

Efficacy of anti-PD-1 antibody nivolumab in Japanese patients with metastatic renal cell carcinoma: A retrospective multicenter analysis

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Abstract. To evaluate the efficacy and safety of anti-PD1 therapy (nivolumab) in advanced renal cell carcinoma (RCC) in a clinical setting. Between March 2013 and January 2018, 33 patients with RCC (27 men and 6 women) were treated with nivolumab. Before anti-PD1 treatment, 12, 9 and 12 patients received one, two, and three or more therapies, respectively. Objective response, survival rate, and clinical adverse events were evaluated by the revised RECIST criteria (version 1.1). The median patient age was 68 years (range: 37-79). In total, 14 (42%) patients had an Eastern Cooperative Oncology Group performance status (ECOG PS) of 0 while 17 (52%) and two (6%) had an ECOG PS of 1 and 2 or higher, respectively. One (3%), 24 (73%) and eight (24%) were classified as having favorable, intermediate, and poor risk, respectively. The median follow-up duration after nivolumab initiation was 26 months (range: 1-131). The median progression-free and overall survival were 10.3 months and 45.9 months, respectively. Nivolumab was associated with a disease control rate of 58%, with an objective response of 24% (complete response, 1; partial response, 7; stable disease, 11; progressive disease, 10; not assessed, 4). A total of 15 (46%) patients experienced adverse events, of which six were severe (grade 3 or more) and 10 were immunotherapy-related. This study examined the initial experience of nivolumab administration in Japanese patients with advanced RCC. Our results suggest that nivolumab can achieve acceptable outcomes in a real clinical setting, with outcomes that are comparable to those of clinical trials.

Introduction

An estimated 20-30% of all patients with renal cell carcinoma (RCC) have metastatic disease at the time of diagnosis. Although multiple guidelines, including those of The National Comprehensive Cancer Network and the European Association of Urology, recommend first-line vascular endothelial growth factor (VEGF)-targeted therapy for patients with clear cell metastatic RCC (mRCC) (1,2), the response to first-line VEGF-targeted therapy is generally not durable, and most patients require additional therapy (3). The role of immune checkpoint inhibitors is increasingly of interest in cancer immunotherapeutics, and nivolumab, a novel anti-PD1 antibody, has been increasingly used in patients with several types of cancer. Although nivolumab has been most frequently used for melanoma, it is currently under evaluation for a number of other cancers. In terms of overall survival (OS), nivolumab has been shown to be superior to everolimus in a phase III randomized trial, and thus offers promising potential for use in patients with mRCC after treatment failure. However, unfamiliar toxicities have also been recognized with nivolumab use, such as immune-related adverse events (irAEs) (4). There has been no initial report about mRCC treated by nivolumab in the Japanese population yet.

The present study aimed to analyze the clinical outcomes and toxicities related to nivolumab in Japanese patients with mRCC in a real-world setting.

Patients and methods

Eligibility criteria. The present study included patients who were diagnosed with advanced RCC and treated with nivolumab from two institutes between March 2013 and January 2018. All patients with histologically-proven mRCC, regardless of Eastern Cooperative Oncology Group (ECOG) performance status (PS), were eligible for inclusion. All patients received at least two cycles of nivolumab and were assessed for treatment efficacy and toxicity. This study was approved by both institutional review boards, The Research

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Key words: renal cell carcinoma, nivolumab, prognosis, adverse events, initial case

Table I. Patient characteristics.

Characteristic	Result
Total patient number	33
Age, median year (range)	68 (37-79)
Sex, male:Female	27:6
Nephrectomy, yes:No	32:1
Histology	
Clear cell	29
Xp11.2 translocation	2
Papillary	1
Chromophobe	1
Sarcomatoid	1
MSKCC risk classification, Favorable:	
Intermediate:Poor	
At the beginning of systemic therapy	4:22:7
At the starting of nivolumab	1:24:8
Metastatic site, 1:2:≥3	12:11:10
ECOG PS, 0:1:≥2	
At the beginning of systemic therapy	24:8:1
At the starting of nivolumab	14:17:2
Number of prior therapy, 1:2:≥3	12:9:12

MSKCC, Memorial Sloan-Kettering Cancer Center; ECOG, Eastern Cooperative Oncology Group; PS, performance status.

Ethics board of Akita University Hospital (approval no. 1993) and Japanese Red Cross Akita Hospital, and all procedures were performed in accordance with the 1964 Declaration of Helsinki and its later amendments. Written informed consent was obtained from each patient.

