

# Dynamics of the neutrophil-to-lymphocyte ratio during lenvatinib treatment for unresectable hepatocellular carcinoma

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**Abstract.** Lenvatinib is an approved therapy for advanced hepatocellular carcinoma (HCC). Recently, immune checkpoint inhibitors have been approved as frontline chemotherapies for HCC, and the tumor immune microenvironment (TIME) has been demonstrated to significantly affect HCC treatment. The neutrophil-to-lymphocyte ratio (NLR) is associated with the TIME, and the dynamics of the NLR are associated with prognosis or treatment efficacy in various cancer types. The present study investigated the dynamics of the TIME using the NLR in 101 patients with HCC treated with lenvatinib. Immunostaining for CD8<sup>+</sup> tumor-infiltrating lymphocytes (TILs) was also performed in 9 patients who underwent liver tumor biopsy prior to subsequent chemotherapy for progression or discontinuation due to adverse events on lenvatinib treatment. The NLR values measured at the start of treatment (SOT), after 1 month of treatment and after 3 months of treatment were  $2.78 \pm 2.20$ ,  $2.61 \pm 1.86$  and  $2.66 \pm 2.36$ , respectively ( $P=0.733$ ). Among the patients with no reduction in the initial dose, there was no significant difference between the NLR after 1 month ( $2.34 \pm 0.25$ ) and that at the SOT ( $2.86 \pm 2.33$ ) ( $P=0.613$ ). In patients who achieved a complete or partial response, the NLR at the time of the best tumor response was  $1.65 \pm 0.56$ , which was significantly lower than that at the SOT ( $2.05 \pm 0.78$ ) ( $P=0.023$ ). In patients who did not respond to lenvatinib, the NLR at the time of disease progression was

$3.68 \pm 3.19$ , which was significantly higher than that at the SOT ( $2.78 \pm 1.79$ ) ( $P=0.043$ ). Overall, 5 out of the 6 patients who did not respond to lenvatinib had low CD8<sup>+</sup> TIL counts at disease progression. Although the present study included a limited number of patients, the NLR was associated with the therapeutic effects of lenvatinib. These findings suggest the potential of lenvatinib as an immunomodulator.

## Introduction

With an estimated 900,000 new cases and 830,000 associated deaths in 2020, hepatocellular carcinoma (HCC) ranks as the sixth most common neoplasm and the third leading cause of cancer-related mortality worldwide (1,2). Recent advancements in systemic chemotherapy for advanced HCC, including immune checkpoint inhibitors (ICIs) and molecular targeted agents, have enhanced patient outcomes (3-8). The main elements of the tumor immune microenvironment (TIME) include cancer cells, antigen, immune cells and cytokines. These components interact with each other to determine the tendency of antitumor immunity (9). ICIs exhibit antitumor effects by reactivating the immune cells in TIME and it is imperative to elucidate the TIME in HCC.

Lenvatinib, an oral multi-kinase inhibitor targeting vascular endothelial growth factor receptors 1-3, fibroblast growth factor receptors 1-4, platelet-derived growth factor receptor  $\alpha$ , rearranged during transfection and stem cell factor receptor, has demonstrated anticancer efficacy (10). A global, randomized, multicenter, open-label trial assessing the non-inferiority of lenvatinib compared with sorafenib (REFLECT; NCT01761266) revealed that lenvatinib significantly improved progression-free survival (PFS) versus sorafenib in patients with previously untreated, metastatic or advanced HCC (3). Lenvatinib is currently approved for the treatment of HCC. Recently, Yamauchi *et al* (11) described the capability of lenvatinib to modulate the TIME in HCC.

