NCT Number	Title	Status	Results	Interventions	Pharmacogenomic analyses	Phase	Participants	Study Type
NCT00598858	Neoadjuvant Docetaxel on	Withdrawn (lack of	No	docetaxel +	Genomic factors affecting	Π	0	Interventional
	Newly Diagnosed	funding)		prednisone	responses to chemotherapy.			
	Intermediate and High							
	Grade Cancer of the							
	Prostate							
NCT01087580	Docetaxel + Prednisone	Withdrawn	No	docetaxel +	Identification of genes important	II	0	Interventional
	With or Without Radiation			prednisone or	in governing the response to			
	for Castrate Resistant			docetaxel +	chemotherapy and radiation			
	Prostate Cancer			prednisone +	therapy. Expression and mutation			
				radiotherapy	of p53 and AR.			
NCT02793219	Provenge Followed by	Withdrawn (company	No	sipuleucel-T +	Association of immunological	II	0	Interventional
	Docetaxel in Castration-	decided not to fund)		docetaxel	biomarkers with clinical results			
	Resistant Prostate Cancer				for therapy. Expression of CD3,			
					CD4, CD8, CD25/FOX3P,			
					CD56, CTLA-4, PD-1, and Ki67			
					in prostate cancer infiltrates.			
NCT02793765	Docetaxel Followed by	Withdrawn (company	No	docetaxel +	Association of immunological	II	0	Interventional
	Provenge in Metastatic	decided not to fund)		sipuleucel-T	biomarkers with clinical results			
	Prostate Cancer				for therapy. Expression of CD3,			
					CD4, CD8, CD25/FOX3P,			
					CD56, CTLA-4, PD-1, and Ki67			
					in prostate cancer infiltrates.			

Supplementary table 1. Withdrawn clinical trials and clinical trials with unknown status for docetaxel treatment in prostate cancer (ClinicalTrials.gov)

NCT00104715	Hormone Therapy and	Unknown status	No	hormonal therapy +	Tumor profiles of gene	III	378	Interventional
	Docetaxel or Hormone			docetaxel or	expression		(estimated)	
	Therapy Alone in Treating			hormonal therapy				
	Patients With Metastatic			alone				
	Prostate Cancer							
NCT02208583	Molecular Phenotype	Unknown status	No	docetaxel +	Molecular phenotypic changes	NA	150	Interventional
	Changes and Personalized			prednisone + targeted	after acquired ADT resistance		(estimated)	
	Treatment for CRPC			drugs or cisplatin +	and their effect on OS (AR,Ki-			
				etoposide + targeted	67, CD56, Syn, P53, AURKA,			
				drugs	N-myc, retinoblastoma			
					susceptibility, E-cadherin,			
					vimentin, hotspot mutation for 48			
					cancer related genes)			

AR, androgen receptor ADT, androgen deprivation therapy OS, overall survival