

# 日本人の去勢抵抗性前立腺癌患者に対するPSMA標的の内用療法の治療経験

Experience of PSMA radioligand therapy in Japanese patients with metastatic castration-resistant prostate cancer

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**Purpose:** We have partnered with a medical team in Australia to perform <sup>177</sup>Lu-/<sup>225</sup>Ac-PSMA radioligand therapy (RLT) for Japanese castration-resistant prostate cancer (CRPC) patients overseas. In this article, we report our initial experience with <sup>177</sup>Lu-/<sup>225</sup>Ac-PSMA RLT in Japanese patients and discuss its efficacy and indications.

**Conclusions:** Although PSMA RLT is expected to be highly effective in Japanese CRPC patients, its efficacy in advanced cancers with already high PSA levels may be limited. Further investigation is warranted.

## Results:

### Characteristics (n = 26), March 2018 – March 2020

	Median or N	Range
Age (years)	68	50 - 79
Gleason score	9	6 - 10
Time to induction (months)	40.9	5.8 - 150.3
PSA (ng/ml), initial	42.8	3.4 - 7504.0
PSA (ng/ml), at induction	69.4	0.0 - 3052.3
WBC (/μL), mean	6664	2960 - 13990
Hb (g/μL), mean	11.5	7.8 - 14.4
Plt (x10 <sup>4</sup> /μL), mean	22.6	12.5 - 37.1
ALP (U/L), mean	421	138 - 2644
LDH (U/L), mean	305	107 - 1379
Cr (mg/dL), mean	0.8	0.6 - 1.3
<b>ECOG performance status</b>		
	0	15 (57.7%)
	1	10 (38.5%)
	2	1 (3.8%)
<b>Metastases</b>		
Bone	26 (100%)	
Lymph nodes	19 (73.1%)	
Visceral	4 (15.4%)	
<b>Prior treatment</b>		
<b>For prostate</b>		
RP	3 (1.2%)	
Irradiation	11 (42.3%)	
Others	1 (3.8%)	
No	11 (42.3%)	
<b>General</b>		
Hormone	26 (100%)	
Chemo	17 (65.4%)	