Inflammation has an important role in cancer, and neutrophils suppress T cell function by secreting myeloperoxidase and arginase-1, and upregulating programmed cell death ligand 1 (12). Therefore, neutrophils create an immunosuppressive tumor microenvironment that reduces the efficacy of immunotherapy (13). Lymphocytes also have a role in cytotoxic cell death, and they produce cytokines to inhibit

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*Abbreviations:* HCC, hepatocellular carcinoma; ICI, immune checkpoint inhibitor; TIME, tumor immune microenvironment; PFS, progression-free survival; NLR, neutrophil-to-lymphocyte ratio; CR, complete response; PR, partial response; DCR, disease control rate; ORR, objective response rate; IHC, immunohistochemistry; TIL, tumor-infiltrating lymphocyte; CI, confidence interval; SOT, start of treatment

*Key words:* HCC, lenvatinib, NLR, immune microenvironment

tumor cell growth (14). The neutrophil-to-lymphocyte ratio (NLR) is considered a systemic marker of the balance between adaptive immune surveillance and the inflammatory status. A high NLR at baseline is associated with a poor prognosis in numerous types of cancer, such as lung, thyroid, biliary tract and colon cancer, and the dynamics of the NLR are associated with prognosis or treatment efficacy in various cancer types, such as lung cancer, renal cell carcinoma and gastrointestinal cancer, treated with systemic chemotherapies such as ICIs (15-22). It has been reported that the dynamics of the NLR reflect changes in the TIME and capture antitumor immune responses, ultimately being associated with clinical outcomes following immune checkpoint blockade (23). To the best of our knowledge, to date, no reports have evaluated the dynamics of the NLR as a biomarker of the TIME during lenvatinib therapy in HCC. The present study therefore investigated the dynamics of the NLR in this context.

### Patients and methods

**Patients.** The current prospective, single-center study analyzed the dynamics of the NLR in patients with HCC who were treated with lenvatinib at Aso Iizuka Hospital (Iizuka, Japan) between May 2018 and February 2023. In total, 130 patients with unresectable HCC who received lenvatinib treatment as first-line treatment or post-progression treatment after other therapies, including transarterial chemoembolization, sorafenib, and atezolizumab plus bevacizumab, were identified. Finally, 101 patients were evaluated, after excluding 29 patients who were observed for <12 weeks and did not have images to assess treatment efficacy. Additionally, liver tumor biopsy samples were obtained with consent from 9 patients treatment to assess the TIME prior to subsequent chemotherapy treatment for progression or discontinuation due to adverse events on lenvatinib treatment. This study adhered to the Declaration of Helsinki guidelines and received approval from the Iizuka Hospital Ethics Committee (approval no. 18070). All patients provided written informed consent. Specific written informed consent was obtained from 2 patients for the publication of their immunohistochemistry results.

**Biomarker analysis.** Peripheral blood (2 ml) was obtained from the patients at the start of treatment and at each hospital visit during lenvatinib treatment. The NLR was a calculation based on the absolute neutrophil count divided by the absolute lymphocyte count determined by complete blood count differential in the peripheral blood.

**Treatment protocol.** Patients received oral lenvatinib (Eisai Co., Ltd.) based on body weight (8 mg/day for those weighing <60 kg and 12 mg/day for those weighing  $\geq$ 60 kg). Reduction of the initial dose was permitted according to the performance status (by assessment of the level of function and capability of self-care) and the presence of proteinuria at the start of treatment (SOT) (4-8 mg/day). Dose adjustment, including interruption and reduction (to 8 mg/day, 4 mg/day or 4 mg every other day), was permitted during treatment according to the performance status and adverse events. The protocols outlined in the REFLECT trial, as prescribed by Eisai Co., Ltd., were followed (3). Adverse events were graded using the

Common Terminology Criteria for Adverse Events, version 4.0 (24). Grade 3 or higher adverse events or any unacceptable grade 2 events led to a reduction in the drug dose or interrupted treatment according to the lenvatinib administration guidelines. Following the occurrence of an adverse event, the lenvatinib dose was reduced or treatment was temporarily halted until symptoms improved to grade 1 or 2, in line with Eisai Co., Ltd., guidelines.

