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ORIGINAL ARTICLE

Randomised, multicentre prospective trial of transarterial chemoembolisation (TACE) plus sorafenib as compared with TACE alone in patients with hepatocellular carcinoma: TACTICS trial

Masatoshi Kudo ,¹ Kazuomi Ueshima,¹ Masafumi Ikeda,² Takuji Torimura,³ Nobukazu Tanabe,⁴ Hiroshi Aikata,⁵ Namiki Izumi,⁶ Takahiro Yamasaki,⁷ Shunsuke Nojiri,⁸ Keisuke Hino,⁹ Hidetaka Tsumura,¹⁰ Teiji Kuzuya,¹¹ Norio Isoda,¹² Kohichiroh Yasui,¹³ Hajime Aino,¹⁴ Akio Ido,¹⁵ Naoto Kawabe,¹⁶ Kazuhiko Nakao,¹⁷ Yoshiyuki Wada,¹⁸ Osamu Yokosuka,¹⁹ Kenichi Yoshimura,²⁰ Takuji Okusaka,²¹ Junji Furuse,²² Norihiro Kokudo,²³ Kiwamu Okita,²⁴ Philip James Johnson ,²⁵ Yasuaki Arai,²⁶ on behalf of the TACTICS study group

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For numbered affiliations see end of article.

Correspondence to

Dr Masatoshi Kudo, Department of Gastroenterology and Hepatology, Kindai University Faculty of Medicine, Osaka-Sayama, Japan; m-kudo@med.kindai.ac.jp

MK and KU contributed equally.

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ABSTRACT

Objective This trial compared the efficacy and safety of transarterial chemoembolisation (TACE) plus sorafenib with TACE alone using a newly established TACE-specific endpoint and pre-treatment of sorafenib before initial TACE.

Design Patients with unresectable hepatocellular carcinoma (HCC) were randomised to TACE plus sorafenib (n=80) or TACE alone (n=76). Patients in the combination group received sorafenib 400 mg once daily for 2–3 weeks before TACE, followed by 800 mg once daily during on-demand conventional TACE sessions until time to untreatable (unTACEable) progression (TTUP), defined as untreatable tumour progression, transient deterioration to Child-Pugh C or appearance of vascular invasion/extrahepatic spread. Co-primary endpoints were progression-free survival (PFS), which is not a conventional one but defined as TTUP, or time to any cause of death plus overall survival (OS). Multiplicity was adjusted by gatekeeping hierarchical testing.

Results Median PFS was significantly longer in the TACE plus sorafenib than in the TACE alone group (25.2 vs 13.5 months; p=0.006). OS was not analysed because only 73.6% of OS events were reached. Median TTUP (26.7 vs 20.6 months; p=0.02) was also significantly longer in the TACE plus sorafenib group. OS at 1 year and 2 years in TACE plus sorafenib group and TACE alone group were 96.2% and 82.7% and 77.2% and 64.6%, respectively. There were no unexpected toxicities.

Conclusion TACE plus sorafenib significantly improved PFS over TACE alone in patients with unresectable HCC. Adverse events were consistent with those of previous TACE combination trials.

Trial registration number NCT01217034.

INTRODUCTION

Transarterial chemoembolisation (TACE) has shown survival benefits in patients with intermediate stage hepatocellular carcinoma (HCC).¹

Significance of this study

What is already known about this subject?

- Three trials conducted previously (Korea-Japan post-TACE trial, SPACE trial and TACE-2 trial) comparing PFS or TTP in TACE plus sorafenib with that in TACE alone could not show any clinical benefit by the addition of sorafenib to TACE.
- It was demonstrated in the exploratory analysis of Korea-Japan post-TACE trial that longer treatment duration (31 weeks) of sorafenib in Korean patients is associated with significantly favourable TTP (HR=0.38, 95% CI, 0.18 to 0.81) as compared with TTP (HR=0.94, 95% CI, 0.75 to 1.19) in Japanese patients with shorter sorafenib treatment duration (16 weeks). Similarly, in SPACE trial, it was also shown that longer treatment duration with sorafenib in Asian patients (33.6 weeks) is associated with more favourable HR of TTP (HR=0.72, 95% CI, 0.457 to 1.135, p=0.078) than HR of TTP (HR=0.865, 95% CI, 0.576 to 1.300, p=0.243) in Non-Asian patients with shorter treatment duration (26.0 weeks). The design of these previous trials was based on RECIST1.1 or modified RECIST as a definition of the progression. Therefore, when tumour status becomes progression per RECIST1.1/mRECIST such as appearance of intrahepatic new lesions, protocol treatment had to be terminated at the time of progression, leading to a short treatment duration of sorafenib.

Most clinical practice guidelines, including those of the Barcelona Clinic Liver Cancer (BCLC),² the European Association for the Study of the Liver,³ the American Association for the Study of Liver Diseases (AASLD),^{4,5} the Asian Pacific Association

Significance of this study

What is already known about this subject?

- ▶ However, even when intrahepatic new lesion appears, repeated transarterial chemoembolisation (TACE) is still effective in intermediate stage hepatocellular carcinoma (HCC) in the routine clinical practice. In other words, progression per Response Evaluation Criteria in Solid Tumors (RECIST) 1.1/modified RECIST (mRECIST) does not always mean TACE treatment failure or does not suggest switching to next line of treatment in case of intermediate stage HCC different from advanced stage HCC, for which only systemic therapy is performed. Since intrahepatic metastasis/multicentric occurrence is a natural tumour biology in HCC, different endpoint to prolong the treatment duration may be necessary to demonstrate the clinical benefit of the addition of sorafenib. Authors felt that RECIST or mRECIST are not adequate for progression registration in the TACE realm and that we used a different progression endpoint, which was based on Response Evaluation Criteria in Cancer of the Liver (RECICL).
- ▶ In addition, pre-treatment with molecular targeted agent before TACE had been proven in preclinical studies to normalise tumour vessels and suppress hypoxia (caused by TACE) inducible factor-1 α , upregulating vascular endothelial growth factor, platelet-derived growth factor, or angiopoietin 2, which further enhance the HCC progression. Previous trials failed to show progression-free survival (PFS)/time to progression benefit due to the lack of pre-treatment of sorafenib as well as short treatment duration of sorafenib based on inadequate trial design for adjusting the TACE combination trial.