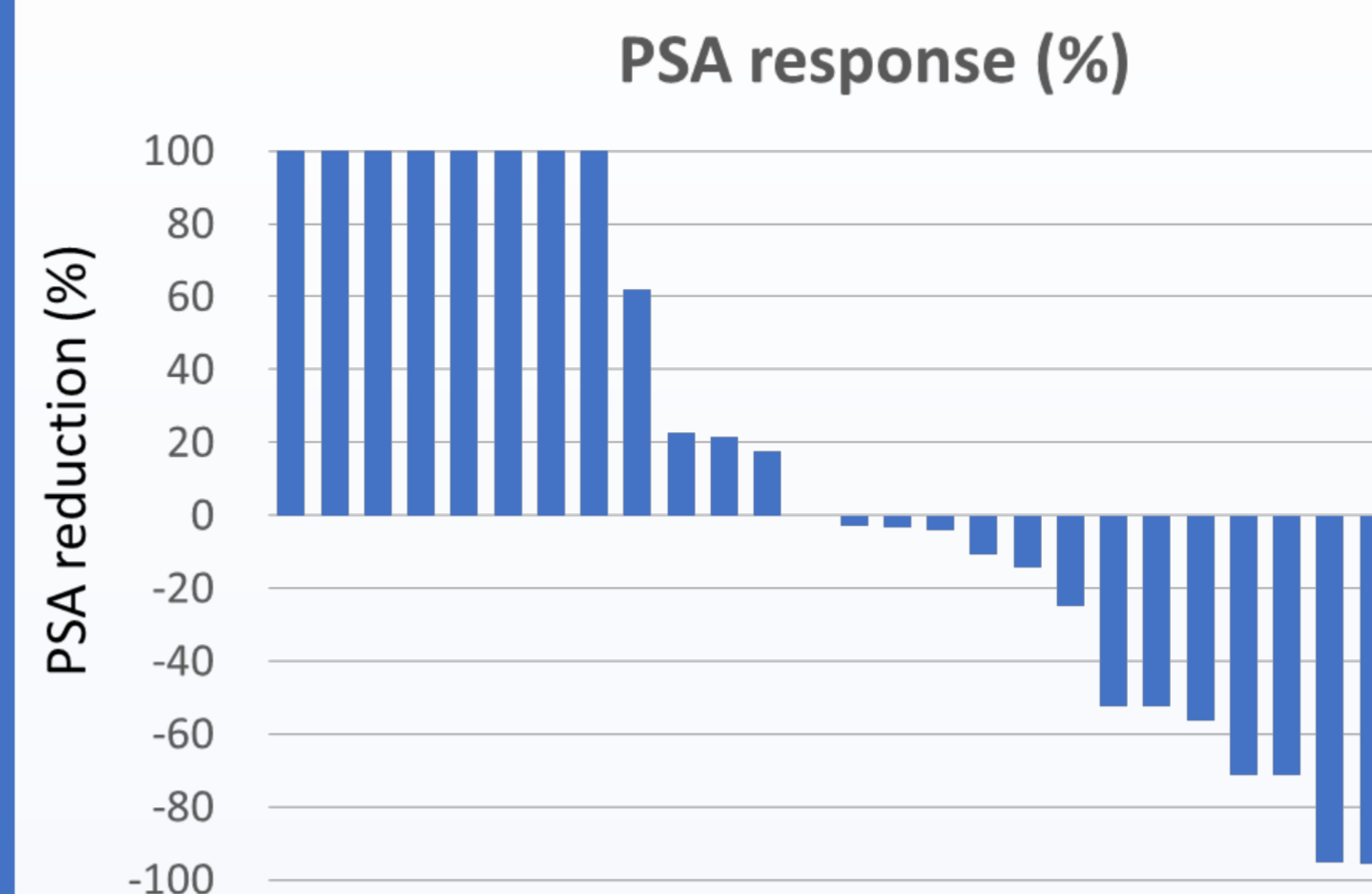
### Overview (n = 26)

	Mean or N	Range
No. of cycle	2.0	1 - 7
<b>Treatment nuclei</b>		
<sup>177</sup> Lu only	17 (65.4%)	
<sup>225</sup> Ac only	3 (11.5%)	
<sup>177</sup> Lu + <sup>225</sup> Ac	6 (23.1%)	
<b>Target</b>		
PSMA only	25 (96.1%)	
Dotatate only	0 (0%)	
PSMA + Dotatate	1 (3.8%)	

