



Comparison of liver function and safety in hepatocellular cancer patients treated with DEB-TACE and cTACE: a multi-center, retrospective cohort study

Hua Xiang^{1#}, Bin Xiong^{2#}, Haiping Li³, Chang Zhao⁴, Zishu Zhang⁵, Cong Ma⁵, Chuansheng Zheng², Chao Luo⁶, Huaiming Qiu⁶, Yuanhui Yao¹, Hongyao Hu⁷, Hui Zhao⁷, Qingyun Long⁸, Jun Zhou⁸, Changyong Chen³, Yilong Ma⁴

¹Department of Intervention, Hunan Provincial People's Hospital, Changsha 410005, China; ²Department of Interventional Radiology, Union Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan 430022, China; ³Department of Radiology, Xiangya Hospital, Central South University, Changsha 410008, China; ⁴Department of Interventional Therapy, Affiliated Tumor Hospital of Guangxi Medical University, Nanning 530021, China; ⁵Department of Radiology, The Second Xiangya Hospital of Central South University, Changsha 410011, China; ⁶Department of Radiology, Wuhan General Hospital of Guangzhou Military, Wuhan, China; ⁷Department of Radiology, Renmin Hospital of Wuhan University, Hubei General Hospital, Wuhan 430070, China; ⁸Department of Radiology, Zhongnan Hospital of Wuhan University, Wuhan 430071, China

Contributions: (I) Conception and design: H Xiang, B Xiong, Y Ma, C Chen; (II) Administrative support: H Li, Z Zhang, C Ma; (III) Provision of study materials or patients: C Zhao, H Qiu, H Hu; (IV) Collection and assembly of data: C Zheng, C Luo, Y Yao, J Zhou; (V) Data analysis and interpretation: H Zhao, Q Long; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

[#]These authors contributed equally to this work.

Correspondence to: Yilong Ma. Department of Interventional Therapy, Affiliated Tumor Hospital of Guangxi Medical University, 71 Hedi Road, Qingxiu District, Nanning 530021, China. Email: mayilong66@aliyun.com; Changyong Chen. Department of Radiology, Xiangya Hospital Central South University, 87 Xiangya Road, Changsha 410008, China. Email: rongdaolous@163.com.

Background: This study aimed to compare the live function change and adverse events (AEs) between drug-eluting beads (DEB) transarterial chemoembolization (TACE) with CalliSpheres[®] microspheres (CSM) and conventional TACE (cTACE) in treating hepatocellular carcinoma (HCC) patients.

Methods: Three hundred and thirty-five HCC patients underwent DEB-TACE with CSM (n=171, DEB-TACE group) or cTACE (n=164, cTACE group) were consecutively enrolled in this multi-center, retrospective cohort study. Liver function indexes were reviewed before treatment (W0), at 1 week (W1) and 1 month (M1) post treatment. Moreover, AEs during operation and hospitalization were retrieved as well.

Results: The changes of albumin (ALB) [-3.55 (-6.25 to -0.43) *vs.* -2.20 (-4.63-0.00), P=0.043] and total protein (TP) [-4.62 (-10.18-0.43) *vs.* -2.50 (-7.08-1.08), P=0.013] from W1 to W0 were lower in DEB-TACE group compared to cTACE group, while no difference was observed referring to the change of alanine aminotransferase (ALT) (P=0.494), aspartate aminotransferase (AST) (P=0.747), alkaline phosphatase (ALP) (P=0.895), total bilirubin (TBIL) (P=0.059), total bile acid (TBA) (P=0.491) from W1 to W0. And the changes of these seven indexes from M1 to W0 were all similar between DEB-TACE group and cTACE group (all P>0.05). Besides, the occurrence of pain during treatment (19.3% *vs.* 11.0%, P=0.034) and the occurrence of fever during hospitalization (18.1% *vs.* 9.1%, P=0.017) were both increased in DEB-TACE group compared to cTACE group.

Conclusions: DEB-TACE with CSM is non-inferior to cTACE in terms of liver function change, while DEB-TACE with CSM leads to higher incidences of pain and fever.