What are the new findings?

- ▶ In this trial, RECIST1.1 or mRECIST was not used as a response evaluation criterion, but RECICL was used. PFS based on untreatable (unTACEble) progression plus death was defined as a trial endpoint based on a line of routine clinical practice in intermediate stage HCC. In this trial, intrahepatic new lesion was not regarded as a 'progression' and trial was continued until newly defined progression, resulting in very long treatment duration with sorafenib (38.7 weeks) as compared with previous trials (17–21 weeks).
- ▶ Pre-treatment with sorafenib 2–3 weeks before initial TACE procedure was introduced in this trial, resulting in significantly longer interval between each TACE procedure in TACE plus sorafenib group than that in TACE alone group. In Sorafenib or Placebo in combination with TACE for intermediate stage hepatocellular carcinoma (SPACE) trial, pre-treatment with sorafenib before TACE was already introduced, but treatment duration before TACE was only 3–7 days, which is a bit different trial design as compared with current trial design.
- ▶ This trial met its primary endpoint: PFS in TACE plus sorafenib group showed significantly longer (25.2 months) than that in TACE alone group (13.5 months) (HR, 0.59; 95% CI, 0.41 to 0.87). Furthermore, time to extrahepatic spread (EHS), time to vascular invasion and time to stage progression were significantly longer in TACE plus sorafenib group than those in TACE alone group.
- ▶ This is the first ever positive trial of TACE in combination with a molecular targeted agent, sorafenib, in patients with unresectable HCC without vascular invasion or EHS. Thus, the combination of TACE plus sorafenib can improve clinical outcomes and may be a choice of treatment in patients with unresectable HCC.

How might it impact on clinical practice in the foreseeable future?

- ▶ This trial clearly showed that TACE in combination with sorafenib should not be terminated at the point of intrahepatic tumour progression, when TACE is still deemed effective. The results in this trial also suggest that pre-treatment of sorafenib before TACE and continued use of sorafenib after TACE will prolong the PFS and prolong the interval between each TACE session, providing the prevention of liver function deterioration often caused by TACE repetition. Improvement of PFS and preservation of liver function will eventually lead to the prolongation of overall survival. However, further large-scaled validation randomised controlled trial is warranted to establish the real benefits of the combination use of targeted agents with TACE.

for the Study of the Liver⁶ and the Japan Society of Hepatology (JSH),⁷ have recommended the use of TACE for these patients, with TACE becoming the standard of care. Because of the high tumour recurrence rate after TACE, this procedure is usually repeated many times. However, the repetition of TACE may lead to deterioration of liver function, resulting in poor patient prognosis.⁸ Furthermore, TACE increases tumour hypoxia, leading to the upregulation of hypoxia inducible factor-1 α (HIF-1 α).^{9–11} Increased HIF-1 α , in turn, upregulates the expression of vascular endothelial growth factor (VEGF) and platelet-derived growth factor (PDGF) and increases tumour angiogenesis.^{9–11} That is, administration of TACE to patients with unresectable HCC has been shown to lead to a spike in the intratumoural concentration of VEGF, suggesting that blockade of VEGF receptors may prevent the effects of a surge in proangiogenic factors.^{11–12} A preclinical model has shown that the combination of antiangiogenic therapy with TACE reduces tumour volume and vessel density, as well as prolonging survival, when compared with TACE alone.¹²

Sorafenib is an oral multikinase inhibitor targeting, among others, VEGF receptor, RAF and PDGF receptor, thereby

exerting both antiangiogenic and direct antitumour effects. This activity profile suggests that sorafenib may suppress the surge of proangiogenic factors after TACE. Sorafenib has been shown to significantly prolong overall survival (OS) in patients with advanced HCC, resulting in a significant advance in the treatment of this disease.^{13–14} Sorafenib has since been approved worldwide and has become the standard treatment for patients with advanced unresectable HCC. Because TACE and sorafenib have each been shown to prolong survival in patients with unresectable HCC,^{13–17} their combination may improve clinical outcomes.^{18–21} To date, however, all five randomised controlled trials testing combinations of TACE with molecular targeted agents such as sorafenib, brivanib and orantinib have failed to show clinical benefits of these combinations.^{22–27}

Trials assessing the combination of TACE plus sorafenib in patients with unresectable HCC have yielded inconsistent results. Several single-arm trials have shown that this combination is safe, effective and feasible in patients with unresectable HCC.^{19–21} In contrast, three previous randomised controlled trials failed to show the benefits of this combination of TACE plus sorafenib.^{22–24} Subgroup analyses of two of these trials,

however, the Japan-Korea Post-TACE trial and the SPACE trial, suggested that longer duration of treatment with sorafenib in combination with TACE may improve clinical outcomes.^{22 23} Based on the three negative clinical trials, the parameter TACE-specific progression was defined as a study outcome in the current trial. The present study reports the results of the TACE Therapy In Combination with Sorafenib as compared with TACE alone in patients with hepatocellular carcinoma (TACTICS) trial, comparing the safety and efficacy of the combination of TACE plus sorafenib with TACE alone in patients with unresectable HCC who were not candidates for resection or ablation.

PATIENTS AND METHODS

Patients

This randomised, open label, multicentre trial was conducted at 33 institutions in Japan and included patients who had been diagnosed with unresectable HCC by biopsy, cytology or diagnostic imaging, such as dynamic CT or MRI, according to the criteria of the AASLD. Patients were included if they were aged ≥ 20 years, had a life expectancy ≥ 12 weeks and had tumours confined to the liver without vascular invasion/extrahepatic spread (EHS) and treatable by TACE. Maximum tumour diameter was 10 cm, and maximum number of nodules was 10. The maximum number of TACE procedures allowed before enrolment was 2, with at least 6 months since prior TACE. All of those patients who had a history of prior TACE achieved complete response by previous TACE, but recurred more than 6 months later at the study entry. All patients had an Eastern Cooperative Oncology Group (ECOG) performance status of 0 or 1, Child-Pugh scores ≤ 7 points and adequate organ function. Prior resection or ablation was allowed; however, any prior systemic therapy was not allowed.