Treatment and follow-up examinations. Complete medical history, physical examination, ECOG PS, CBC with differential and platelet count, biochemical profile (including electrolytes, renal and hepatic function, coagulation, pancreatic amylase, and lipase), urinalyses, and chest radiography were recorded for all patients before treatment was started; these tests were repeated during therapy according to the attending physician's decision. Toxicity was graded using the National Cancer Institute Common Toxicity Criteria version 2.0. Tumors were measured by computed tomography scans within 4 weeks prior to starting nivolumab. After starting the drug, the assessment interval was scheduled for individual patients by attending physicians. We defined that patients experienced tumor enlargement or appearance of a new lesion after initiating nivolumab and intolerable adverse event (AE) as 'nivolumab failure'. Tumor response was evaluated using the Response Evaluation Criteria in Solid Tumors (RECIST) guidelines v.1.1.

Results

Patient characteristics. A total 33 patients who were diagnosed with advanced RCC and eligible for the present study

Table II. Results of nivolumab management.

Factor	Result
Dose, mg/kg	3
Median (range)	11 (3-65)
Duration, median month (range)	23 (1-131)
Current status	
Continue	9
Stop	19
Pending	4
Unknown	1
Best response	
CR	1
PR	7
SD	11
PD	10
Not assessed	4
Objective response rate, %	24.2
Time to best response, median month (range)	3.2 (0.5-5.7)
Duration of best response, median month (range)	2.3 (2.0-26.6)

CR, complete response; PR, partial response; SD, stable disease; PD, progressive disease.

were enrolled. Patients' baseline characteristics are summarized in Table I. All patients were Japanese, and the cohort included 27 (81.8%) men and six (18.2%) women. The median patient age was 68 (range: 37-79) years. Thirty-two (97.0%) patients underwent radical nephrectomy before starting systemic therapies. Twenty-nine (87.9%) patients had clear cell histology, two (6.1%) patients had Xp11.2 translocation, and one patient each (3.0%) had papillary, chromophobe, and sarcomatoid histology. Twelve (36.4%) patients were administered nivolumab as second-line systemic therapy, nine (27.3%) as third-line therapy, and 12 (36.4%) as fourth-line or later therapy. Fourteen (42.4%) patients had an ECOG PS of 0, 17 (51.5%) patients had an ECOG PS of 1, and two (6.1%) patients had an ECOG PS of 2 or higher. Using the Memorial Sloan-Kettering Cancer Center (MSKCC) risk classification system, one (3.0%) patient was classified as being at favorable risk, 24 (72.7%) patients were classified as being at intermediate risk, and eight (24.2%) patients were classified as being at poor risk. Twelve (36.4%) patients had one metastatic site, 11 (33%) patients had two, and 10 (30%) patients had three or more.

Antitumor effect. An objective response (OR) was found in eight (24.2%) patients (Table II). The median progression-free survival was 12.3 months (range: 0-30.2), and the OS was 45.9 months (range: 0-53.9) (Fig. 1). These survival rates might be equivalent to the median PFS 4.6 months and OS 25.0 months in CheckMate-025 trial (4). The median time to best response after starting nivolumab was 3.2 months (range: 0.5-5.7 months). The median duration after achieving best response was 2.3 months (range: 2.0-26.6 months) (Table II).