Keywords: CalliSpheres[®] microspheres (CSM); drug-eluting bead transarterial chemoembolization (DEB-TACE); conventional transarterial chemoembolization (cTACE); hepatocellular carcinoma (HCC); liver function; safety

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Introduction

Hepatocellular carcinoma (HCC), the most frequent primary liver cancer, is the sixth most common neoplasm and the third leading cause of cancer death in the world (1). As a dominant therapy, surgical resection is preferred and exhibits good efficacy in the treatment of HCC. However, due to the lack of typical symptoms, most patients are diagnosed with HCC at an advanced stage with distant metastases or other severe complications, making them unsuitable for surgery; thus, the best time to perform surgical therapy has often passed (2-5). Transarterial chemoembolization (TACE), a minimally invasive method, has been reported to be the standard treatment for patients with unresectable liver cancer or a poor hepatic function reserve; TACE is performed by applying mixed formulations of embolization and chemotherapy drugs into the hepatic arteries to not only cut off of the tumor blood supply but also kill tumor cells (6-9). Conventional TACE (cTACE) with emulsions of lipiodol and chemotherapy drugs (such as doxorubicin) is widely applied for treating HCC; however, owing to the poor stability of drug carriers, the chemotherapy drugs easily diffuse into the systemic blood circulation, resulting in high systemic toxicity (10). To solve this problem, TACE with drug-eluting beads (DEB-TACE) is carried out using drug-loaded microspheres as drug carriers to precisely release chemotherapy drugs, thereby maximizing tumor necrosis and minimizing adverse effects (11,12). CalliSpheres[®] microspheres (CSMs) (Jiangsu Hengrui Medicine Co., Ltd., Jiangsu Province, China), as the first invented and produced drug-eluting beads in China, are characterized by biocompatibility and nondegradability and have physical properties that allow them to localize in tumor blood vessels; these CSMs release the adsorbed chemotherapy drugs in a controlled and predictable manner in tumor blood vessels (13). Although there have been some studies revealing the good efficacy of DEB-TACE with CSMs in treating HCC, there is limited information regarding its safety in HCC patients (14). Therefore, the purpose of this study was to compare liver function changes and adverse events (AEs) in HCC patients treated with DEB-TACE with CSMs and cTACE to further prepare DEB-TACE with CSMs for clinical application.

Methods

Study design

The present study compared the comprehensive safety profile and resulting liver function between DEB-TACE with CSMs and cTACE in the treatment of HCC as part of the DECTH study (Drug-Eluting beads transarterial chemoembolization versus Conventional Transarterial chemoembolization for Hepatocellular carcinoma). The DECTH study was a multicenter, retrospective cohort study aimed at comparing the efficacy and safety of DEB-TACE treatment and cTACE treatment in Chinese HCC patients. It included eight medical centers in China, as shown in *Table S1*, and was approved by the Institutional Review Board at each participating center.

Participants

Three hundred thirty-five HCC patients who underwent DEB-TACE or cTACE treatment between 2014/9/24 and 2017/8/28 were consecutively enrolled in this multicenter, retrospective cohort study. The inclusion criteria were as follows: (I) diagnosis of primary HCC confirmed by clinical and pathological findings; (II) age of at least 18 years old; (III) underwent DEB-TACE or cTACE treatment; (IV) complete data regarding demographics, medical history, diagnosis, clinical features, pathology, treatment, measurements and assessments. The exclusion criteria were as follows: (I) diagnosis of diffuse HCC, hepatobiliary carcinoma, mixed cell carcinoma or lamellar cell carcinoma; (II) a history of liver transplantation or other malignancies; (III) loss to follow-up without any follow-up data; and (IV) switched between DEB-TACE and cTACE treatment within 6 months.

Data collection

After written informed consent was obtained from the eligible patients or their statutory guardians, patient data were extracted from electronic medical records and the Medical Records Department, which included demographic characteristics, medical history, clinical features, laboratory

indexes of routine blood tests, liver function and kidney function, tumor marker indexes, previous treatments, records of equipment and drugs used in DEB-TACE and cTACE procedures, assessment of treatment response, documentation of AEs and follow-up data regarding patient survival. The baseline patient information collected included the following: (I) demographic characteristics: age and sex; (II) medical history: drinking history, hepatitis B (HB), hepatitis C (HC) and cirrhosis; (III) clinical features: tumor location (unilobar or bilobar), tumor distribution (multifocal disease or unifocal disease), largest nodule size, portal vein invasion, hepatic vein invasion, Eastern Cooperative Oncology Group (ECOG) performance status, Child-Pugh stage and Barcelona Clinic Liver Cancer (BCLC) stage; (IV) laboratory indexes of blood routine tests, liver function and kidney function: white blood cell (WBC) count, red blood cell (RBC) count, absolute neutrophil count (ANC), and hemoglobin (Hb), platelet (PLT), albumin (ALB), total protein (TP), total bilirubin (TBIL), total bile acid (TBA), alanine aminotransferase (ALT), aspartate aminotransferase (AST), alkaline phosphatase (ALP), blood creatinine (BCr) and blood urea nitrogen (BUN) levels; (V) tumor marker indexes: alpha fetoprotein (AFP), carcinoembryonic antigen (CEA) and carbohydrate antigen 199 (CA199) levels; and (VI) previous treatments: cTACE, surgery, systematic chemotherapy, radiofrequency ablation and targeted therapy.