Patients were excluded if they had another previous or current malignant tumour, except for early-stage cancer with low risk of recurrence or a malignant tumour curatively treated > 3 years prior to enrolment with no recurrence. Patients were also excluded if they had cardiac disease or a serious and active infection, except for hepatitis B virus (HBV) and hepatitis C virus (HCV). Patients with diffuse tumour lesions, extrahepatic metastases, vascular invasion, hepatic encephalopathy, uncontrolled ascites or pleural effusion were also excluded, as were patients previously treated for advanced HCC, including systemic chemotherapy, and those treated with CYP3A4-inducing agents. Patients with uncontrollable ascites or decompensated liver function were also excluded.

All patients provided written informed consent prior to enrolment.

Randomisation and masking

Patients were randomised 1:1 to treatment with TACE plus sorafenib or TACE alone. Randomisation was performed centrally via an interactive Web response system involving a computer-generated sequence and Electric Data Capture System Software (Viedoc, Uppsala, Sweden). Patients were stratified by study centre, by their meeting or not meeting Milan criteria (one nodule ≤ 5 cm in diameter or ≤ 3 nodules ≤ 3 cm in diameter) and by number of prior TACE sessions (0 vs 1–2).

Treatment protocol

Patients randomised to TACE plus sorafenib were started on 400 mg/day sorafenib 2–3 weeks prior to first TACE to confirm tolerability to sorafenib, to normalise tumour neovasculature for efficient TACE response and to suppress the VEGF increase

after the TACE procedure. Sorafenib was discontinued for 2 days before and 2 days after each TACE session.

TACE in both groups consisted of intra-arterial injection of lipiodol plus epirubicin or miriplatin,²⁸ followed by injection of an embolic agent (Gelpart) to interrupt blood flow. Selection of anticancer agent (epirubicin or miriplatin) was decided by the sites/investigators; however, the same agent must be used at the repeated TACE sessions. Therefore, sites were included in the one of the stratification factors to avoid any imbalance. When necessary in treating very large tumours, split TACE was allowed within 4–6 weeks of the first TACE session. First image was taken 4 weeks after split TACE was performed. Repeat TACE with the same anticancer agent was recommended when the two-dimensional measurement of the viable lesion was $> 50\%$ that of the baseline tumour. TACE was repeated for intrahepatic new lesions measuring > 10 mm, which show arterial enhancement with venous washout. If venous washout is not associated with arterial enhancement even in lesions > 10 mm, TACE was waited until the time venous washout is confirmed.

Patients in the TACE plus sorafenib group resumed taking 400 mg/day sorafenib 3 days after TACE. If this dose was tolerated, dose increases were allowed to 800 mg/day in a stepwise manner at the discretion of the investigator. At the same dose before re-TACE sorafenib was started after the on-demand TACE. The dose of sorafenib in patients who experienced adverse events (AEs) due to this agent was reduced to 400 mg/day or 400 mg every other day, with patients requiring further dose reduction undergoing dose interruption or discontinued from sorafenib treatment.

Beginning 4 weeks after TACE, tumours were assessed by dynamic CT or MRI every 8 weeks, with tumour marker tests performed at the same times. Treatment was continued until untreatable (unTACEable) progression, progression to meet the TACE refractoriness criteria, unacceptable toxicity or withdrawal of consent.

Study endpoints

The co-primary endpoint of this trial consisted of progression-free survival (PFS)/OS, an endpoint determined using a gate-keeping strategy for multiplicity adjustments, such that OS was calculated only if PFS was statistically significant.²⁹ PFS was defined as the time from randomisation to progression or death from any cause, and OS was defined as the time from randomisation to death from any cause.

Progression in this trial was determined as untreatable (UnTACEable) progression, defined as the inability of a patient to further receive or benefit from TACE for reasons that included intrahepatic tumour progression (25% increase vs baseline) according to Response Evaluation Criteria in Cancer of the Liver (RECICL),³⁰ transient deterioration of liver function to Child-Pugh C right after TACE, macrovascular invasion (MVI) or EHS. When ascites is not controllable or liver function is decompensated, TACE was not performed in both arms since TACE is contraindicated in such cases. In this trial, new intrahepatic lesions were NOT regarded as 'progressive disease', as they are indicative of the natural tumour biology of HCC and do not imply treatment failure or moving to the next line of treatment. Therefore, neither Response Evaluation Criteria in Solid Tumors (RECIST) 1.1 nor modified RECIST criteria were used in this trial. Rather, tumour response was evaluated using RECICL.³⁰ Lipiodol retention area in the nodule after 1 month following TACE was regarded as necrosis.³¹ Therefore, complete tumour response to TACE was defined as complete lipiodol retention

within the nodule and partial tumour response was defined as 50%–100% lipiodol retention within the nodule throughout the trial. Progression in this trial also included progression that met the JSH criteria for TACE failure/refractoriness,³² defined as two or more consecutive insufficient responses of the treated tumour (viable lesion >50%) or two or more consecutive unnumbered increases in liver tumour number, even after changing the chemotherapeutic agents and/or reanalysis of the feeding artery, as determined by CT/MRI 8 weeks after adequate performance of selective TACE.

Secondary endpoints of this trial included time to untreatable (unTACEable) progression (TTUP), defined as the time from randomisation to Child-Pugh grade C, intrahepatic tumour progression (>25% increase vs baseline), with new lesions NOT defined as tumour progression, MVI or EHS (diameter >10 mm); and time to progression (TTP), defined as the time from randomisation to unTACEable progression or progression to TACE failure/refractoriness. Other secondary endpoints included objective response rate (ORR) after the first session of TACE; serum concentrations of the tumour markers AFP, AFP-L3 and PIVKA-II; and safety.

Statistical methods

All statistical analyses were performed using the data obtained by the cut-off date of 2 November 2017, at which time point 118 patients had experienced disease progression. Efficacy data were analysed on an intention-to-treat basis. To adjust for multiple comparisons, hierarchical statistical testing was performed in a pre-specified fixed-order, first for PFS and then for OS. OS was formally compared only when the difference in PFS was significant ($p < 0.15$). The primary endpoints were compared using stratified log-rank tests. The Kaplan-Meier method was used to estimate survival curves, from which medians were calculated. HRs and CIs were estimated using

Cox proportional hazards models. Safety analysis included all patients who received at least one dose of the assigned treatment, irrespective of eligibility or duration of treatment. All statistical analyses were performed using SAS (SAS Institute, Cary, NC, USA), V.9.4.

