A combination of portal vein stent insertion and endovascular iodine-125 seed-strip implantation, followed by transcatheter arterial chemoembolization with sorafenib for treatment of hepatocellular carcinoma-associated portal vein tumor thrombus

Shuangxi Li, BS, Baohua Li, MS, Lei Li, MS, Fangyu Xu, BS, Xujun Yang, MS, Wenhui Wang, MD, PhD
Interventional Department, The First Hospital of Lanzhou University, Lanzhou City, China

Abstract

Purpose: This study aimed to assess efficacy of portal vein stent (PVS) insertion and endovascular iodine-125 (125I) seed-strip implantation, followed by transcatheter arterial chemoembolization (TACE) with sorafenib (PVS-125I TACE-S) in patients with hepatocellular carcinoma (HCC)-associated type II or type III portal vein tumor thrombus (PVTT).

Material and methods: A retrospective study was performed on 53 consecutive patients with HCC and type II or type III PVTT, from May 2014 to July 2018. Patients were divided into 2 groups, including group A with 28 patients treated with PVS-125I TACE-S, and group B with 25 patients treated with TACE-S. Primary end-point was overall survival (OS), while secondary endpoints were hepatic function and disease control rate (DCR). Albumin-bilirubin (ALBI) score approach was used for evaluating liver function. Cox regression analysis was applied to identify factors associated with treatment outcomes.

Results: No pre-operative differences were found in ALBI scores between group A and group B (–2.57 ±0.42 vs. –2.61 ±0.38, p = 0.724), or in these scores at 1 month post-operatively (–2.62 ±0.46 vs. –2.20 ±0.59, p = 0.666). However, these scores were significantly different at 3 (–2.17 ±0.59 vs. –1.69 ±0.48, p = 0.007) and 6 (–2.28 ±1.23 vs. –1.47 ±0.31, p = 0.044) months post-operatively. In addition, group A exhibited higher DCR (71.4% vs. 44.0%, p = 0.043) after 6 months of treatment and extended OS duration (11.4 vs. 7.7 months, p = 0.007). A stratified analysis revealed that OS in patients with type II PVTT did not differ significantly (10.4 vs. 10.7 months, p = 0.689), but OS with type III varied significantly (11.5 vs. 7.5 months, p = 0.002). Multivariate analysis revealed that tumor size > 10 cm (p = 0.002) and multiple tumors (p = 0.022) were independent predictors for poor prognosis, whereas PVS-125I TACE-S was predictor for favorable patient’s prognosis (p = 0.040).

Conclusions: PVS-125I TACE-S represents a potentially viable strategy for improving hepatic functionality, DCR, and OS in HCC with type III PVTT compared with TACE-S alone.

Key words: hepatocellular carcinoma, portal vein, iodine-125, transcatheter arterial chemoembolization, sorafenib.

Purpose

Hepatocellular carcinoma (HCC) is diagnosed when the disease has progressed to mid- or late-stage cancer due to lack of early symptoms [1]. Approximately 44.0-62.2% of patients exhibit macroscopic portal vein tumor thrombus (PVTT) when initially diagnosed [2]. The presence of PVTT can result in severe adverse effect on hepatic blood supply, leading to decreased liver function and poor patient’s prognosis. Median survival time of patients with HCC without treatment affected by PVTT is reduced to 2.5-4.0 months, as compared to 10-24 months in those without this comorbidity [3, 4].
At present, however, the optimal means of treating HCC complicated by PVTT remains uncertain. The Barcelona Clinic Liver Cancer (BCLC) group recommends that sorafenib is the therapeutic agent of choice for patients with advanced HCC, irrespective of PVTT status. However, sorafenib hepatocellular carcinoma assessment randomized protocol with BCLC recommendations, suggests that vascular invasion is present in only 38.4% of patients with HCC [5, 6]. As such, only sorafenib may not necessarily improve survival in patients with HCC and PVTT. In addition, studies on Asian populations with vascular invasion or extrahepatic metastasis have suggested that treatment with sorafenib was associated with poor survival time (median, 6.5 months) [6]. Transcatheter arterial chemoembolization (TACE) with sorafenib (TACE-S) exhibited therapeutic synergy when employed in treatment of HCC patients with PVTT, and provided good patency for the main portal vein or sufficient collateral circulation [7]. This combination is associated with some increase in patients’ survival. However, prolongation of survival in patients with main portal vein invasion is limited [6, 8]. Hence, a superior combination strategy is needed.

The use of a portal vein stent (PVS) to reduce portal vein occlusion has been explored in the context of PVTT [9]. Its’ combination with iodine-125 (125I) seed brachytherapy offers a potentially optimal means of improving PVTT management in patients with HCC [10]. PVS insertion and endovascular 125I seed-strip implantation combination with TACE-S had a potential therapeutic effect on HCC PVTT patients, but so far, that has not been fully elucidated [11]. In the present study, a retrospective analysis was performed to assess the efficacy of PVS insertion and endovascular 125I seed-strip implantation, followed by TACE-S (PVS-125I TACE-S) as well as the factors associated with patients’ outcomes.