## Results:

### PSA response\* (n = 26)

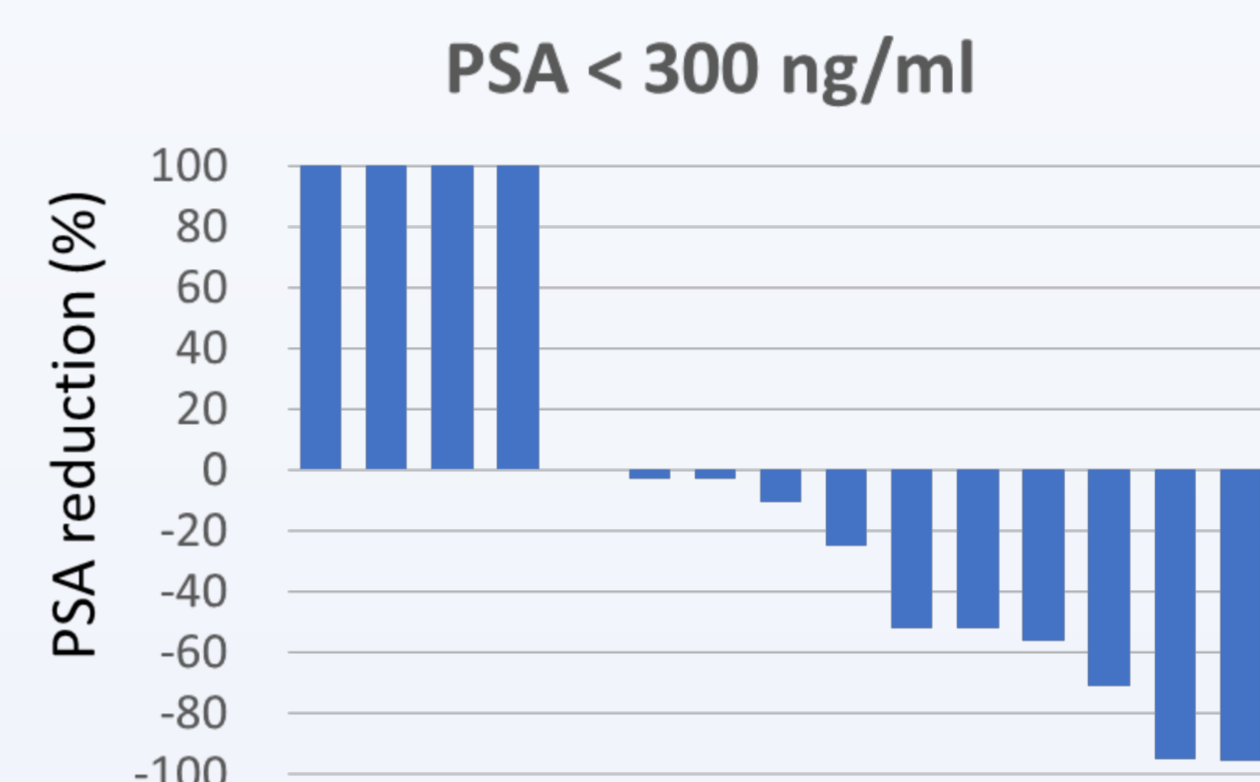
\*PSA at 7 weeks after last PSMA RLT



