Abstract. Background/Aim: Cabazitaxel is recommended as first-line treatment after docetaxel for metastatic castration-resistant prostate cancer. However, the efficacy, adverse events and prognostic factors associated with cabazitaxel are unclear. Patients and Methods: This single-centre retrospective study including 30 patients with CRPC treated with cabazitaxel between 2014 and 2020 investigated efficacy, outcomes and prognostic factors. Results: Fourteen patients had visceral metastases. The median cabazitaxel dose was 20 mg/m². The prostate-specific antigen response rate, time to prostate-specific antigen response, and overall survival were 13.3%, 3.48 months, and 7.92 months, respectively. The rates of grade 3 or more neutropenia and febrile neutropenia were 20% and 6.7%, respectively. By multivariate analysis, sarcopenia and visceral metastasis at the time of cabazitaxel initiation were independent and significant factors conferring a poor prognosis. Conclusion: The early introduction of cabazitaxel, prior to the development of sarcopenia and visceral metastasis, might contribute to improved prognosis in CRPC.

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Key Words: Sarcopenia, visceral metastasis, castration-resistant prostate cancer, cabazitaxel.

Sarcopenia and Visceral Metastasis at Cabazitaxel Initiation Predict Prognosis in Patients With Castration-resistant Prostate Cancer Receiving Cabazitaxel Chemotherapy

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We retrospectively reviewed the charts of all patients and collected medical data, including age, serum PSA level, bone scan index (BSI), levels of C-reactive protein, alkaline phosphatase and lactate dehydrogenase, neutrophil-to-lymphocyte ratio, prostate biopsy pathology, adverse events, sarcopenia, clinical stage and treatment progress. BSI was calculated as the percentage of bone metastatic hotspots based on artificial neural network values using BONENAVI version 2 (FujiFilm RI Pharma, Tokyo, Japan; Exini Bone, Exini Diagnostics, Lund, Sweden) (17, 18). Sarcopenia was defined as a skeletal muscle index (SMI) <43 cm²/m² in patients with a body mass index (BMI) <25 cm²/m² or an SMI <53 cm²/m² in patients with a BMI ≥25 cm²/m² (19). SMI and BMI were calculated as follows: BMI (kg/m²)=weight/height²; SMI (cm²/m²)=skeletal muscle cross-sectional area at L3/height² (20, 21).

Clinical cancer stage was determined based on the eighth edition of the TNM classification published in 2017 (22). Diagnostic imaging studies such as computed tomography, magnetic resonance imaging, and bone scintigraphy were performed at the time of PC diagnosis or CRPC occurrence and before changes in treatment. Thereafter, the imaging studies were left to the discretion of the attending physician. The interval between subsequent imaging studies and all therapeutic decisions were left to the discretion of the attending physician.

The follow-up was terminated on December 31, 2020. Survival was measured from the time of cabazitaxel administration until death or last follow-up. Time to PSA progression (TTP) and OS were retrospectively analysed using the Kaplan-Meier method, and the log-rank test was used for the comparison of survival distributions. The Cox proportional hazards model was used for multivariate analyses. All statistical analyses were performed using the commercially available SPSS software, version 25.0 (SPSS, Chicago, IL, USA) and Prism 5 (GraphPad, San Diego, CA, USA). In all analyses, a p-value of less than 0.05 was considered to indicate statistical significance. The present study was approved by the Institutional Review Board of Kanazawa University Hospital (2016-328).

### Results

The characteristics of the study cohort of 30 patients with PC treated with cabazitaxel are shown in Table I. The median age at the time of cabazitaxel administration was 69.5 (range=48-80) years. The median PSA at the start of cabazitaxel treatment was 63.75 (range=0.24-22141) ng/ml. In the study cohort, 18 patients had a Gleason score ≥9, two patients had neuroendocrine PC, and 14 patients were in stage M1c indicating visceral metastasis (VM). A median of six (range=3-8) pretreatment lines and a median of four (range=1-10) cabazitaxel cycles were administered, with a
median cabazitaxel dose of 20 (range=15-20) mg/m². Grade 3 or higher neutropenia and febrile neutropenia (FN) were observed in six and two patients, respectively.