Material and methods

This was a retrospective analysis of 53 patients with HCC with type II or type III PVTT, from May 2014 to July 2018, who underwent PVS-125I TACE-S or TACE-S. This study received approval from the ethics committee of our institution (No. LDYYLL2019-204), and all participants provided written informed consents.

In all patients, HCC diagnosis was confirmed either via histology or by a combination of two imaging approaches and elevated alpha-fetoprotein, as per the American Association for the study of liver diseases guidelines [11]. PVTT diagnosis was confirmed following an identification of low-attenuation intraluminal mass, partially or completely occluding the portal vein, or a filling defect in the portal vein, which were observed from three-phase dynamic computed tomography (CT) or magnetic resonance imaging (MRI) [12]. Patients were stratified into types I-IV PVTT according to Cheng’s PVTT classification system as follows: type I – tumor thrombi located at the segmental branches of the portal vein or above; type II – tumor thrombi extending and involving the right or left portal vein, type III – the main portal vein trunk involved; type IV – tumor thrombus extending to the main portal vein and the superior mesenteric vein [13].

Study inclusion criteria were as follows: 1. Diagnosis with HCC and either type II or type III PVTT; 2. No previous local treatment of PVTT lesions; 3. The Eastern Cooperative Oncology Group performance status of 0-2; 4. Child-Pugh class A (score 5 or 6) or B (score 7-9). Study exclusion criteria were as follows: 1. Complete portal vein occlusion with a lack of collateral circulation; 2. Bleeding of the esophagus or gastric fundus; 3. Intractable coagulation disorders; 4. Macroscopic hepatic vein tumor thrombi or extrahepatic tumor metastasis; 5. Lack of baseline imaging results.

Treatment schedule, including TACE, sorafenib, TACE-S, PVS-125I TACE-S, and stereotactic body radiotherapy (SBRT) was conveyed in detail to all participants, and the advantages and disadvantages of all treatments were discussed with the patients, who themselves chose the treatment method independently. During this study period, 72 patients received the afore-mentioned treatment, 12 patients with TACE, 5 patients with sorafenib, and 2 patients who chose SBRT were excluded. Finally, 53 patients were included in this study, of these, 28 patients who decided upon the combination of PVS-125I TACE-S formed group A, while the remaining 25 patients who chose TACE-S alone formed group B.

**Portal vein stent insertion and endovascular 125I seed-strip implantation**

The required number (N) of 125I seeds (GMS Pharmaceutical Co., Ltd., Shanghai, China) was determined based upon the length (L) of the obstructed segment of the portal vein in millimeters, using the following formula: \( N = L/4.5 + 2 \) [10]. The estimated radiation doses were roughly 40-50 Gy, as determined at specific dose reference points based on calculations conducted by a computerized treatment planning system (FTT Technology Co., Ltd., Beijing, China). Iodine-125 seeds used in this study were 0.8 mm in diameter and 4.5 ±0.5 mm in length, with 25.9 MBq of radioactivity and 59.4-day half-life, primarily emitting 27.4 and 31.4 keV X-rays and 35.5 keV γ-rays. Given the local tissue half-value thickness of 17 mm, these seeds were associated with an initial dose rate of 0.07 Gy/h. Prior to implantation, seeds were loaded in a linear arrangement in a 4-Fr flexible stiffening cannula (Boston Scientific; Marlborough, MA, USA) to construct 125I seed-strip. With ultrasound guidance, a Neff percutaneous access set (Cook Medical; Bloomington, IN, USA) was then used to puncture the patent second-order branch of the portal vein, following which, both a vascular stent (Bard Peripheral Vascular Inc.; Tempe, AZ, USA) with 12-14 mm diameter and 60-100 mm length and 125I seed-strip were implanted in a sequence. After PVS insertion and endovascular 125I seed-strip implantation, a 3-3 spring coil (Cook Medical; Bloomington, IN, USA) was used for blocking the intra-hepatic puncture. Subsequently, subcutaneous low-molecular-weight heparin (Changshen Biochemical Pharmaceutical Co., Ltd.; Shijiazhuang, China) 4,100 IU was administered twice daily, for a 5-day period. Next, warfarin (Shanghai Sine Pharmaceutical Laboratories Co., Ltd.; Shanghai, China) was administered orally to achieve international normalized ratio of 2.0-2.5.
TACE procedure

In the group A patients, TACE was conducted 3-7 days after PVS insertion and endovascular 125I seed-strip implantation, whereas in the group B patients, this procedure was performed directly. A 5-Fr hepatic-curve catheter (Terumo Corporation; Tokyo, Japan) was placed into the celiac artery and then, hepatic arterial angiography and indirect portography were performed. A 2.7-Fr microcatheter (Progreat™, Terumo Corporation; Tokyo, Japan) was placed into tumor-feeding arteries, and a 5-20 ml of lipiodol (Wh-Medical Apparatus and Instruments Co., Ltd.; Beijing, China) mixed with 50-75 mg/m² doxorubicin hydrochloride (Hisun Pfizer Pharmaceutical Co., Ltd.; Shanghai, China) was injected into the arteries. If the entire 20-ml volume was administered without substantially impairing blood flow in vessels, polyvinyl alcohol (PVA) particles (ALICON Pharmaceutical Science and Technology Co., Ltd.; Hangzhou, China) were used for vessel embolization, and administered until a limited slow flow was evident only. In patients with arterioportal shunts, initial embolization using 350-1,000 μm PVA particles was conducted prior to lipiodol/doxorubicin infusion to ensure shunt occlusion. After a month, the effects of TACE were assessed by contrast-enhanced abdominal CT/MRI, and a next treatment plan was determined.