**Evaluation of efficacy.** The treatment response was assessed every 8-12 weeks after treatment initiation using computed tomography or magnetic resonance imaging. The antitumor response was evaluated by the treating physician based on the modified Response Evaluation Criteria in Solid Tumours version 1.1 (25). The disease control rate (DCR) was defined as the sum of the rates for complete response (CR), partial response (PR) and stable disease lasting at least 4 months. The objective response rate (ORR; also referred to as the best response) was defined as the sum of the PR and CR rates. Patients were followed up every 4 weeks, and long-term treatment was continued until disease progression or intolerable side effects occurred.

**Immunohistochemistry (IHC).** Liver tumor biopsy specimens were fixed in 10% formalin at room temperature for 10-48 h and embedded in paraffin. Serial sections (5  $\mu$ m) were cut from the paraffin blocks and stained with hematoxylin and eosin (hematoxylin for 3 min and eosin for 45 sec at room temperature). CD8<sup>+</sup> T-cell staining was performed with a Leica Bond-III, which is an automatic and continuous access slide-staining system that simultaneously processes IHC protocols, using a Bond Polymer Refine Detection Kit (Leica Biosystems). Specimens were then incubated for 30 min with the primary antibody mouse anti-human monoclonal CD8 antibody (clone C8/144B; 1:50; Dako; Agilent Technologies, Inc.), followed by visualization with the Leica Bond Polymer Refine Detection kit for 20 min at room temperature. The sections were counterstained with hematoxylin, dehydrated and mounted. The slides were examined under the BZ-X700 fluorescence microscope (Keyence Corporation). CD8<sup>+</sup> cell infiltration was quantified according to the number of positively stained CD8<sup>+</sup> tumor-infiltrating lymphocytes (TILs) at x400 magnification, focusing on areas with the densest CD8<sup>+</sup> TIL presence. A cutoff of 15.9 cells/high-power field was utilized to classify high and low CD8<sup>+</sup> TIL infiltration, consistent with a previous report (26).

**Statistical analysis.** JMP Pro version 11 (SAS Institute Inc.) was utilized for all statistical analyses. Data are presented as the median (interquartile range) or mean (standard deviation). The Kaplan-Meier method was applied for statistical testing to evaluate overall survival (OS) time, PFS time and first objective response time. NLR was compared at different time points using Friedman's test with Dunn's post hoc test or a paired t-test.  $P < 0.05$  was used to indicate a statistically significant difference.

### Results

**Patient characteristics.** The characteristics of the 101 patients who received lenvatinib are presented in Table I. A total of

Table I. Baseline and overall characteristics of patients who received lenvatinib.

Characteristics	Value
Number of patients	101
Age, years <sup>a</sup>	73.0 (68.3-80.0)
Males/females, n	77/24
MVI-positive, n	20
EHS-positive, n	30
Intrahepatic max tumor size, cm <sup>a</sup>	3.1 (2.0-5.2)
Patients with >5 tumors, n	50
Etiology, n	
HBV	19
HCV	42
NBNC	40
Child-Pugh score, n	
A	83
B/C	18
Alb, g/dl <sup>a</sup>	3.7 (3.3-4.1)
T.Bil, g/dl <sup>a</sup>	0.8 (0.6-1.2)
ALBI score <sup>a</sup>	-2.39 (-2.75 to -2.01)
BCLC stage, n	
A	12
B	45
C	44
Tumor markers <sup>a</sup>	
AFP, ng/ml	23.7 (4.2-4392.1)
PIVKA-II, mAU/ml	189.0 (29.0-2086.0)
Initial dose reduction, n (%)	54 (53.5%)
ORR (CR + PR), n (%)	26 (25.7)
DCR (CR + PR + SD), n (%)	59 (58.4)
Median PFS, months <sup>b</sup>	6.0 (4.9-7.5)
Median OS, months <sup>b</sup>	27.9 (16.5-32.8)

<sup>a</sup>Data are expressed as median (interquartile range). <sup>b</sup>Data are expressed as median (95% confidence interval). HBV, hepatitis B virus; HCV, hepatitis C virus; NBNC, non-B non-C; MVI, microvascular invasion; EHS, extrahepatic spread; Alb, albumin; T.Bil, total bilirubin; ALBI score, albumin-bilirubin score; BCLC, Barcelona Clinic Liver Cancer; AFP,  $\alpha$ -fetoprotein; PIVKA-II, vitamin K absence or antagonist-II; ORR, objective response rate; CR, complete response; PR, partial response; DCR, disease control rate; SD, stable disease; PFS, progression free survival; OS, overall survival.