The waterfall plot of the maximum percentage change in PSA level from baseline after cabazitaxel administration is shown in Figure 1. Specifically, 16 out of the 30 patients (53.3%) exhibited a decrease in PSA and four patients (13.3%) exhibited PSA response. The waterfall plot of the maximum percentage change in BSI from baseline after cabazitaxel administration is shown in Figure 2. Briefly, 13 out of the 18 patients (72.2%) for whom BSI data were available exhibited a decrease in BSI. Finally, the waterfall plot showing the relationship between the maximum percentage change in PSA and the maximum percentage change in BSI (Figure 3) revealed that there was no correlation between the decrease in BSI and PSA.

Table II shows the results of univariate and multivariate analyses of factors associated with OS from the time of cabazitaxel administration. Multivariate analysis showed that the presence of sarcopenia and VM at the time of cabazitaxel initiation were independent and significant factors indicating a poor prognosis. The median TTP and OS from the initiation of cabazitaxel administration were 3.48 months and 7.92 months, respectively (Figure 4A and B). As shown in Figure 4C, the median OS rates of patients with and without sarcopenia were 5.45 and 16.82 months, respectively ($p<0.0001$, log-rank test). As shown in Figure 4D, the median OS rates of patients with and without VM were 5.45 and 12.02 months, respectively ($p=0.017$, log-rank test). These results indicate that the OS was significantly shorter in patients with sarcopenia and in those with VM.

Discussion

Based on the results of the phase III TROPIC trial, 25 mg/m² cabazitaxel in combination with prednisone was approved in 2010 for the treatment of patients with CRPC after docetaxel treatment (9). In the TROPIC trial, the median OS rates were 15.1 and 12.7 months for the cabazitaxel-treated and mitoxantrone-treated groups, respectively, showing a significantly longer OS for the cabazitaxel-treated group ($p<0.0001$) (9). The PSA response rate for the cabazitaxel group was 39.2%. Based on these results, the 2020 National Comprehensive Cancer Network Clinical Practice Guidelines in Oncology for Prostate Cancer recommend cabazitaxel as first-line treatment for metastatic CRPC after docetaxel (23). In the TROPIC trial, 82% of patients had grade 3 or higher neutropenia and 8% of patients had FN (9). In a phase I trial from Japan, 100% of patients had grade 3 or higher neutropenia and 24.6% of patients had FN (24). Compared with patients from Western countries, disease in patients from Asian countries has generally been reported to exhibit stronger resistance to chemotherapeutic agents, and the cabazitaxel dosage for Japanese patients should be considered with caution (9, 24, 25). The phase III PROSELICA trial was conducted to investigate the safety and efficacy of low-dose cabazitaxel [20 mg/m² (C20)] to 25 mg/m² cabazitaxel (C25) (26). The PSA response rate was significantly higher for the C25 group compared to the C20 group (42.9% vs. 29.5%;
but there was no significant difference in TTP (3.5 and 2.9 months, respectively; hazard ratio=1.099) and OS (14.5 and 13.4 months, respectively, hazard ratio=1.024) (26).

The C20 group had a lower rate of grade 3 or higher neutropenia (42% vs. 73% in the C25 group) and FN (2.1% vs. 9.2% in the C25 group). Based on these results, treatment

Table II. Univariate and multivariate analyses of factors associated with overall survival from the initiation of cabazitaxel administration.