Grouping

Depending on the treatment options, patients who underwent DEB-TACE treatment were assigned to the DEB-TACE group (N=171), and the others, who underwent cTACE treatment, were assigned to the cTACE group (N=164). The treatment decisions were mainly made by the patients (or their guardians) according to their disease conditions (explained by the physicians), financial status, and physical functions. In addition, for patients who had a history of cTACE therapy, the previous treatment response to cTACE was also considered.

DEB-TACE procedure

In the DEB-TACE procedure, CSMs with a diameter of 100–300 or 300–500 μm were used as chemoembolization reagent carriers and embolization agents. Before initiation of the operation, the CSMs were loaded with pirarubicin (60 or 80 mg) using the following methods: first, one

bottle of CSMs was shaken gently to equally distribute the CSMs in the bottle. Then, the CSMs and normal saline were extracted using a 20 mL syringe, which was placed erectly at room temperature (RT) for 1–2 min until the CSMs had completely precipitated. Meanwhile, the chemoembolization reagent was dissolved in solution at 20 mg/mL, mixed with the CSMs using a T joint and then stored in a syringe. Then, the syringe containing the mixture of CSMs and chemoembolization reagent solution was placed at RT and shaken gently every 5 for 15 min until the CSMs were loaded with the chemoembolization reagent. Subsequently, a contrast agent at a high concentration was added to the mixture at a 1:1, 1:1.1 or 1:1.2 ratio, after which the mixture was kept still for 5 min for further application. For large HCC tumors, if the embolization point was not reached after a bottle of CSMs was emptied, another bottle of CSMs was used.

All DEB-TACE procedures were conducted in the digital subtraction angiography (DSA) room. Before the initiation of DEB-TACE, the targeted tumor was assessed by triphasic computerized tomography (CT) or magnetic resonance imaging (MRI) according to the Milan criteria: if the diameter of the targeted tumor was less than 5 cm or the multiple targeted tumors all had diameters less than 3 cm, the CSMs were loaded with 60 mg of pirarubicin per procedure; otherwise, the CSMs were loaded with 80 mg of pirarubicin (15,16). On initiation of the DEB-TACE procedure, hepatic angiography was performed to detect the tumor-supplying vessels using segment or subsegment super-selective catheterization, which was conducted as follows: (I) if an area was found with no or few vessels, the potential tumor-supplying vessel would be identified in this area; (II) the femoral artery was punctured using the Seldinger technique, and a microcatheter with a diameter ranging from 2.4 to 5 F (Merit Maestro, Merit Medical System, Inc., Utah, USA) was used for the puncture; (III) the CSMs were injected through the microcatheter by pulse injection, during which the syringe was rotated or a T joint was used to avoid deposition of the CSMs. Embolization was stopped when the flow of contrast agent stagnated. After embolization, the microcatheter was pulled out, and the wound was compressed for hemostasis and then bandaged. In addition, for patients with a large HCC tumor, DEB-TACE was performed multiple times.

cTACE procedure

All cTACE procedures were performed in the DSA room

as well. Similar to the DEB-TACE procedure, hepatic angiography was first performed to detect the tumor-supplying vessel, as described above. Second, once the tumor-supplying vessel was selected, the femoral artery was percutaneously punctured using the Seldinger technique. Third, a 2.4 to 5 F microcatheter (Merit Maestro, Merit Medical System, Inc., Utah, USA) was subsequently used for catheterization, and the chemotherapy drug solution (pirarubicin, 60 or 80 mg, 20 mg/mL), along with ethiodized poppy seed oil (EPO) (Jiangsu Hengrui Medicine Co., Ltd., Jiangsu Province, China) as the drug carrier and polyvinyl alcohol (PVA) particles (Cook Medical LLC, Bloomington, USA) as the embolization agent, was injected into the tumor-supplying vessel. Finally, embolization was stopped when stenosis of the vessel occurred. Angiography was performed again to ensure that the EPO/PVA particles were deposited and to detect incomplete embolization.

Preprocedural and postprocedural treatments

Preprocedural and postprocedural treatments were administered to all patients treated with DEB-TACE or cTACE. The preprocedural treatments were as follows: before DEB-TACE or cTACE, antiemetic treatment was performed using tropisetron (Chia Tai Tianqing Pharmaceutical Group Co., Ltd., Shandong Province, China), analgesic treatment was performed using dezocine (Yangtze River Pharmaceutical Group, Jiangsu Province, China) and anti-infection treatment was performed. The postprocedural treatments were as follows: all patients were told to lie on one side and extend the punctured leg for 6–12 h postembolization. Patients with postoperative nausea and vomiting were treated with tropisetron (IV), and analgesic treatment was performed using pethidine, dexamethasone or lidocaine.