54 patients (53.5%) required a reduction of the initial dose of lenvatinib. The ORR was 25.7% (26/101 patients) and the DCR was 58.4% (59/101 patients). The median PFS time was 6.0 months [95% confidence interval (CI), 4.9-7.5] and the median OS time was 27.9 months (95% CI, 16.5-32.8). Median time to first objective response was 3.1 months (95% CI, 2.3-3.6 months).

**Dynamics of the NLR after treatment with lenvatinib.** The NLR values at the SOT, after 1 month of treatment and after 3 months of treatment were  $2.78 \pm 2.20$ ,  $2.61 \pm 1.86$  and

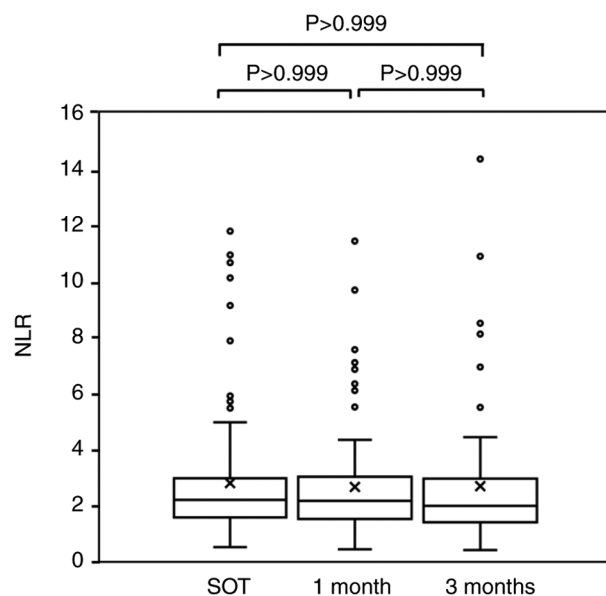


Figure 1. NLR dynamics after treatment with lenvatinib in all patients. NLR, neutrophil-to-lymphocyte ratio; SOT, start of treatment.

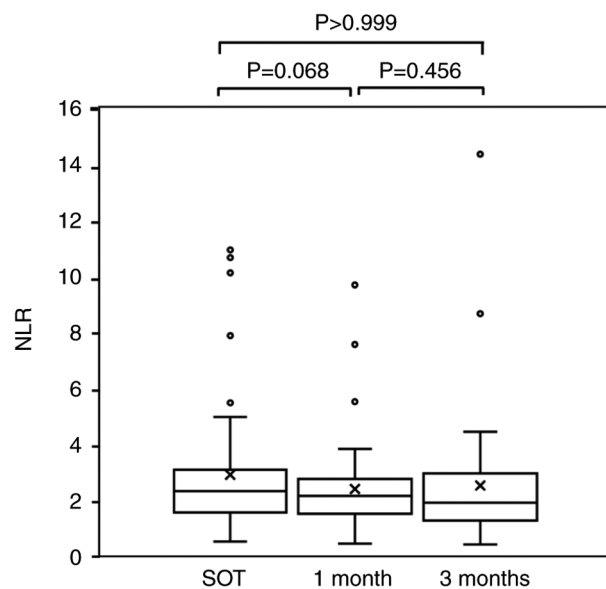


Figure 2. NLR dynamics after treatment with lenvatinib in patients with no reduction of the initial dose. NLR, neutrophil-to-lymphocyte ratio; SOT, start of treatment.