We originally designed this trial such that a total of 125 TTUP events would give the trial an 80% power to detect an HR of 0.71 with a one-sided alpha of 0.15. The required number of events was anticipated to be reached by monitoring 168 patients. In November 2015, with the investigators and statisticians still blinded to all patient data, the protocol was amended due to slow recruitment. The primary endpoint was amended to the co-primary endpoints, PFS and OS, because extension of the recruitment period made it possible to observe sufficient numbers of PFS and OS events, not only TTUP events that do not require long-term follow-up. As such, a total of 125 PFS and OS events were required to maintain an 80% power to detect a HR of 0.71 with a one-sided alpha of 0.15.

RESULTS

Patient characteristics

A total of 197 patients were screened from February 2011 to March 2016; of these, 156 met the inclusion criteria and were enrolled and randomised. In contrast, 41 patients did not meet the inclusion criteria, for reasons that included Child-Pugh score, prior TACE number and laboratory test results (figure 1). Of the 156 enrolled patients, 60 (38%) were categorised as BCLC stage A, with most having a single large (>5 cm) unresectable tumour and regarded as good candidates for TACE. Similarly, 18 patients (12%) were categorised as BCLC stage C because they had performance status 1, but without EHS or vascular invasion. The remaining 78 patients (50%) were categorised as BCLC stage B.

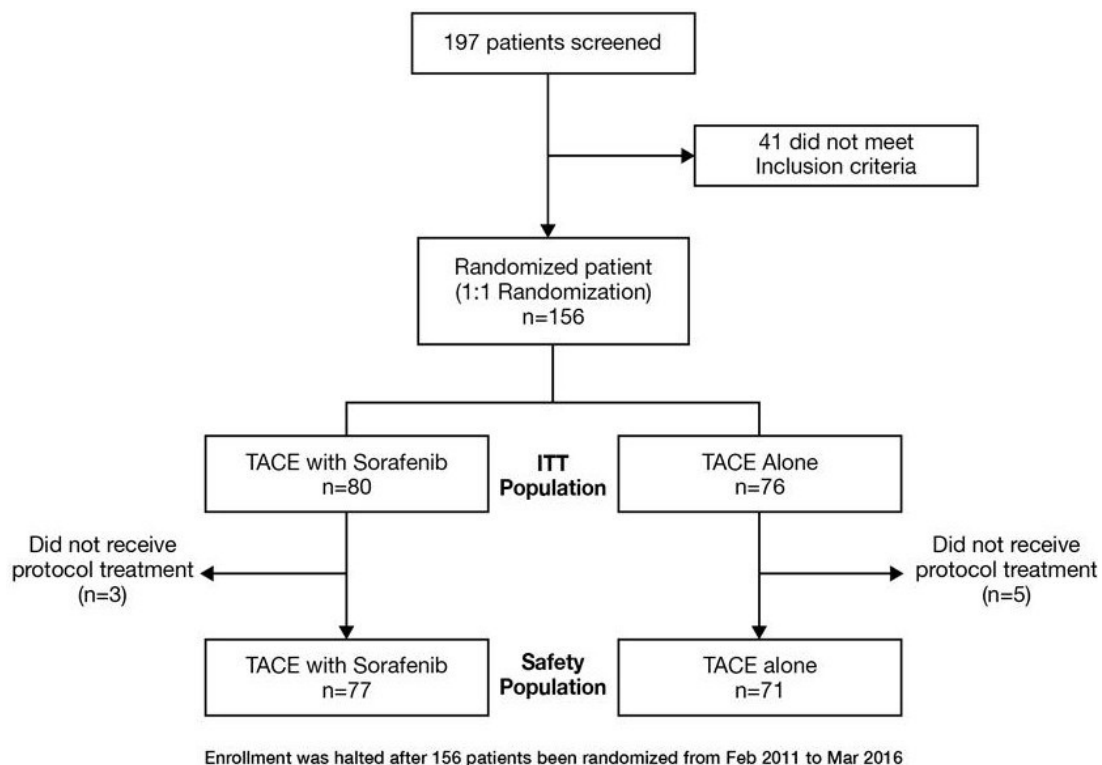


Figure 1 Patientflow chart (consort diagram). TACE, transarterial chemoembolisation.

Table 1 Baseline demographic and clinical characteristics of patients enrolled in this study results reported as N (%), unless otherwise indicated

Characteristic	TACE plus sorafenib (n=80)	TACE alone (n=76)
Age, median (range), years	72.0 (36–85)	73.0 (53–86)
Sex		
Male	63 (78.8)	55 (72.4)
Female	17 (21.2)	21 (27.6)
Performance status		
0	71 (88.8)	67 (88.2)
1	9 (11.3)	9 (11.8)
Aetiology		
Hepatitis B	10 (12.5)	2 (2.6)
Hepatitis C	38 (47.5)	53 (69.7)
Non-B Non-C	32 (40.0)	21 (27.6)
Child-Pugh score		
5	64 (80.0)	54 (71.1)
6	15 (18.8)	17 (22.4)
7	1 (1.3)	5 (5.6)
Ascites	0 (0)	0 (0)
Treatment with diuretics	10 (12.5)	9 (11.8)
AFP		
<200 ng/mL	64 (80.0)	60 (78.9)
≥200 ng/mL	16 (20.0)	16 (21.1)
Tumour burden		
Within Milan criteria	28 (35.0)	35 (46.1)
Outside Milan criteria	52 (65.0)	41 (53.9)
Up to T7 criteria		
Within	54 (67.5)	50 (65.8)
Outside	26 (32.5)	26 (34.2)
BCLC stage		
A	27 (33.8)	33 (43.4)
B	44 (55.0)	34 (44.7)
C	9 (11.3)	9 (11.8)
Prior TACE		
0	45 (56.3)	48 (63.2)
1–2	35 (43.8)	28 (36.8)

BCLC, Barcelona Clinic Liver Cancer; TACE, transarterial chemoembolisation.

In all, 80 patients were randomised to the TACE plus sorafenib group and 76 to the TACE alone group; of these, three and five patients, respectively, were withdrawn prior to protocol treatment. The three patients in the TACE plus sorafenib and two of the five patients in the TACE alone group did not receive protocol treatment (TACE) because of worsening liver function; the other three patients in the TACE alone group were withdrawn because they were found not to meet eligibility criteria after randomisation. The baseline demographic and clinical characteristics were similar in the TACE plus sorafenib and TACE alone groups (table 1).