Sorafenib

Sorafenib (Nexavar®, Bayer HealthCare; Leverkusen, Germany) is a small molecule that inhibits tumor cell proliferation and tumor angiogenesis to achieve therapeutic effect [14]. In this study, all 53 patients received sorafenib. The patients were administered 400 mg twice per day, for 3 to 7 days after TACE treatment, when liver function had stabilized. Sorafenib doses were reduced in many patients suffering from grade 1 or 2 adverse events (AEs), or with serum bilirubin > 34.2 μmol/l. Sorafenib treatment was temporarily halted in patients suffering from grade 3 AEs or hyperbilirubinemia (serum bilirubin > 51.3 μmol/l) events, and was resumed only if AE grades and serum bilirubin levels returned within the acceptable treatment ranges. The patients received sustained-release sorafenib tablets as long as possible, even when disease progression was noted, or until death.

Follow-up and treatment evaluation

Patients’ follow-up was conducted at 1 and 3 months after the procedure, and every 3 months thereafter. Follow-up assessment included a combination of physical examination, laboratory testing, and a contrast-enhanced abdominal CT or MRI. The Common Terminology Criteria for Adverse Events (CTCAE) v. 4.0 assessment was used to rate complications associated with the treatment [15]. Albumin-bilirubin (ALBI) scores were applied as an objective means of gauging liver function, with scores being calculated solely based upon albumin and bilirubin levels [ALBI score = log10 bilirubin (μmol/l) / 0.66 + (albumin (g/l) x -0.0852)] [16]. The modified response evaluation criteria in solid tumors (mRECIST) criteria for HCC were applied for tumor responses, with possible responses consisting of complete response (CR), partial response (PR), stable disease (SD), or progressive disease (PD) (Figure 1). A disease control rate (DCR) after 6 months of treatment was calculated based on the following formula: (CR + PR + SD)/total cases × 100% [17]. Overall survival (OS) was the primary end-point defined as the time duration between start of treatment and either death or last follow-up.

Statistical analyses

SPSS 21.0 (SPSS Inc.; Chicago, IL, USA) was used for all analyses. Quantitative results were expressed as means ± standard deviation, and t-test was applied to compare these values. Qualitative results were expressed as a number (%), and Pearson χ² test or Fisher’s exact test were used to compare these results, whenever appropriate. Paired-samples t-test was used for comparison of ALBI scores in the patients of group A. Survival was analyzed via Kaplan-Meier approach and using log-rank tests. P < 0.05 was the significance threshold. Cox univariate and multivariate regression analyses were used to assess the relationship between specific factors and treatment outcomes.

Results

Baseline information

Baseline characteristics of the 53 patients are shown in Table 1, with no significant clinically relevant differences between groups. For the patients of group A, on average, 18.2 ±1.7 (range, 16-20) 125I seeds were implanted.

Complications and clinical outcomes

The most frequent AEs in the group A during PVS insertion and endovascular 125I seed-strip implantation were fever (3 patients, 10.7%), abdominal pain (20 patients, 71.4%), and a transient decrease in liver functionality (7 patients, 25%). Patients who experienced these AEs were recovered after conservative management. Furthermore, no serious procedure-related complications, such as abdominal hemorrhage, biloma, stent migration, or puncture site bleeding, were observed. Table 2 shows AEs occurring immediately during TACE-S, including nausea or vomiting, fever, and abdominal pain as short-term side effects in group A in 5 (15.5%), 3 (4.6%), and 7 (23%) patients, respectively, and in group B in 5 (15.5%), 3 (4.6%), and 7 (2.3%) cases, respectively. All of these AEs were resolved after symptomatic treatments. Long-term side effects, such as fatigue, diarrhea, hypertension, hand-foot syndrome, alopecia, pruritus, rash or desquamation, voice change, anorexia, and abscess occurred in group A in 9 (32.1%), 12 (42.9%), 10 (35.7%), 16 (57.1%), 2 (7.1%), 4 (14.3%), 3 (10.7%), 1 (3.6%), 4 (14.3%), and 1 (3.6%) patients, respectively, and in group B in 8 (32.0%), 10 (40.0%), 9 (36.0%), 13 (52.0%), 2 (8.0%), 5 (20.0%), 3 (12.0%), 0 (0%), 3 (12.0%), and 0 (0%) patients, respectively. Usually, all of these long-term side-effects began 1-2 weeks after the treatment and alleviated after sorafenib dose adjustment, short-term interruption, medications, or
Fig. 1. A) Results from a 63-year-old male patient who had HCC with type III PVTT. Contrast-enhanced CT scan exhibiting a hepatic arterial phase hyper-attenuation lesion at segment 6. B) Segment 7 and PVTT extending to the main portal vein. C) Histopathological examination of the biopsy tissue sampling showed obvious cell atypia, different sizes and shapes, and deep nuclear staining. Some of the cells were acidophilic, with diffuse arrangement and necrotic tissue around, consistent with the morphological features of HCC. D) Hematoxylin and eosin stain, 100× results from the 3-month follow-up following the combination of PVS-125I TACE-S, lipiodol accumulation in the tumor. E) The observed satisfactory patency of the stent. F) Results from the 6-month follow-up, with the treated lesion decreased in size.
puncture drainage. TACE-S-associated AEs showed no significant difference between the two groups ($p > 0.05$). All patients had no procedure-related mortality.