$2.66 \pm 2.36$ , respectively ( $P=0.733$ ; Friedman's test) (Fig. 1). Among the patients with no reduction of the initial dose, the NLR values at the SOT, after 1 month of treatment and after 3 months of treatment were  $2.86 \pm 2.33$ ,  $2.34 \pm 0.25$  and  $2.48 \pm 2.26$  ( $P=0.613$ ; Friedman's test) (Fig. 2). There was no significant difference between the NLR after 1 month and that at the SOT. Among the patients with an objective response, the NLR at the time of the best tumor response was  $1.65 \pm 0.56$ , which was significantly lower than that at the SOT ( $2.05 \pm 0.78$ ) ( $P=0.023$ ; Fig. 3). Among the non-responders, the NLR was significantly higher at the time of disease progression ( $3.68 \pm 3.19$ ) compared with that at the SOT ( $2.78 \pm 1.79$ ) ( $P=0.043$ ; Fig. 4).

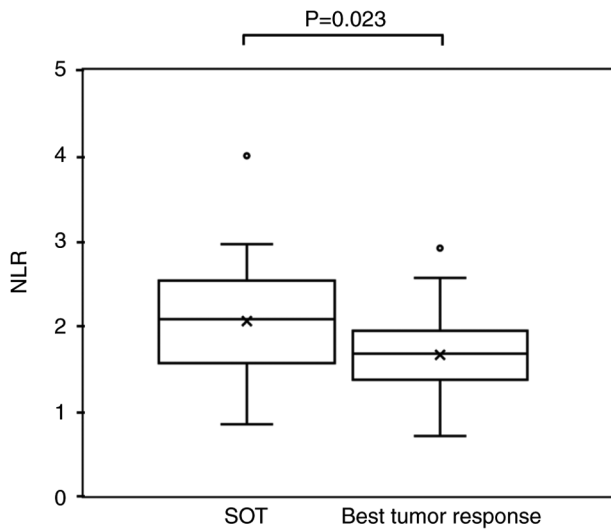


Figure 3. NLR dynamics after treatment with lenvatinib in patients who achieved a complete or partial response (best tumor response). NLR, neutrophil-to-lymphocyte ratio; SOT, start of treatment.

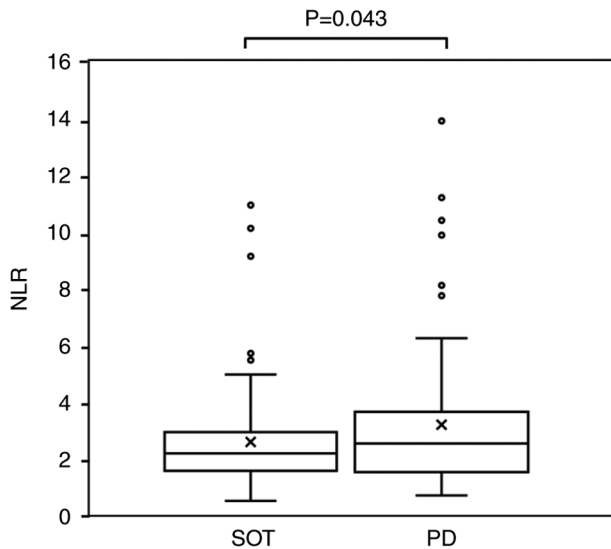


Figure 4. NLR dynamics after treatment with lenvatinib in patients who did not respond to treatment. NLR, neutrophil-to-lymphocyte ratio; SOT, start of treatment; PD, progressive disease.

**IHC for CD8<sup>+</sup> TILs in HCC tissues.** CD8<sup>+</sup> TIL counts were assessed by IHC after lenvatinib treatment and prior to subsequent-line chemotherapy in 9 patients (Table II). In total, 5 out of 6 patients who did not respond to lenvatinib had low CD8<sup>+</sup> TIL counts at disease progression. A typical case is presented in Fig. 5A (case 6). Furthermore, 2 out of the 3 patients who discontinued treatment due to adverse events had high CD8<sup>+</sup> TIL counts. A typical case is presented in Fig. 5B (case 7).