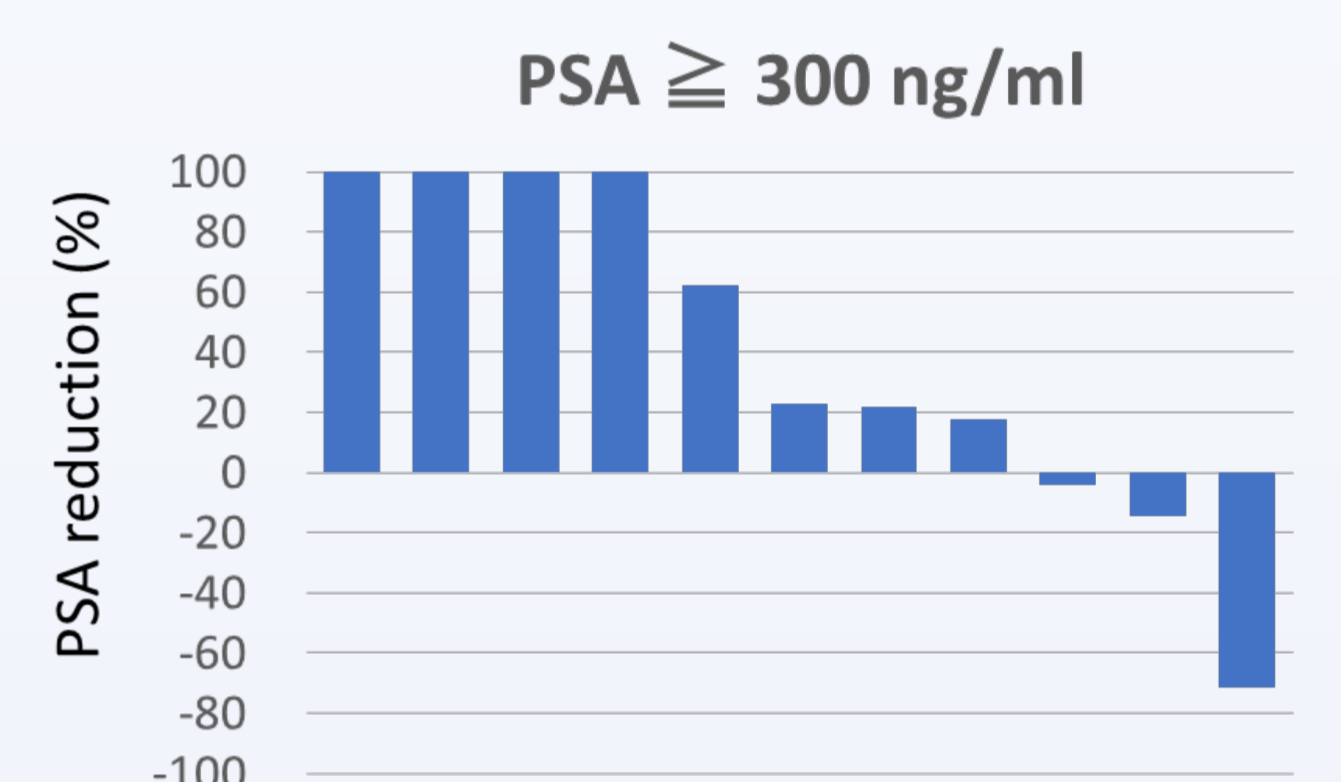
PSA progression	12 (46.1%)
Stable	1 (3.8%)
PSA reduction	13 (50.0%)
50% PSA reduction	7 (26.9%)