Pre-operative ALBI scores were not significantly different between group A and group B ($-2.57 \pm 0.42$ vs. $-2.61 \pm 0.38$, $p = 0.724$), nor did these scores differ at 1 month post-operatively ($-2.62 \pm 0.46$ vs. $-2.20 \pm 0.59$, $p = 0.666$). However, these scores were significantly different at 3 months ($-2.17 \pm 0.59$ vs. $-1.69 \pm 0.48$, $p = 0.007$) and 6 months ($-2.28 \pm 1.23$ vs. $-1.47 \pm 0.31$, $p = 0.044$) post-operatively (Table 3). DCR was 71.4% in group A and 44.0% in group B ($p = 0.043$) after 6 months of treatment.

The median survival of group A was 11.4 ± 0.7 months (range, 10.1-12.7), while that of group B was 7.7 ± 0.8 months (range, 6.1-9.3 months) ($p = 0.007$). A stratified analysis demonstrated that the median survival in those with type II PVTT was 10.4 ± 2.0 months (range, 6.5-14.4 months) and 10.7 ± 1.7 months (range, 7.3-14.1 months) in group A and group B, respectively ($p = 0.689$), whereas in patients with type III PVTT, the survival time was 11.5 ± 0.8 months (range, 10.0-12.9 months) and 7.5 ± 0.9 months (range, 5.8-9.2 months), respectively ($p = 0.002$) (Figure 2).

Univariate Cox model analyses suggested that tumor size and number were potentially associated with treatment outcomes, and as such, these were incorporated into a multivariate model. This analysis, in turn, determined that tumor size > 10 cm ($p = 0.002$) and multiple tumors ($p = 0.022$) were independent predictors of poor prognosis, whereas PVS-$^{125}$I TACE-S was a predictor of favorable patient’s prognosis ($p = 0.040$) (Table 4).

**Discussion**

Nowadays, effective treatments for HCC with PVTT are limited and controversial. Liver resection can cure some patients with types I-III HCC [18]. However, post-operative long-term survival outcomes of these patients are poor due to high HCC recurrence rates, especially for early recurrence within 5 years of surgery [19]. Therefore, selection of the most appropriate treatment approach for these patients is critically important.

One approach to improve OS of patients with HCC/PVTT is radiofrequency ablation. However, this approach is associated with a relatively high-risk of injury.
of the portal vein. In addition, PVTT ablation can also cause bile vessel injury due to adjacent location of PVTT and bile vessels [4]. Also, transarterial radio-embolization has been used more commonly for patients with HCC and PVTT due to lower risk of hepatic ischemia and infarction, but it is not yet commercially available in certain regions, such as mainland China or Japan [20]. SBRT achieved a better therapeutic effect because the tumor could receive a higher dose of radiation directly. It was relatively safe, as the organ could also be protected from substantial radiation. However, tumor size was the main limiting factor with local control rates of 91% (< 5 cm tumors) and 74% (≥ 5 cm tumors) [21]. Some studies suggested that high-dose-rate brachytherapy (HDR-BRT), as an ablation technique using gamma irradiation of iridium-192 (192Ir) source, has successfully been used in HCC. In addition, HDR-BRT is not restricted to tumor size or tumors close to blood vessels or sensitive structures [22, 23]. However, research on whether HDR-BRT can benefit various types of PVTT is lacking.