Doses of study drugs

At a median follow-up of 122.3 weeks, the median duration of sorafenib treatment in the TACE plus sorafenib group was 38.7 weeks (range, 0.3–245.9 weeks), and the mean±SD duration of sorafenib treatment was 57.1±53.3 weeks. Only in 42 of 80 patients in combination group, sorafenib dose was increased up to 800 mg after initial TACE procedure; however, sorafenib dose was eventually reduced during the protocol treatment in all of these 42 patients and also other patients who do not increase

the sorafenib dose up to 800 mg due to AEs, patients' requests or physician's decision. The median actual daily dose of sorafenib was 355.2 mg (range, 86.8–792.9 mg), and the mean±SD actual daily dose of sorafenib was 353.6±172.0 mg. The median interval between TACE sessions was 21.1 weeks (range, 15.8–39.0 weeks) in the TACE plus sorafenib group and 16.9 weeks (range, 10.1–32.1 weeks) in the TACE alone group (p=0.018).

Efficacy outcomes

The first co-primary endpoint, median PFS, specifically defined for TACE-specific PFS based on unTACEable progression was significantly longer in the TACE plus sorafenib than in the TACE only group (25.2 vs 13.5 months; HR=0.59; 95% CI, 0.41 to 0.87; p=0.006; figure 2A). In TACE naïve group, there was no statistical difference between median PFS based on unTACEable progression in the combination group and that in TACE alone group (HR=0.741; 95% CI, 0.45, 1.22; p=0.23) (online supplementary figure 1A). However, in prior TACE 1–2 group, median PFS based on unTACEable progression in the combination group was significantly longer than that in TACE alone group (HR=0.570; 95% CI, 0.33 to 0.99; p=0.04) (online supplementary figure 1B). The second co-primary endpoint, median OS, was not analysed because only 92 of 125 (73.6%) of OS events were mature on the study cut-off date of 2 November 2017. Independent review committee declared that OS was not mature at the PFS manuscript submission on 15 September 2019; therefore, final analysis could not have been performed yet. However, survival rates in TACE plus sorafenib group and TACE alone group were 96.2% and 82.7% at 1 year and 77.2% and 64.6% at 2 years at the last data cut-off (2 November 2017), respectively, although statistical analysis was not allowed based on the protocol regulation.

Analysis of secondary efficacy outcomes showed that median TTUP (26.7 vs 20.6 months; HR=0.57; 95% CI, 0.36 to 0.92; p=0.02; figure 2B) and median TTP (26.7 vs 16.4 months; HR=0.54; 95% CI, 0.35 to 0.83; p=0.005; figure 2C) were also significantly longer in the TACE plus sorafenib than in the TACE alone group. Exploratory subgroup analyses showed that PFS based on unTACEable progression was longer in all subgroups treated with TACE plus sorafenib than with TACE alone (figure 3). In contrast, tumour responses, disease control rate (DCR) and ORR 4 weeks after the first TACE session were similar in the two groups (table 2).

Pattern of unTACEable progression

The patterns and frequency of progression did not differ significantly in the TACE plus sorafenib and TACE alone groups (online supplementary table 1). However, in the analysis considering the timing of event occurrence, time to MVI (31.3 months vs 4.0 months; HR=0.26, 95% CI, 0.09 to 0.75, p=0.008) and time to EHS (15.7 months vs 6.9 months; HR=0.21, 95% CI, 0.006 to 0.70, p=0.006) were significantly longer in the TACE plus sorafenib than in the TACE alone group (online supplementary figure 2). Time to stage progression, which was defined as appearance of MVI and/or EHS was also significantly longer in the TACE plus sorafenib (22.5 months) than in the TACE alone (6.3 months) group (HR=0.31, 95% CI, 0.15 to 0.63, p=0.001) (online supplementary figure 3). Discontinuation of protocol treatment due to transient liver function deterioration to Child-Pugh C right after TACE was seen only in four cases; two in combination group and two in TACE alone group. Liver function in those