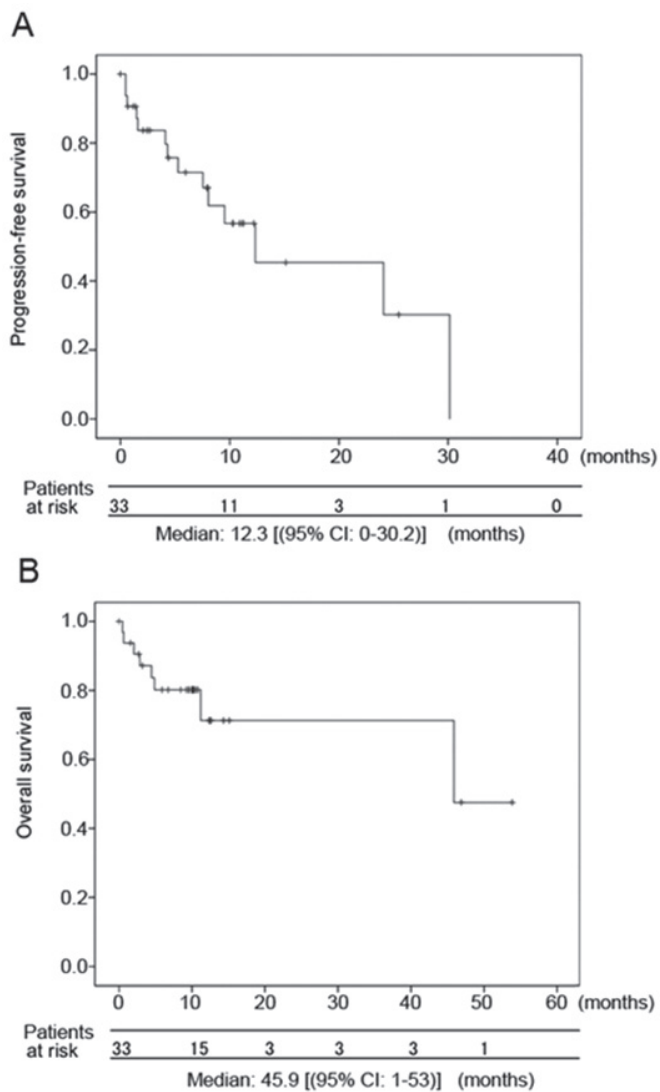


Figure 1. Kaplan-Meier curves for progression-free survival and OS. OS, overall survival.

Adverse events. At the data cutoff, 15 patients (45.5%) had experienced an AE (Table III). Six patients (18.2%) discontinued nivolumab treatment because of AEs. The most common AE was renal dysfunction (n=3). Grade 3 or higher AEs presented in six (18.2%) patients. IrAEs (any grade) included two cases each of interstitial pneumonia, myasthenia gravis, liver dysfunction, and hypothyroidism, and one case each of hypopituitarism and organizing pneumonia. Corticosteroid was administered as primary treatment for all irAEs, except for one patient with myasthenia gravis. Only one irAE, a case of hypopituitarism, was not resolved, and the patient has since required continuous steroid and thyroid hormone supplemental therapy. There was no relationship between irAE development and clinical response. Five patients died within 6 months after initiating nivolumab treatment. Of these five, three patients were already administered multiple therapies (at least four) and had advanced RCC before starting nivolumab treatment, and two patients had an ECOG PS of 2 or worse at the time of starting nivolumab.

Table III. Treatment-related adverse events.

Factor	Result
Patient number	33
Number of patients with AEs	15 (45.5%)
Cause of discontinuation	
PD	13
AE	6
Detail of discontinuation	
G3 or more	6
Myasthenia gravis	2
Liver dysfunction	1
General edema	1
Hypopituitarism	1
Heart failure	1
General AE, any grade	
Renal dysfunction	3
Pleural effusion	2
Heart failure	2
General edema	1
General fatigue	1
Hyper Kalemia	1
Rash	1
irAE, any grade	
Myasthenia gravis	2
Interstitial pneumonia	2
Liver dysfunction	2
Hypothyroidism	2
Hypopituitarism	1
Organizing pneumonia	1

PD, progressive disease; AE, adverse event; G, grade; ir, immune-related.

Table IV. Treatment after nivolumab failure.

Status	Patient number	Best response
Dead	3	
Axitinib	5	PR 2, SD 2, PD 1
Sunitinib	4	PR 1, SD 1, PD 2
Pazopanib	3	SD 2, PD1
Sorafenib	1	SD 1
Treatment pending	4	SD 2, PD 1, unknown 1
Move to other hospital	1	Unknown 1

PR, partial response; SD, stable disease; PD, progressive disease.

Treatments after nivolumab failure. Thirteen patients were administered further treatment after nivolumab failure during the period of this investigation. Of these, all patients received VEGF receptor inhibitor, with five receiving axitinib,

four receiving sunitinib, three receiving pazopanib, and one receiving sorafenib (Table IV). Only three patients (OR rate: 20.8%) showed partial response (PR), but six patients achieved stable disease (SD) even after nivolumab failure, indicating some clinical benefit.

Discussion

This study found that nivolumab was relatively effective and showed acceptable results in Japanese patients with mRCC. The OR rate and survival rate in our cohort were comparable to those of previous clinical trials (4), despite the fact that our cohort included patients with worse PS, MSKCC-determined poor risk, and non-clear cell histology. Nine patients had SD at the time of this study, and four patients showed durable responses for over 12 months; these findings indicate the potential utility of immune checkpoint blockade for the treatment of patients with mRCC (5). This relatively better outcome than we expected before nivolumab launched despite unfavorable patient clinical backgrounds could support the assertion that immune checkpoint inhibitors are a revolutionary therapy (6).