| Table 1. Patients’ demographic and clinical characteristics |
|------------------------------------------------------------|
| **Variable** | **Group A (n = 28)** | **Group B (n = 25)** | **P-value** |
| Sex, n (%) | | | 0.857* |
| Male | 23 (82.1) | 21 (84.0) | |
| Female | 5 (17.9) | 4 (16.0) | |
| Age (years), mean ±SD | 57.3 ±12.2 | 57.5 ±8.0 | 0.956* |
| Child-Pugh score, n (%) | | | 0.925* |
| 5 | 17 (60.7) | 15 (60.0) | |
| 6 | 8 (28.6) | 8 (32.0) | |
| 7 | 3 (10.7) | 2 (8.0) | |
| ECOG performance, n (%) | | | 0.991* |
| 1 | 19 (67.9) | 17 (68.0) | |
| 2 | 9 (32.1) | 8 (32.0) | |
| Etiology, n (%) | | | 0.902* |
| HBV | 23 (82.1) | 20 (80.0) | |
| HCV | 3 (10.7) | 2 (8.0) | |
| Alcohol | 1 (3.6) | 2 (8.0) | |
| Other | 1 (3.6) | 1 (4.0) | |
| Classifications of PVTT, n (%) | | | 0.983* |
| Type II | 10 (35.7) | 9 (36.0) | |
| Type III | 18 (64.3) | 16 (64.0) | |
| AFP (mg/ml), n (%) | | | 0.958* |
| > 400 | 11 (39.3) | 10 (40.0) | |
| ≤ 400 | 17 (60.7) | 15 (60.0) | |
| Ascites, n (%) | | | 0.786* |
| Present | 8 (28.6) | 8 (32.0) | |
| Absent | 20 (71.4) | 17 (68.0) | |
| TACE times, mean ±SD | 2.8 ±0.9 | 1.8 ±0.8 | < 0.001# |
| Tumor size (cm), mean ±SD | 9.0 ±3.5 | 9.2 ±2.6 | 0.837# |
| Tumor size, n (%) | | | 0.834* |
| ≥ 10 cm | 10 (35.7) | 9 (36.0) | |
| < 10 cm | 18 (64.3) | 16 (64.0) | |
| Multiple tumors, n (%) | | | 0.991* |
| ≥ 3 | 9 (32.1) | 8 (32.0) | |
| < 3 | 19 (67.9) | 17 (68.0) | |
| Fistula, n (%) | | | 0.806* |
| Present | 4 (14.3) | 3 (12.0) | |
| Absent | 24 (85.7) | 22 (88.0) | |
| Total bilirubin (μmol/l), mean ±SD | 24.7 ±10.4 | 25.3 ±11.1 | 0.833# |

*Data obtained with Pearson χ² test; # Data obtained with independent sample t-test; ECOG – Eastern Cooperative Oncology Group; HBV – hepatitis B virus; HCV – hepatitis C virus; PVTT – portal vein tumor thrombus; AFP – α-fetoprotein; TACE – transcatheter arterial chemoembolization; value of p < 0.05 was considered statistically significant difference
In this study, PVS-125I TACE-S was employed. It was a well-tolerated and viable strategy for improving hepatic functionality and prolonging survival, compared with traditional TACE-S in patients with HCC with type III PVTT. The AEs related to PVS insertion and endovascular 125I seed-strip implantation, including fever, abdominal pain, and transient decrease in liver functionality, which were resolved after conservative managements. Decreased liver function is likely to result from injury to the bile duct upon puncture of the portal vein [24]. TACE-S-related AEs, such as nausea or vomiting, fever, and abdominal pain were relatively mild and were resolved after symptomatic treatments. Fatigue, diarrhea, hypertension, hand-foot syndrome, alopecia, pruritus, rash or desquamation, voice change, and anorexia were the most common sorafenib-related AEs, which could be managed by sorafenib dose adjustment, short-term interruption, and medications. Abscess rarely occurred and was resolved through puncture drainage. All AEs showed no significant differences between groups, which suggested that PVS-125I TACE-S was a well-tolerated intervention strategy.

In the present study, pre-operative ALBI score differences between groups were not significantly different (p = 0.724), nor did these scores differ at 1 month post-operatively (p = 0.666). However, those scores were significantly different at 3 (p = 0.007) and 6 (p = 0.044) months post-operatively. Patients with HCC and PVTT had relatively poor basal liver function due to portal vein invasion, and liver function was aggravated after repeated TACE-S. It was postulated that this might be due to multiple reasons, including liver function damage caused by chemotherapy and embolization that could cause damage to normal liver tissue, further aggravating the loss of the remaining liver function. Additionally, sorafenib itself has a potential to reduce portal blood flow, increasing the risk of liver failure [18, 25, 26]. However, in group A, PVS insertion and endovascular 125I seed-strip implantation were performed before TACE-S. PVS provided immediate restoration of the blood flow of obstructed portal vein, improving the hepatic blood supply. In addition, as 125I seed-strip was used for sustained intravascular brachytherapy, a suitable means of counteracting neointimal hyperplasia and improving stent patency duration is required [24]. The combined therapy can lead to long-term re-canualization of the portal vein, and hence, liver function is improved to varying degree.

Also, better DCR was observed in group A rather than in group B (71.4% vs. 44.0%) after 6 months of treatment. TACE was reported to block the flow through arteries, which supplied blood to the tumor, thus allowing for the control of tumor growth and PVTT progression, since the hepatic artery was the primary source of blood for tumor cells and thrombi [17]. Complete local necrosis was often difficult to achieve because this region had blood supplied by both the artery and portal vein, and it had the potential for collateral arterial development [27, 28].

