when required (Maximum of 30mg per day including 20mg pre-chemo dose). Do not take for more than 5 days continuously		

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