Research efforts to determine a predictive marker, such as PD-L1 positivity (4), mutation burden (7), and profile of immune-related genes (8), are currently ongoing. Most previous studies have focused on identifying exceptional responders to immune checkpoint inhibition; however, even patients with lower expression of PD-L1 showed an OR rate of over 20% in the CheckMate-025 trial (4). Therefore, immunohistological evaluation of PD-L1 expression, would not change our clinical decision, because a substantial number of patients without PD-L1 expression can still benefit from nivolumab therapy.

The timing of nivolumab administration may be another important consideration for appropriate clinical decisions. Although our sample size was small, and we could not assess statistical significance, we saw a trend between worse survival and both poor ECOG PS and multiple prior therapies in our cohort. This suggests that the maximum benefit from nivolumab is likely to occur early in the sequential course of multiple treatments and prior to cancer advancement or decline in a patient's general status. Early exposure to immune checkpoint inhibitors can be reinforced by the possibility of late immune response with anti-tumor effect (9). Although in this study there were no patients who met the criteria for 'pseudo-progression' (10) with subsequent response, one patient with SD did eventually develop PR. Determining the appropriate time to examine the 'delayed response' (11) of an immune-related anti-tumor effect can be difficult, which emphasizes the importance of earlier introduction and good PS in nivolumab treatment (10).

Although immune checkpoint-blocking antibodies are effective in many types of cancers, irAEs may be unavoidable (12). In addition to relatively common AEs, such as interstitial pneumonia, colitis, hepatitis, kidney insufficiency, and hypo-endocrinology, uncommon AEs have also been reported, such as cardiotoxicity (13), type 1 diabetes (14), and myasthenia gravis (15). In this study, several irAEs were observed in response to nivolumab despite the small size of the patient cohort. In the important phase III trial of nivolumab (4), grade 3-4 toxicities were reported at a rate of 21%. The rate of grade 3-4 toxicities observed in this study (18.2%) was similar,

despite the short observation period. Importantly, all grade 3-4 irAEs that occurred in patients in this study, with the exception of hypopituitarism, resolved with prompt initiation of corticosteroid administration followed by a slow steroid taper. This is similar to what has been described in the management of irAEs in other cancers, and underscores the necessity for vigilance by the attending physician concerning these potential events (16). Our results indicate that nivolumab might be a relatively safe drug for the treatment of mRCC in real clinical practice.

The proper endpoint for nivolumab treatment has been controversial (17,18). In several patients in our study, no tumor progression was observed after cessation of nivolumab treatment. Based on these cases, we suggest that nivolumab treatment not be stopped prematurely, even after an initial therapeutic effect is achieved. However, it is not clear from our study whether prolongation of treatment would further improve clinical outcome. It has been reported that long-term application of anti-PD1 antibody can also cause drug resistance (19). Hence, the duration of nivolumab application for mRCC treatment needs to be further studied.

Several limitations of this study should be noted, including the retrospective nature and limited number of patients. However, our results show promising relevance for nivolumab in clinical practice and warrant further research of immune checkpoint inhibitor therapy in advanced RCC.

The present study demonstrated the safety and acceptable outcomes of anti-PD1 therapy with nivolumab in patients with advanced RCC. Although nivolumab precipitated unique AEs, these were treated promptly and all but one were reversible. Our findings suggest that nivolumab is a reasonable therapeutic option for patients with mRCC after multiple lines of treatment. Future prospective studies with larger sample sizes are required to better understand the appropriate indications of nivolumab for patients with RCC.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

KN was involved in the project development, data collection and management, data analysis, and manuscript writing. YH collected the data and edited the manuscript. SK was involved in data collection. AK, TN, SK, MS, SN, TI, TO and SC collected the data. NS collected the data and edited the manuscript. TH was involved in the project development, data collection, and manuscript editing.

Ethics approval and consent to participate

The present study was conducted in accordance with the Declaration of Helsinki and approved by the Research Ethics board of Akita University Hospital (approval no. 1993) and Japanese Red Cross Akita Hospital. Written informed consent was obtained from each patient.

Patient consent for publication

Written informed consent was obtained from each patient for the publication of their data.

Competing interests

Dr Habuchi received honoraria from Novartis International AG, Pfizer, Inc., GlaxoSmithKline and Ono Pharmaceutical Co., Ltd. All other authors declare that they have no competing interests.

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