### Table 2. Transcatheter arterial chemoembolization-sorafenib (TACE-S)-associated adverse events (AEs) for the two groups (%)

| Complications                  | Group A (n = 28) | Group B (n = 25) | P-value |
|-------------------------------|-----------------|-----------------|---------|
|                               | Grade 1-2 (%)   | Grade 3-5 (%)   |         |
| Nausea or vomiting            | 42.9            | 0.0             | 0.707   |
| Fever                         | 35.7            | 0.0             | 0.776   |
| Abdominal pain                | 39.3            | 0.0             | 0.728   |
| Abscess                       | 0.0             | 3.6             | 0.340   |
| Fatigue                       | 25.0            | 7.1             | 0.871   |
| Diarrhea                      | 35.7            | 7.1             | 0.958   |
| Hypertension                  | 32.1            | 3.6             | 0.997   |
| Hand-foot syndrome            | 42.9            | 14.3            | 0.878   |
| Alopecia                      | 7.1             | 0.0             | 0.906   |
| Pruritus                      | 14.3            | 0.0             | 0.580   |
| Rash or desquamation          | 7.1             | 3.6             | 0.989   |
| Voice change                  | 3.6             | 0.0             | 0.340   |
| Anorexia                      | 14.3            | 0.0             | 0.806   |

*P<0.05 was considered statistically significant difference

**Table 3.** Pre- and post-operative albumin-bilirubin (ALBI) scores in the two treatment groups

| ALBI score                  | Group A (n = 28) | Group B (n = 25) | P-value |
|-----------------------------|-----------------|-----------------|---------|
| Pre-operative               | −2.57 ±0.42     | −2.61 ±0.38     | 0.724   |
| 1-month post-operative      | −2.62 ±0.46     | −2.20 ±0.59     | 0.666   |
| 3 months post-operative     | −2.17 ±0.59     | −1.69 ±0.48     | *0.007  |
| 6 months post-operative     | −2.28 ±1.23     | −1.47 ±0.31     | *0.044  |

*Significance, ALBI – albumin-bilirubin
TACE was also associated with elevated levels of vascular endothelial growth factor, resulting in a higher risk of local recurrence [27]. Sorafenib combination treatment reduced TACE-associated risks and improved outcomes. However, some reports suggested that sorafenib was less effective in those with Vp3/4 PVTT (< 10% of response rate) [6, 9]. In contrast, 125I seed implantation in the portal vein allowed for effective PVTT control owing to sustained low-dose X- and γ-ray release throughout the tumor area, damaging tumor cell DNA and disrupting proliferation [11, 29, 30].

In a sub-group analysis, a significant difference in OS was observed in group A only for patients with type III PVTT (11.5 vs. 7.5 months). This might be because the near-total portal vein blockage in patients with type III PVTT was linked to a rapid decrease in liver function [31]. The liver function could be aggravated by repeated TACE and long-term sorafenib application, as described earlier.

This, in turn, required modifications or discontinuations of TACE-S treatment regimens, potentially constraining their therapeutic value [32, 33]. PVTT in the main portal vein can also increase portal vein pressure, causing fatal acute variceal bleeding [10]. The PVS insertion and endovascular 125I seed-strip implantation strategy improved portal vein obstruction, thereby reducing the risk of esophageal and gastric bleeding [34, 35]. In addition, 125I seed implantation improved local tumor control and delayed PVTT progression, and possibly reducing the risk of local and distant metastases [36]. No significant differences in OS were observed in patients with type II PVTT (p = 0.689); C) Patients with type III PVTT (p = 0.002)

The multivariate analyses showed that tumor size and multiple tumors were independent predictors of OS, and
this was consistent with previous findings [37]. Additionally, the treatment strategy was an independent predictor. However, no positive results were obtained for PVTT typing, probably because of PVS-125I TACE-S treatment protocol, which could be more important for prolonged survival.

This study had certain limitations. First, the sample size was small, which undermined the conclusion. Second, this study was retrospective in nature. Despite efforts to control potential confounding factors, a future randomized controlled trial is needed to validate the results. Lastly, a skin surface dosimeter was not employed, probably because of PVS-125I TACE-S treatment protocol, which could be more important for prolonged survival.

Table 4. Univariate and multivariate analyses of variables associated with overall survival (OS) in hepatocellular carcinoma (HCC) patients suffering from portal vein tumor thrombus (PVTT)

| Variables                        | Univariate analysis | P-value | Multivariate analysis | P-value |
|----------------------------------|---------------------|---------|-----------------------|---------|
|                                  | HR (95% CI)         |         | HR (95% CI)           |         |
| Gender (male vs. female)         | 1.476 (0.710-3.068%)| 0.298   | --                    | --      |
| ECOG scores (1 vs. 2)            | 1.170 (0.646-2.119%)| 0.604   | --                    | --      |
| Etiology (HBV/HCV/alcohol/other) | 0.846 (0.583-1.228%)| 0.379   | --                    | --      |
| Child-Pugh score (5/6/7)         | 1.205 (0.812-1.787%)| 0.355   | --                    | --      |
| AFP (≤ 400 vs. > 400 ng/ml)      | 0.672 (0.378-1.193%)| 0.175   | --                    | --      |
| Ascites (absent vs. present)     | 1.014 (0.562-1.832%)| 0.962   | --                    | --      |
| Tumor size (≤ 10 cm vs. > 10 cm) | 3.575 (1.836-6.962%)| < 0.001*| 2.920 (1.478-5.768%) | 0.002*  |
| Multiple tumors (< 3 vs. ≥ 3)    | 1.897 (1.028-3.500%)| 0.041*  | 2.186 (1.122-4.256%) | 0.022*  |
| Type of PVTT (II vs. III)        | 1.422 (0.783-2.583%)| 0.247   | 1.301 (0.684-2.475%) | 0.423   |
| Fistula (absent vs. present)     | 1.006 (0.449-2.252%)| 0.988   | --                    | --      |
| Treatment strategy               | 0.576 (0.332-1.002%)| 0.510   | 0.547 (0.308-0.972%) | 0.040*  |