## Discussion

The immune response serves a crucial role in the progression of cancer. The most recent immunogenomic classification of HCC was published in 2022 (27). This study reported that ICI treatment was likely to initiate a response in 65% of HCC

cases in the non-inflammatory group and in 35% of cases in the inflammatory group. The inflammatory group was characterized by robust interferon signaling and cytolytic activity, upregulated effector molecules of cytotoxic T cells, and increased checkpoint molecule levels and CD8<sup>+</sup> T-cell counts. HCC is influenced by the TIME, which has been reported to benefit from immune checkpoint blockade treatment (28).

Clinical trials and preclinical studies investigating the immunomodulatory effects of antiangiogenic agents on the tumor microenvironment have highlighted enhanced maturation of dendritic cells, improved trafficking and function of T cells, and reversal of immunosuppression that is induced by hypoxia or immunosuppressive cells (29-31). Further *in vivo* and *in vitro* studies have illustrated that molecular targeted agents enhance antitumor immunity by promoting the polarization of tumor-associated macrophages to an M1 phenotype (32-34), enhancing the infiltration and functions of CD4<sup>+</sup> and CD8<sup>+</sup> T cells (35,36), reducing the numbers of regulatory T cells (37-39), and reversing the suppressive functions of myeloid-derived suppressor cells in the tumor microenvironment (40,41). Lenvatinib has also been demonstrated to modulate the TIME (11,42-44). It is important to evaluate the TIME in the treatment of HCC. However, previous studies have required liver tumor biopsy tissues to be obtained, highlighting the need for a non-invasive biomarker for predicting treatment response.

The NLR is a simple and inexpensive measure of the balance between adaptive immune surveillance and the inflammatory status (16). Tada *et al.* (45) reported that a high NLR was associated with negative outcomes (PFS, ORR, and DCR) in patients who received lenvatinib for HCC. However, the dynamics of the NLR in patients with unresectable HCC treated with lenvatinib have not been thoroughly investigated.

In the present study, the NLR decreased at the time of the best tumor response among the patients with an objective response, indicating an inflammatory condition, whereas NLR elevation at the time of disease progression suggested a non-inflammatory condition. There was notably less CD8<sup>+</sup> TIL infiltration in liver tumor tissue at the time of disease progression in patients who did not respond to lenvatinib. The results suggest that NLR may be useful for assessing the TIME and treatment efficacy.

Recently, the combination of an ICI and a vascular endothelial growth factor inhibitor (atezolizumab plus bevacizumab) and the combination of the ICIs tremelimumab and durvalumab were approved as systemic therapy options for patients with advanced HCC (4,8). If the tumor is inflamed prior to ICI treatment, the response to ICI treatment might be improved. Therefore, switching to ICI treatment early or before disease progression after lenvatinib administration could improve prognosis.

The limitations of the present study include the small number of patients due to the single-center design and the lack of observation of tumor tissue over time. The NLR can be influenced by numerous factors, including age, body mass index, steroidal drugs, viral hepatitis, alcoholic fatty liver and diabetes (46,47). The present study encompassed advanced HCC cases with varying stages and levels of liver function. Matching patients according to these factors is not feasible when analyzing a small case series. These factors warrant consideration, and randomized controlled trials should be

Table II. CD8<sup>+</sup> TIL levels after lenvatinib treatment and prior to subsequent-line chemotherapy.

