

Table I Patient characteristics

	No	Age (y)	sex	Location	Pathology	stage(TNM)	IRS-Group	<u>registered clinical study group</u>
Group non-Cx	1	0.6	M	Bladder	embryonal	<u>2</u>	III	JRSG
	2	4.4	M	Prostate	embryonal	<u>2</u>	III	JRSG
Group Cx	3	0.9	M	Prostate	unknown	<u>3</u>	III	non
	4	2.7	F	Bladder	embryonal	<u>3</u>	III	non
	5	2.8	M	Prostate	embryonal	<u>3</u>	III	JRSG

Abbreviations; IRS: intergroup rhabdomyosarcoma study group, M: male, F: female, JRSG: Japan Rhabdomyosarcoma Study Group

Table2

Table II Treatment summary and patient outcome

Pt no	Cx	PBT						<u>Radiotherapy other than PBT</u>	<u>HDC ±</u>	Surgery	Outcome	Months to last f/u	Complications at last visit	
		GyE (Fr)	Combined Cx	Lowest WBC (/uL)	BTF	Adverse event	PBT							
1	<u>1st: VAC</u>	41.4	none	2,100	none	none	<u>No</u>	No	+	1CR	37	none	RCP	
	<u>2nd: VID</u>	(23)											SC*	
Group non-Cx	<u>1st: VAC</u>	41.4	none	2,400	none	Catheter related	<u>Yes:</u>	Yes;	Tepa 960mg/m2	+	1CR	93	hydronephrosis**	RCP
	<u>2nd: irinotecan</u>	(23)				infection	<u>local</u>							CUR
	<u>3rd: CDDP-based</u>													
214	3	<u>1st: VAC</u>	50.4	VC	500	RBC	dermatitis	<u>No</u>	No	total	1CR	10	none	Gross resection
		(28)												
	Group Cx	<u>1st: VAC</u>	45	VAC	700	none	Cystitis, vaginal fistula	<u>No</u>	No	+	1CR	56	mild hydronephrosis	RC
		<u>2nd: CDDP-based</u>					CUR							
		<u>3rd: irinotecan+VCR</u>												
	5	<u>4th: reduced VAC</u>	50.4	VC	100	RBC	Local pain with phentanyl	<u>No</u>	No	Biopsy	VGPR ⇒SD	11	none	
		<u>1st: VAC</u>												
		<u>2nd: VDC/IE</u>												
		<u>3rd: back to VAC</u>												

Abbreviations: Pt no: patients' number, BTF: Blood Transfusion, HST: hematopoietic stem cell transfusion, CR: complete response, VGPR: very good partial response, SD: stable disease, RCP: radical cystoprostatectomy, SC: sigmoid colon conduit, CUR: continent urinary reservoir, RC: radical cystectomy, Abbreviations: Pt no: patients' number, PBT: proton beam therapy, Cx: chemotherapy, WBC: white blood cell count, VAC: Consists of

Vincristine 1.5mg/m² day 0,7,14, Actinomycine-D 0.045 mg/kg day 0 and Cyclophosphamide 2,200 mg/m² day1, VC: Consists of Vincristine 1.5mg/m² day 0,7,14 and Cyclophosphamide 2,200 mg/m² day1, VID: Consists of VCR 1.0 mg/m² + IFO 1,200 mg/m² and DOX 25 mg/m², VDS/IE: Consists of Vincristine 1.5 mg/m² day0, Doxorubicin 37.5mg/m² day0,1, Cyclophosphamide 1,200 mg/m² day0, Ifosfamide 1,800 mg/m² days 14 through 18 and Etoposide 100mg/m² days14 through 18.

*Diversion to continent urinary reservoir being under consideration. **Due to ureterointestinal anastomosis stricture.