*Significance; OS – overall survival; HCC – hepatocellular carcinoma; PVTT – portal vein tumor thrombus; ECOG – Eastern Cooperative Oncology Group; HBV – hepatitis B virus; HCV – hepatitis C virus; AFP – α-fetoprotein

Conclusions

The combination of PVS-125I TACE-S is a safe and effective therapeutic strategy for treating patients with HCC and type III PVTT compared with TACE-S alone.

Disclosure

The authors report no conflict of interest.

References

1. Park JW, Kim YJ, Kim DY et al. Sorafenib with or without concurrent transarterial chemoembolization in patients with advanced hepatocellular carcinoma: a phase III STAH trial. J Hepatol 2019; 70: 684-691.
2. Sun J, Yang L, Shi J et al. Postoperative adjuvant IMRT for patients with HCC and portal vein tumor thrombus: an open-label randomized controlled trial. Radiother Oncol 2019; 140: 20-25.
3. Sun JX, Shi J, Li N et al. Portal vein tumor thrombus is a bottleneck in the treatment of hepatocellular carcinoma. Cancer Biol Med 2016; 13: 452-458.
4. Yuan D, Gao Z, Zhao J et al. 125I seed implantation for hepatocellular carcinoma with portal vein tumor thrombus: A systematic review and meta-analysis. Brachytherapy 2019; 18: 521-529.
5. Hwa KP, Hyun CS, Hyoung KJ et al. Comparison of radioembolization and sorafenib for the treatment of hepatocellular carcinoma with portal vein tumor thrombosis: a systematic review and meta-analysis of safety and efficacy. Korean J Radiol 2019; 20: 385-398.
6. Zhu K, Chen J, Lai L et al. Hepatocellular carcinoma with portal vein tumor thrombus: treatment with transarterial chemoembolization combined with sorafenib – a retrospective controlled study. Radiology 2014; 272: 284-293.
7. Pan T, Li XS, Xie QK et al. Safety and efficacy of transarterial chemoembolization plus sorafenib for hepatocellular carcinoma with portal venous tumour thrombus. Clin Radiol 2014; 69: e553-561.
8. Lei Z, Jun-Hui S, Zhong-Heng H et al. Prognosis nomogram for hepatocellular carcinoma patients with portal vein invasion undergoing transarterial chemoembolization plus sorafenib treatment: a retrospective multicentre study. Cardiovasc Intervent Radiol 2021; 44: 63-72.
9. Chan SL, Chong CC, Chan AW et al. Management of hepatocellular carcinoma with portal vein tumor thrombosis: Review and update at 2016. World J Gastroenterol 2016; 22: 7289-7300.
10. Sun JH, Zhou T, Zhu T et al. Portal vein stenting combined with iodine-125 seeds endovascular implantation followed by transcatheter arterial chemoembolization for treatment of hepatocellular carcinoma patients with portal vein tumor thrombus. Biomed Res Int 2016; 2016: 3048261.
11. Li S, Li L, Li B et al. Safety and efficacy of endovascular implantation of a portal vein stent combined with iodine-125 seed-strips followed by transcatheter arterial chemoembolization with sorafenib for the treatment of hepatocellular carcinoma with portal vein tumor thrombosis. Br J Radiol 2020; 93: 20190279.
12. Lu J, Guo JH, Zhu HD et al. Safety and efficacy of irradiation stent placement for malignant portal vein thrombus com-
bined with hepatocellular carcinoma: a single-center experience. J Vasc Interv Radiol 2017; 28: 786-794.
13. Zhang XP, Gao YZ, Chen ZH et al. An Eastern hepatobiliary surgery hospital/portal vein tumor thrombus scoring system as an aid to decision making on hepatectomy for hepatocellular carcinoma patients with portal vein tumor thrombus: a multicenter study. Heptology 2019; 69: 2076-2090.
14. Llovet JM, Ricci S, Mazzaferro V et al. Sorafenib in advanced hepatocellular carcinoma. N Engl J Med 2008; 359: 378-390.
15. Cheng AL, Kang YK, Chen Z et al. Efficacy and safety of sorafenib in patients in the Asia-Pacific region with advanced hepatocellular carcinoma: a phase III randomised, double-blind, placebo-controlled trial. Lancet Oncol 2009; 10: 25-34.
16. Zhang W, Liu C, Tan Y et al. Albumin-bilirubin score for predicting post-transplant complications following adult-to-adult living donor liver transplantation. Ann Transplant 2018; 23: 639-646.
17. Lv WF, Liu KC, Lu D et al. Transarterial chemoembolization for hepatocellular carcinoma combined with portal vein tumor thrombosis. Cancer Manag Res 2018; 10: 4719-4726.