Case	Age, years	Sex	Etiology	Reason for discontinuation of treatment	NLR at the start of treatment	NLR at PD or discontinuation of treatment	CD8 <sup>+</sup> TIL infiltration
1	52	M	HBV	PD	1.65	2.17	Low
2	59	M	ALC	PD	5.99	1.10	Low
3	76	M	HCV	PD	2.28	3.52	Low
4	87	M	NBNC	PD	1.87	5.09	Low
5	73	F	HBV	PD	0.71	0.85	Low
6	68	M	HBV	PD	2.39	4.27	High
7	79	M	HCV	Proteinuria	2.61	2.11	High
8	61	M	HCV	Proteinuria	7.86	2.71	High
9	69	M	NBNC	Proteinuria	2.62	2.07	Low

TIL, tumor-infiltrating lymphocyte; NLR, neutrophil-to-lymphocyte ratio; HBV, hepatitis B virus; ALC, alcoholic; HCV, hepatitis C virus; NBNC, non-B non-C; PD, disease progression; SOT, start of treatment.

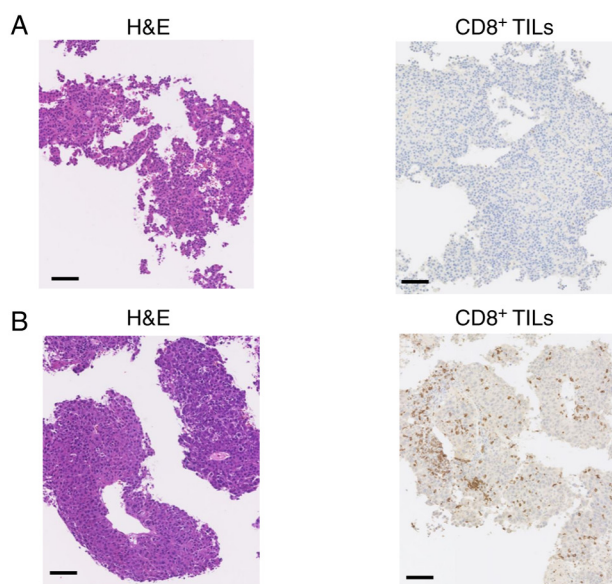


Figure 5. H&E and IHC in liver specimens of two typical cases. (A) H&E and IHC results in a patient who did not respond to lenvatinib. A low CD8<sup>+</sup> TIL count is shown. (B) H&E and IHC results in a patient who discontinued treatment due to adverse events. A high CD8<sup>+</sup> TIL count is shown. HE staining: Magnification, x100; scale bar, 100  $\mu$ m. IHC: Magnification, x100; scale bar, 100  $\mu$ m. IHC, immunohistochemistry; H&E, hematoxylin and eosin; TIL, tumor-infiltrating lymphocyte.

conducted in future. Nevertheless, despite the limitations of the present study, NLR dynamics may be recognized in future as a useful marker of the TIME.

In conclusion, the NLR at the time of the best tumor response was lower than that at the SOT among the patients with a PR or CR. Among the non-responders, the NLR was higher at the time of disease progression than at the SOT. These findings suggest the potential of lenvatinib as an immunomodulator. Further studies exploring the impact of different treatment methods on the TIME of HCC and further studies with larger sample sizes are required to investigate the TIME in patients with advanced HCC.

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### Availability of data and materials

The data generated in the present study may be requested from the corresponding author.

### Authors' contributions

AK, MY, AM and KM designed the study. AK, YK and KT assisted with data analyses. YO performed pathological examinations, including immunostaining. AK wrote the initial draft of the manuscript. MY contributed to data analysis and interpretation. MY, AM and KM assisted in the preparation and critical review of the manuscript. AK and MY confirm the authenticity of all the raw data. All authors read and approved the final version of the manuscript and agree to be accountable for all aspects of the work.

### Ethics approval and consent to participate

The study was performed in accordance with the principles and ethical guidelines of the 1975 Declaration of Helsinki. The study received approval from the Aso Iizuka Hospital Ethics Committee (Iizuka, Japan; approval no. 18070). All patients provided written informed consent.

### Patient consent for publication

Written informed consent was obtained from two patients for the publication of their immunohistochemistry results.

## Competing interests

The authors declare that they have no competing interests.

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