### PSA response\* in PSA group

\*PSA at 7 weeks after last PSMA RLT



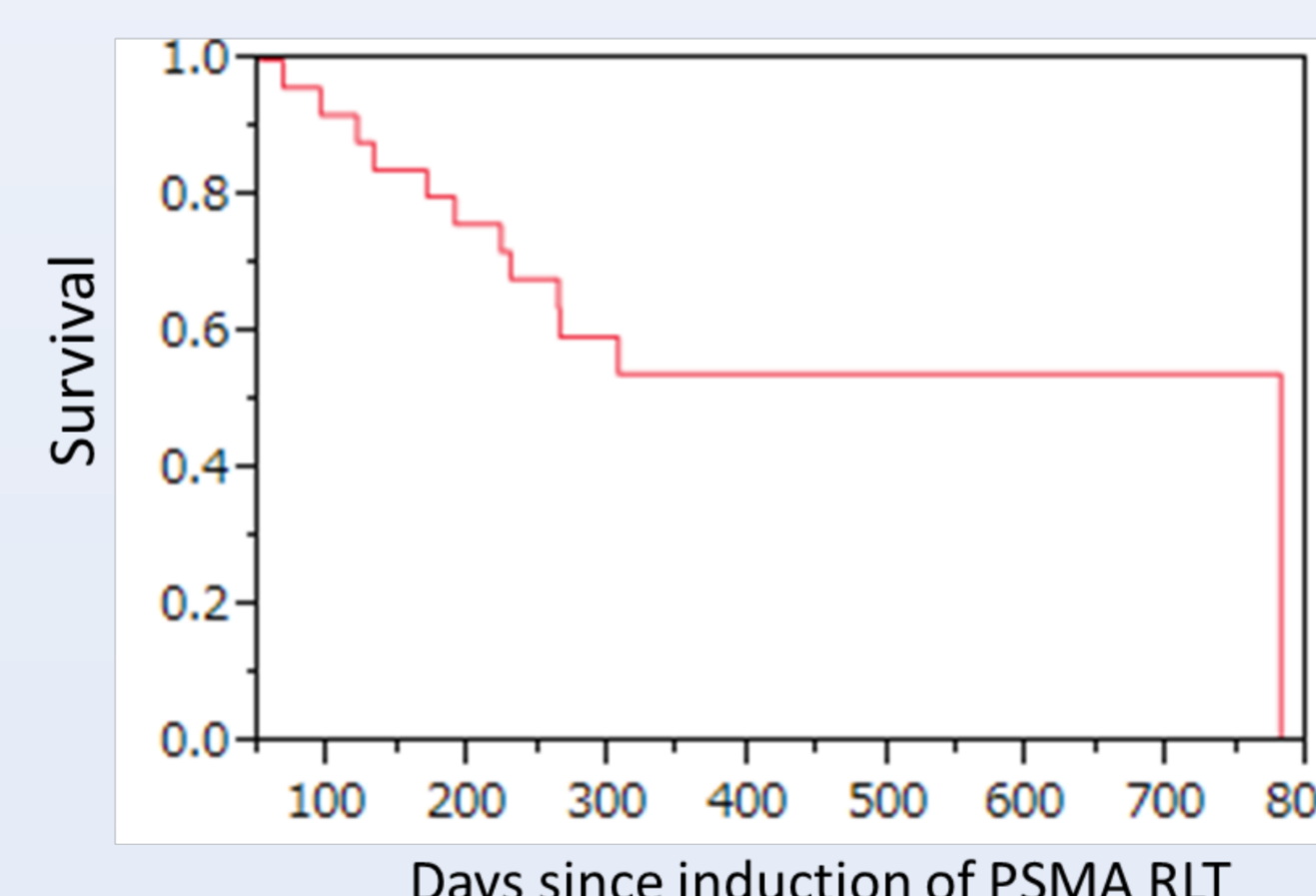
PSA < 300 ng/ml group (n=15)	
Progression	4 (26.7%)
Stable	1 (6.7%)
Reduction	10 (66.7%)
50% PSA reduction	6 (40.0%)