18. Wang JC, Xia AL, Xu Y et al. Comprehensive treatments for hepatocellular carcinoma with portal vein tumor thrombosis. J Cell Physiol 2019; 234: 1062-1070.
19. Chen ZH, Zhang XP, Zhou TF et al. Adjuvant transarterial chemoembolization improves survival outcomes in hepatocellular carcinoma with microvascular invasion: A systematic review and meta-analysis. Eur J Surg Oncol 2019; 45: 2188-2196.
20. Lu J, Zhang XP, Zhong BY et al. Management of patients with hepatocellular carcinoma and portal vein tumour thrombosis: comparing east and west. Lancet Gastroenterol Hepatol 2019; 4: 721-730.
21. Lee J, Shin IS, Yoon WS et al. Comparisons between radiofrequency ablation and stereotactic body radiotherapy for liver malignancies: Meta-analyses and a systematic review. Radiother Oncol 2020; 145: 63-70.
22. Colletti F, Schnapauff D, Poelinger A et al. Hepatocellular carcinoma: computed-tomography-guided high-dose-rate brachytherapy (CT-HDRBT) ablation of large (5-7 cm) and very large (>7 cm) tumours. Eur Radiol 2012; 22: 1101-1109.
23. Mohrike K, Ingo GS, Seidensticker M et al. Radioablation by image-guided (HDR) brachytherapy and transarterial chemoembolization in hepatocellular carcinoma: a randomized phase II trial. Cardiovasc Intervent Radiol 2019; 42: 239-249.
24. Luo JJ, Yan Z, Liu Q et al. Endovascular placement of iodine-125 seed strand and stent combined with chemoembolization for treatment of hepatocellular carcinoma with tumor thrombus in main portal vein. J Vasc Interv Radiol 2011; 22: 479-489.
25. Zhang YF, Shang H, Zeng XL et al. Postoperative adjuvant chemo (embolization) therapy for hepatocellular carcinoma with portal vein tumor thrombosis. Onco Targets Ther 2018; 11: 5407-5417.
26. Yamashita A, Umeno N, Harada S et al. Deteriorated portal flow may cause liver failure in patients with hepatocellular carcinoma being treated with sorafenib. J Gastrointest Oncol 2016; 7: E36-E40.
27. Miki I, Murata S, Uchiyama F et al. Evaluation of the relationship between hepatocellular carcinoma location and transarterial chemoembolization efficacy. World J Gastroenterol 2017; 23: 6437-6447.
28. Sun J, Shi J, Huang B et al. The degree of hepatic arterial blood supply of portal vein tumor thrombus in patients with hepatocellular carcinoma and its impact on overall survival after transarterial chemoembolization. Oncotarget 2017; 8: 79816-79824.
29. Li S, Li B, Li L et al. The efficacy of the combination of percutaneous transhepatic biliary drainage and I stranded seeds for malignant bile duct obstruction treatment. J Contemp Brachytherapy 2020; 12: 225-232.
30. Li J, Zhang L, Sun Z et al. Iodine-125 seed implantation for residual hepatocellular carcinoma or cholangiocellular carcinoma in challenging locations after transcatheter arterial chemoembolization: Initial experience and findings. J Contemp Brachytherapy 2020; 12: 233-240.
31. Su F, Chen KH, Liang ZG et al. Comparison of three-dimenisonal conformal radiotherapy and hepatic resection in hepatocellular carcinoma with portal vein tumor thrombus. Cancer Med 2018; 7: 4387-4395.
32. Yoo SH, Jang JW, Kwon JH et al. Preemptive antiviral therapy with entecavir can reduce acute deterioration of hepatic function following transarterial chemoembolization. Clin Mol Hepatol 2016; 22: 458-465.
33. Chang WT, Lu SN, Rau KM et al. Increased cumulative doses and appearance of hand-foot skin reaction prolonged progression free survival in sorafenib-treated advanced hepatocellular carcinoma patients. Radiosurg J Med Sci 2018; 34: 391-399.
34. Chuan-Xing L, Xu H, Bao-Shan H et al. Efficacy of therapy for hepatocellular carcinoma with portal vein tumor thrombus: Chemoembolization and stent combined with iodine-125 seed. Cancer Biol Ther 2011; 12: 865-871.
35. Wu YF, Wang T, Yue ZD et al. Stents combined with iodine-125 implantation to treat main portal vein tumor thrombus. World J Gastrointest Oncol 2018; 10: 496-504.
36. Sun H, Zhang M, Liu R et al. Endovascular implantation of 125I seed combined with transcatheter arterial chemoembolization for unresectable hepatocellular carcinoma. Future Oncol 2018; 14: 1165-1176.
37. Wang Q, Xia D, Bai W et al. Development of a prognostic score for recommended TACE candidates with hepatocellular carcinoma: A multicentre observational study. J Hepatol 2019; 70: 893-903.