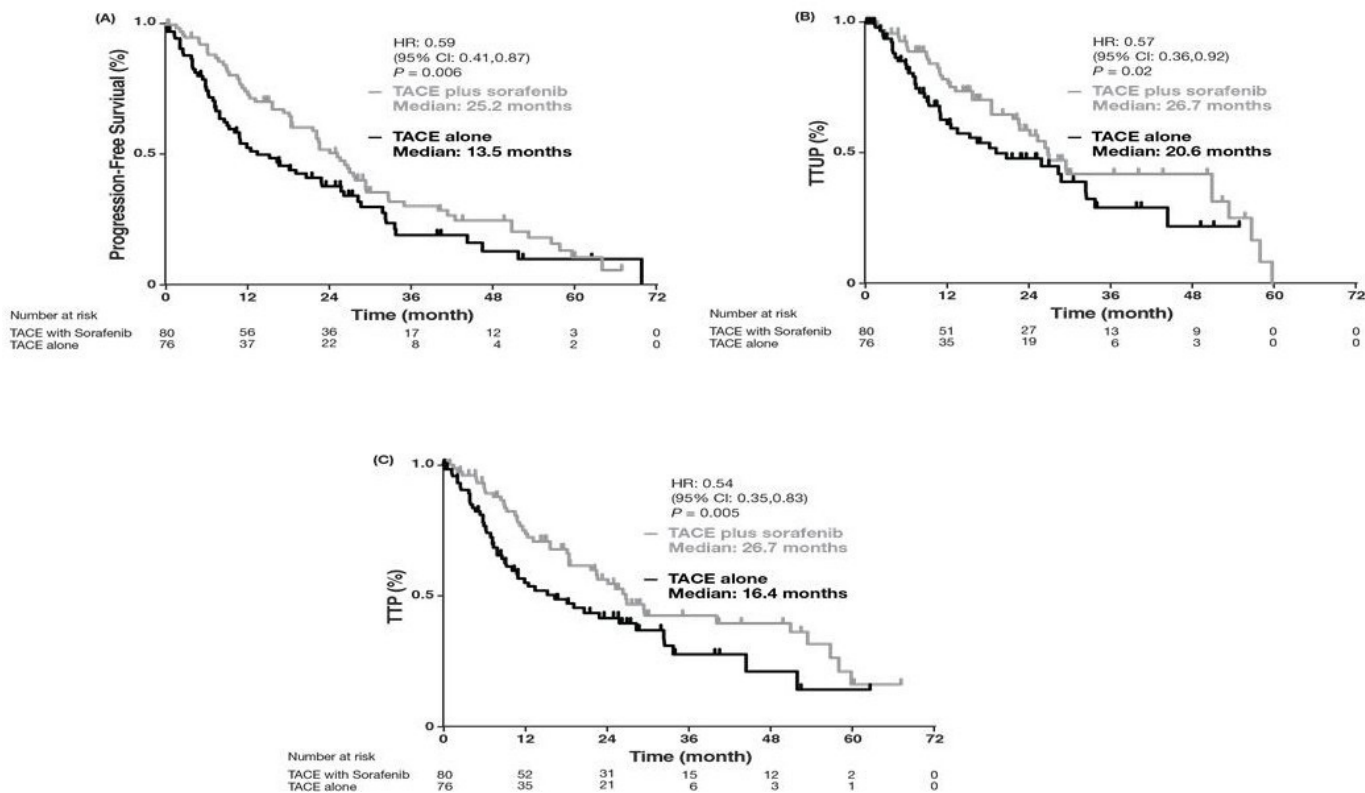


Figure 2 Kaplan-Meier plots of median (A) progression-free survival, (B) TTUP and (C) TTP in the TACE plus sorafenib and TACE alone groups. TACE, transarterial chemoembolisation; TTP, time to progression; TTUP, time to untreatable (unTACEable) progression.

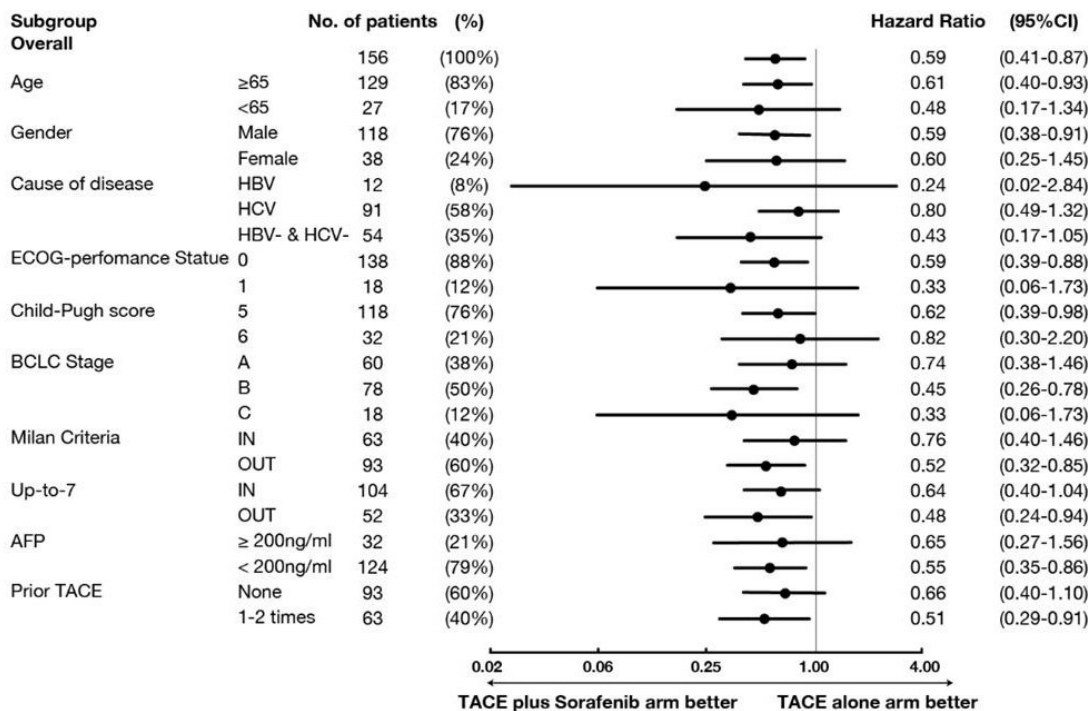


Figure 3 Forest plot of PFS in subgroups of patients treated with TACE plus sorafenib and TACE alone. PFS, progression-free survival; TACE, transarterial chemoembolisation.

Table 2 Tumour responses 4 weeks after first TACE in patients randomised to the TACE plus sorafenib and TACE alone groups results reported as N (%)

	TACE plus sorafenib (n=80)	TACE alone (n=76)	P value*
Best response			0.77
Complete response (CR)	23 (28.8)	21 (27.6)	
Partial response (PR)	34 (42.5)	26 (34.2)	
Stable disease (SD)	10 (12.5)	12 (15.8)	
Progressive disease (PD)	2 (2.5)	3 (3.9)	
Not evaluable	11 (13.8)	14 (18.4)	
ORR (CR+PR)	57 (71.3)	47 (61.8)	0.23
DCR (CR+PR+SD)	67 (83.8)	59 (77.6)	0.42

*By two-sided Fisher's exact tests.

DCR, disease control rate; ORR, objective response rate; TACE, transarterial chemoembolisation.

four patients returned to Child-Pugh B seven or eight later, but dropped out from the trial due to protocol specification.

Safety outcomes

Table 3 shows the treatment emergent AEs with frequency $\geq 10\%$ in either group and their corresponding Grades 3 and 4 AEs within 4 weeks after first TACE. All grade AEs more frequent in the TACE plus sorafenib group than in the TACE alone group included hand-foot skin reaction (HFSR) (53.2% vs 0%), hypertension (51.9% vs 39.4%), increased lipase (49.4% vs 25.4%), fatigue (24.7% vs 9.9%), diarrhoea (14.3% vs 0%), erythema multiforme (11.7% vs 0%), weight loss (11.7% vs 2.8%) and hoarseness (11.7% vs 0%). Grade 3 AEs more frequent in the TACE plus sorafenib group than in the TACE alone group included thrombocytopenia (13.0% vs 2.8%), HFSR (5.2% vs 0%), hypertension (10.4% vs 4.2%), increased lipase (14.3% vs 2.8%), increased amylase (7.8% vs 1.4%), neutropaenia (5.2% vs 0%), fatigue (2.6% vs 0%),

diarrhoea (2.6% vs 0%) and erythema multiforme (2.6% vs 0%). AEs more frequent in the TACE plus sorafenib group were likely due to the effects of sorafenib.