PSA ≥ 300 ng/ml group (n=11)	
Progression	8 (72.7%)
Stable	0 (0%)
Reduction	3 (27.2%)
50% PSA reduction	1 (9.0%)

## Survival

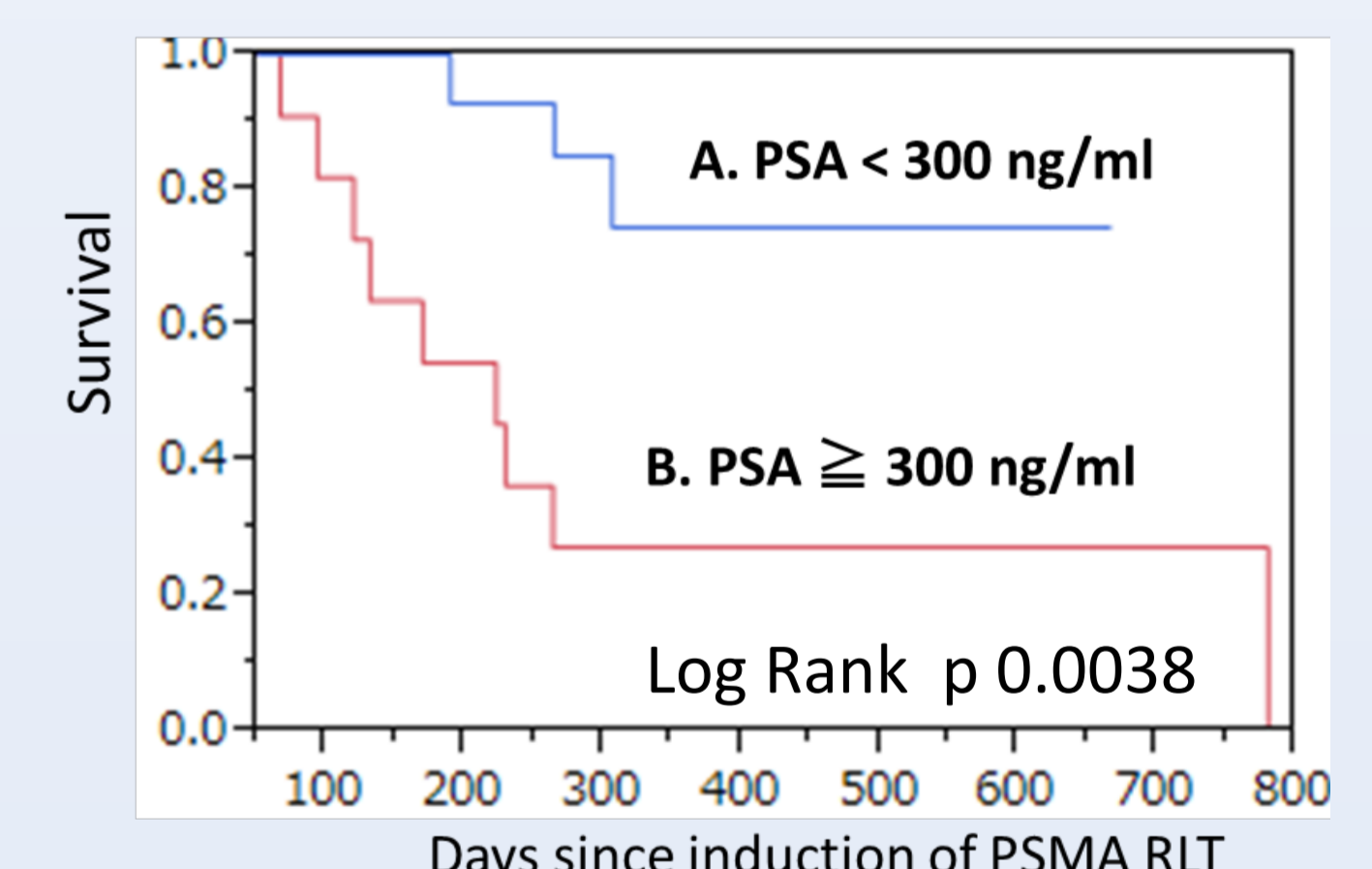
### Overall (n = 26)



Average 509.6, SE 64.0

Median 780, 95% CI 229-780

### PSA group



A. Median 222

B. Median -

## Toxicity (latest 15 patients)

Toxicity	Grade 1	Grade 2	Grade 3	Grade 4
<b>48 hrs</b>				
Dry mouth	2 (13.3%)	0 (0%)	0 (0%)	0 (0%)
Nausea	1 (6.7%)	0 (0%)	0 (0%)	0 (0%)
Dry eyes	0 (0%)	0 (0%)	0 (0%)	0 (0%)
<b>4 weeks</b>				
Dry mouth	2 (13.3%)	0 (0%)	0 (0%)	0 (0%)
Nausea	1 (6.7%)	0 (0%)	0 (0%)	0 (0%)
Dry eyes	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Leukopenia	3 (2.0%)	0 (0%)	0 (0%)	0 (0%)
Thrombocytopenia	1 (6.7%)	0 (0%)	0 (0%)	0 (0%)
Anemia	4 (26.7%)	0 (0%)	0 (0%)	0 (0%)
Increased Cr	3 (2.0%)	0 (0%)	0 (0%)	0 (0%)

CTCAE v5.0

**Patients and methods:** Japanese patients with CRPC underwent treatment with the consent of the patient after prior review of indications with the Australian medical team. The PSMA RLT was performed every 8-10 weeks, and imaging with PSMA PET/CT was performed before each treatment. Blood tests were performed at 4 and 7 weeks post-treatment, and a video interview with the Australian physician was conducted at 5 weeks to assess efficacy and adverse events. Two to four cycles of treatment were administered until COVID-19 travel restrictions were initiated depending on efficacy.

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COI 開示

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演題発表内容に関連し、発表者らに開示すべきCOI関係にある企業などはありません。