| Factor                  | Univariate HR (95% CI) | p-Value | Multivariate HR (95% CI) | p-Value |
|-------------------------|------------------------|---------|--------------------------|---------|
| Age* ≥70 vs. <70 Years  | 0.83 (0.35-2.01)       | 0.68    | 3.80 (1.23-11.75)        | 0.02    |
| Visceral metastasis*    | 2.84 (1.16-6.95)       | 0.02    | 4.40 (0.98-19.82)        | 0.05    |
| PSA* ≥100 vs. <100 ng/ml| 3.68 (1.44-9.43)       | <0.01   | 2.87 (0.78-10.61)        | 0.11    |
| BSI* ≥2 vs. <2         | 1.55 (0.49-4.89)       | 0.46    | 0.66 (0.16-2.75)         | 0.56    |
| CRP* ≥2 vs. <2 mg/l    | 4.00 (1.43-11.12)      | <0.01   | 1.88 (0.42-8.33)         | 0.41    |
| ALD* ≥270 vs. <270 IU/l| 3.74 (1.37-10.25)      | 0.01    | 0.69 (0.24-1.49)         | 0.27    |
| NLR* ≥3 vs. ≤3         | 1.90 (0.69-5.25)       | 0.21    | 2.11 (0.78-5.67)         | 0.14    |
| Total dose of docetaxel| ≥1,000 vs. <1,000 mg    | 0.60    | 1.08 (0.45-2.59)         | 0.86    |
| Neutropenia grade 3/4  | Yes vs. no             | 2.11    | 1.08 (0.45-2.59)         | 0.86    |
| Time to CRPC ≥1 vs. <1 Year | Yes vs. no             | 1.90    | 2.11 (0.78-5.67)         | 0.14    |
| Sarcopenia Yes vs. no  | 9.50 (2.92-30.97)      | <0.001  | 12.19 (2.91-51.06)       | 0.001   |

ALP: Alkaline phosphatase; BSI: bone scan index; CI: confidence interval; CRP: C-reactive protein; LDH: lactate dehydrogenase; NLR: neutrophil-to-lymphocyte ratio; PSA: prostate-specific antigen. *At initiation of cabazitaxel administration.

Figure 4. Kaplan-Meier curves for time to prostate-specific antigen (PSA) progression (A), and overall survival (OS) of the whole cohort (B) and of patients according to the presence of sarcopenia (C) and visceral metastasis (D). HR: Hazard ratio.
with 20 mg/m² cabazitaxel is desirable due to improved prognosis with a lower rate of side-effects. For all patients in the present study, lower cabazitaxel doses of 15-20 mg/m² were initiated.

In the present study, the rates of PSA response, grade 3 or higher neutropenia, and FN were 13.3%, 20%, and 6.7%, respectively, and the OS and TTP were 7.92 and 3.48 months, respectively. The reported rates of PSA response, grade 3 or higher neutropenia, and FN are 19.37%-18-100%, and 6.4-54.6%, respectively, with TTP and OS of 1.4-4.3 and 12.2-20.7 months, respectively (10-13, 15, 16, 24). Therefore, compared with the previous reports, the patients in the present study experienced a lower PSA response rate, shorter OS and lower incidence of grade 3 or higher neutropenia and FN. The lower incidence of grade 3 or higher neutropenia and FN might be due to use of the reduced cabazitaxel dose. In this study, sarcopenia and VM at the time of cabazitaxel initiation were independent and significant factors conferring a poor prognosis. The Eastern Cooperative Oncology Group performance status and VM have been reported to be factors of poor prognosis for OS after treatment with cabazitaxel, consistent with the current study results (27). It is known that cancer cachexia and sarcopenia develop as cancer progresses, and this was observed in about half of patients with advanced cancer (28). VM at the time of prostate cancer diagnosis is very rare but has been found to increase as treatment progresses (5). These results suggest that cabazitaxel should be introduced at an early stage of PC, without sarcopenia and VM. Several studies have reported that cabazitaxel is effective in patients with grade 3 or higher neutropenia, low neutrophil-to-lymphocyte ratio, and low BSI, but in this study, there were no significant differences in OS and TTP related to these factors (27, 29-31). There are several limitations to the current study. This was a retrospective study with a short observation period and included a small number of patients. Furthermore, PC treatment and the interval between imaging assessments were at the discretion of the attending physician. Therefore, the current study findings should be confirmed in large, long-term prospective studies.

Conclusion

In the present study, we found that sarcopenia and VM at the time of cabazitaxel initiation were independent and significant factors indicating a poor prognosis. Early introduction of cabazitaxel might contribute to improved prognosis in patients with CRPC.

Conflicts of Interest

All Authors declare that there are no potential conflicts of interest relevant to this article.

Authors’ Contributions

H.I. designed the experiments. H.I., H.K., T.S., R.N., T.M., S.K., H.Y., S.K., K.I. and Y.K. collected clinical data. H.I., R.N., T.M., S.K., K.I. and A.M. analyzed the data. H.I., K.I., and A.M. drafted and revised the manuscript. All Authors read and approved the final version of the manuscript.

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