DISCUSSION

The TACTICS trial was a randomised, open label, multicentre trial comparing the safety and efficacy of the combination of TACE plus sorafenib with TACE alone in patients with unresectable HCC who were not candidates for resection or ablation. This study, which assessed one of the co-primary endpoints, median PFS based on unTACEable progression, found that the combination of TACE plus sorafenib resulted in significantly longer PFS based on unTACEable progression than TACE alone (25.2 vs 13.5 months; HR=0.59; 95% CI, 0.41 to 0.87; p=0.006). Moreover, times to vascular invasion, EHS and stage progression were much longer in the TACE plus sorafenib than in the TACE alone group, indicating that the combination of TACE plus sorafenib markedly prevented the progression of intermediate to advanced stage HCC. In addition, the interval between pairs of TACE sessions was significantly longer in the TACE plus sorafenib than in the TACE alone group, suggesting that TACE-associated deterioration of liver function is lower in the TACE plus sorafenib than in the TACE alone group. ORR was relatively high in both the TACE plus sorafenib (71.3%) and TACE alone (61.8%) groups, probably because of the high response to TACE performed in a superselective manner. Because OS events are not yet mature in this trial, the second co-primary endpoint, median OS, could not be analysed.

The combination of TACE plus sorafenib was clinically feasible and safe. Many AEs were more frequent in the TACE plus sorafenib than in the TACE alone group, with these AEs likely to result from treatment with sorafenib. None of the AEs in either group was unexpected, and rates of Grades 3 and 4 AEs were relatively low.

Table 3 All-grade treatment-emergent AEs within 4 weeks after first TACE with frequency $\geq 10\%$ in either group and corresponding Grades 3 and 4 AEs

n (%)	TACE plus sorafenib (n=77)				TACE alone (n=71)			
	All grade	Grades 1–2	Grade 3	Grade 4	All grade	Grades 1–2	Grade 3	Grade 4
Elevated AST	72 (93.5)	50 (64.9)	17 (22.1)	5 (6.5)	65 (91.5)	50 (70.4)	14 (19.7)	1 (1.4)
Elevated ALT	69 (89.6)	50 (64.9)	18 (23.4)	1 (1.3)	55 (77.5)	42 (59.2)	13 (18.3)	0 (0.0)
Thrombocytopenia	67 (87.0)	57 (74.0)	10 (13.0)	0 (0.0)	53 (74.6)	51 (71.8)	2 (2.8)	0 (0.0)
Elevated bilirubin	55 (71.4)	54 (70.1)	1 (1.3)	0 (0.0)	39 (54.9)	37 (52.1)	2 (2.8)	0 (0.0)
Anaemia	50 (64.9)	49 (63.6)	1 (1.3)	0 (0.0)	35 (49.3)	34 (47.9)	1 (1.4)	0 (0.0)
Hand-foot skin reaction	41 (53.2)	37 (48.1)	4 (5.2)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)
Hypertension	40 (51.9)	32 (41.6)	8 (10.4)	0 (0.0)	28 (39.4)	25 (35.2)	3 (4.2)	0 (0.0)
Elevated lipase	38 (49.4)	26 (33.8)	11 (14.3)	1 (1.3)	18 (25.4)	16 (22.5)	2 (2.8)	0 (0.0)
Elevated serum amylase	32 (41.6)	26 (33.8)	6 (7.8)	0 (0.0)	19 (26.8)	18 (25.4)	1 (1.4)	0 (0.0)
Neutropaenia	29 (37.7)	25 (32.5)	4 (5.2)	0 (0.0)	29 (40.8)	29 (40.8)	0 (0.0)	0 (0.0)
Decreased WBC count	29 (37.7)	28 (36.4)	1 (1.3)	0 (0.0)	26 (36.6)	26 (36.6)	0 (0.0)	0 (0.0)
Malaise	20 (26.0)	20 (26.0)	0 (0.0)	0 (0.0)	9 (12.7)	9 (12.7)	0 (0.0)	0 (0.0)
Fatigue	19 (24.7)	17 (22.1)	2 (2.6)	0 (0.0)	7 (9.9)	7 (9.9)	0 (0.0)	0 (0.0)
Fever	15 (19.5)	14 (18.2)	1 (1.3)	0 (0.0)	18 (25.4)	18 (25.4)	0 (0.0)	0 (0.0)
Anorexia	11 (14.3)	9 (11.7)	2 (2.6)	0 (0.0)	8 (11.3)	7 (9.9)	1 (1.4)	0 (0.0)
Diarrhoea	11 (14.3)	9 (11.7)	2 (2.6)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)
Erythema multiforme	9 (11.7)	7 (9.1)	2 (2.6)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)
Weight loss	9 (11.7)	9 (11.7)	0 (0.0)	0 (0.0)	2 (2.8)	2 (2.8)	0 (0.0)	0 (0.0)
Hoarseness	9 (11.7)	9 (11.7)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)

AEs, adverse events; ALT, alanine aminotransaminase; AST, aspartate aminotransferase; TACE, transarterial chemoembolisation; WBC, white blood cell.

These findings differ markedly from those of previous trials testing the combination of TACE plus sorafenib in patients with HCC. For example, the Japan-Korea post-TACE and SPACE trials reported median TTPs of 5.4 and 5.6 months, respectively, with the criteria for disease progression determined using RECIST 2004 and mRECIST criteria, respectively.^{22 23} The TACE 2 trial reported a median PFS of 8.5 months, with the criteria for disease progression determined using RECIST 1.1 criteria.²⁴ RECIST and mRECIST criteria may not be good measures of capturing the 'progression' of intermediate stage HCC, as regrowth of the original tumour or appearance of a new intrahepatic lesion is not indicative of treatment (TACE) failure and does not suggest the need to move to a next line of treatment.

In contrast, the TACTICS trial reported a median PFS of 25.2 months, with the criteria for progression being unTACEable progression/TACE failure. New intrahepatic lesions in the TACTICS trial were not regarded as progressive disease. The better outcomes observed in the TACTICS trial may be due to the much longer median duration of sorafenib treatment (38.7 weeks), owing to the TACE-specific trial design. In contrast, the median durations of sorafenib treatment in the post-TACE, SPACE, and TACE 2 trials ranged from 17.0 to 21.0 weeks.^{22–24} Another possible explanation would be sorafenib was given 2–3 weeks before initial TACE was performed. This pre-treatment with sorafenib might have led to the normalisation of tumour vessel, resulting in enhancement of TACE efficacy by dense accumulation of lipiodol mixed with anticancer agent and gelatine sponges. In fact, interval between each TACE session was significantly prolonged in TACE plus sorafenib group than TACE alone group. The third explanation would be sorafenib might have stabilised disease progression after each TACE session, leading to a prolonged PFS.

This study, however, had several limitations, including the relatively small number of patients included. Moreover TACE-specific PFS, PFS based on unTACEable progression, which was applied in this trial based on lessons learnt from previous negative trials, had rarely been used in previous trials, except that 'unTACEable progression' was used in the SPACE trial. Actually, this concept that progression after TACE may not mean treatment failure and that treatment could be maintained until untreatable progression was first proposed by Bruix *et al*³³ and therefore tested in SPACE trial. Thus, we believe that this new TACE-specific endpoint should be tested and validated in future TACE combination trials as well.

In summary, this is the first ever positive trial of TACE in combination with a molecular targeted agent, sorafenib, in patients with unresectable HCC, providing important results in the practice. In contrast to the three negative trials, the present trial assessed the efficacy of TACE plus sorafenib using an endpoint more suitable for a TACE combination trial, with this endpoint being consistent with those used in clinical practice. Furthermore, pre-treatment with sorafenib 2–3 weeks before initial TACE was introduced in this trial similar to, but longer period than SPACE trial. Pre-treatment with sorafenib before TACE will enhance the treatment effect of TACE through tumour vessel normalisation leading to homogeneous distribution of lipiodol mixed with anticancer drugs and gelatine sponge particle within the tumours. Furthermore, pre-treatment with sorafenib before initial TACE will suppress the HIF-1 α induced VEGF or PDGFR, which is regarded as the factor responsible for further tumour progression. The results of the TACTICS trial indicate that new intrahepatic lesions may not be regarded as progressive disease or as a reason for halting treatment in TACE combination trials. Thus, we strongly believe that the definition

of treatment success or failure by TACE should use a different criteria from RECIST or mRECIST.

In TACTICS trial, among unTACEable progression transient liver function deterioration to Child-Pugh C was observed only in 4 of 99 patients, which means many patients were discontinued TACE after TACE refractoriness. This event was included in the category of intrahepatic tumour growth >25% or TACE refractoriness in online supplementary table 2. The reason why for this is that the study protocol clearly specified TACE should not be continued once tumour status become refractory to TACE.

In conclusion, this trial clearly showed that the combination of TACE plus sorafenib can improve clinical outcomes and may be a choice of treatment in patients with unresectable HCC without vascular invasion or EHS, who are good candidates for TACE.

Author affiliations

- ¹Department of Gastroenterology and Hepatology, Kindai University Faculty of Medicine, Osaka-Sayama, Japan
- ²Department of Hepatobiliary and Pancreatic Oncology, National Cancer Center Hospital East, Kashiwa, Japan
- ³Department of Gastroenterology and Hepatology, Kurume University School of Medicine, Kurume, Japan
- ⁴Department of Gastroenterology, National Hospital Organisation Sendai Medical Center, Sendai, Japan
- ⁵Department of Gastroenterology and Metabolism, Hiroshima University, Hiroshima, Japan
- ⁶Department of Gastroenterology, Musashino Red Cross Hospital, Tokyo, Japan
- ⁷Department of Gastroenterology and Hepatology, Yamaguchi University Graduate School of Medicine, Ube-Yamaguchi, Japan
- ⁸Department of Gastroenterology and Metabolism, Nagoya City University Graduate School of Medical Sciences, Nagoya, Japan
- ⁹Department of Hepatology and Pancreatology, Kawasaki Medical School, Kurashiki, Japan
- ¹⁰Department of Gastroenterology and Hepatology, Hyogo Cancer Center, Akashi, Japan
- ¹¹Department of Gastroenterology and Hepatology, Nagoya University Graduate School of Medicine, Nagoya, Japan
- ¹²Department of Medicine, Division of Gastroenterology, Jichi Medical University, Shimotsuke, Japan
- ¹³Department of Gastroenterology and Hepatology, Kyoto Prefectural University of Medicine, Kyoto, Japan
- ¹⁴Division of Gastroenterology, Department of Medicine, Social Insurance Tagawa Hospital, Tagawa, Japan
- ¹⁵Department of Gastroenterology and Hepatology, Kagoshima University Graduate School of Medical and Dental Sciences, Kagoshima, Japan
- ¹⁶Department of Liver, Biliary Tract and Pancreas Diseases, Fujita Health University School of Medicine, Aichi, Japan
- ¹⁷Department of Gastroenterology and Hepatology, Nagasaki University Graduate School of Biomedical Sciences, Nagasaki, Japan
- ¹⁸Department of Hepato-Biliary-Pancreatic Surgery, Clinical Research Institute, National Hospital Organization Kyushu Medical Center, Fukuoka, Japan
- ¹⁹Department of Gastroenterology, School of Medicine, Chiba University, Chiba, Japan
- ²⁰Center for Integrated Medical Research, Hiroshima University, Hiroshima, Japan
- ²¹Department of Hepatobiliary and Pancreatic Oncology, National Cancer Center Hospital, Tokyo, Japan
- ²²Department of Medical Oncology, Kyorin University Faculty of Medicine, Mitaka, Japan
- ²³Department of Surgery, National Center for Global Health and Medicine, Tokyo, Japan
- ²⁴Department of Medicine, Shunan Memorial Hospital, Kudamatsu, Yamaguchi, Japan
- ²⁵Department of Molecular and Clinical Cancer Medicine, University of Liverpool, Liverpool, UK
- ²⁶Department of Diagnostic Radiology, National Cancer Center Hospital, Tokyo, Japan

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ORCID iDs

Masatoshi Kudo <http://orcid.org/0000-0002-4102-3474>

Philip James Johnson <http://orcid.org/0000-0003-1404-0209>

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