Assessment of liver function and AEs

The influence of DEB-TACE or cTACE treatment on liver function was assessed by testing liver function indexes, including the ALT, AST, ALP, TBIL, ALB, TP and TBA levels, which were measured at baseline (W0/M0) and week 1 (W1) and month 1 (M1) after treatment. Moreover, AEs that occurred during the operation and hospitalization were used to evaluate the safety profile; AEs included pain, nausea/vomiting, increased blood pressure and fever. Pain was evaluated using a numeric rating scale (NRS); the NRS

for pain was a 10-point numeric scale, with 0 representing “no pain”; 1–3, “mild pain”; 4–6, “moderate pain”; 7–9, “severe pain”; and 10, “unbearable pain”.

Statistical analysis

Statistical analysis was performed using SPSS 22.0 statistical software (SPSS, Inc., Chicago, USA), and figures were made using GraphPad Prism 6.01 software (GraphPad Software, Inc., San Diego, USA). Count data are expressed as counts (percentage), and comparisons between two groups were performed by chi-square test. Normally distributed continuous data are presented as the mean \pm standard deviation, and comparisons between two groups were performed by *t* test. Skewed distributed continuous data are described as the median (25th–75th quartiles), and comparisons between two groups were performed by Wilcoxon rank-sum test. Univariate and multivariate logistic regression analyses were used to determine factors affecting the occurrence of AEs. A *P* value <0.05 was considered significant, and significant results are shown in boldface.

Results

Characteristics of HCC patients

There was no difference in age (*P*=0.742) or sex (*P*=0.252) between the DEB-TACE and cTACE groups. The mean age in the DEB-TACE and cTACE groups was 54.9 \pm 11.8 and 55.4 \pm 13.2 years, respectively. The median value of the largest nodule size in the DEB-TACE group was larger than that in the cTACE group (*P*=0.004), and a larger percentage of patients had a previous history of cTACE treatment (*P*=0.018) in the DEB-TACE group than in the cTACE group. However, no differences were found in other characteristics between the two groups (all *P*>0.05) (*Table 1*). These results suggest that HCC patients in the DEB-TACE group might have presented with more severe disease.

Comparison of liver function indexes between the DEB-TACE and cTACE groups at baseline, W1 posttreatment and M1 posttreatment

The median TBIL level in the DEB-TACE group was higher than that in the cTACE group (*P*=0.001), and the

Table 1 Baseline characteristics of HCC patients

Parameters	DEB-TACE group (N=171)	cTACE group (N=164)	P value
Age (years)	54.9±11.8	55.4±13.2	0.742
Gender (male/female)	145/26	146/18	0.252
History of drink (n/%)	46 (26.9)	37 (22.6)	0.358
History of HB (n/%)	109 (63.7)	109 (66.5)	0.606
History of HC (n/%)	6 (3.5)	3 (1.8)	0.342
History of cirrhosis (n/%)	72 (42.1)	86 (52.4)	0.058
Tumor location (n/%)			0.795
Unilobar	118 (69.0)	111 (67.7)	
Bilobar	53 (31.0)	53 (32.3)	
Tumor distribution (n/%)			0.458
Unifocal	113 (66.1)	102 (62.2)	
Multifocal	58 (33.9)	62 (37.8)	
Largest nodule size (cm)	7.9 (4.8–12.1)	6.5 (3.4–7.8)	0.004
Portal vein invasion (n/%)	53 (31.0)	38 (23.2)	0.108
Hepatic vein invasion (n/%)	26 (15.2)	22 (13.4)	0.640
ECOG performance status (n/%)			0.087
0	50 (29.2)	59 (36.0)	
1	92 (53.8)	87 (53.0)	
2	29 (17.0)	17 (10.4)	
3	0 (0.0)	1 (0.6)	
Child-Pugh stage (n/%)			0.743
A	136 (79.5)	133 (81.1)	
B	34 (19.9)	29 (17.7)	
C	1 (0.6)	2 (1.2)	
BCLC stage (n/%)			0.379
A	36 (21.1)	40 (24.4)	
B	73 (42.7)	73 (44.5)	
C	54 (31.6)	40 (24.4)	
D	8 (4.7)	11 (6.7)	
Blood routine			
WBC ($\times 10^9$ cell/L)	4.9 (3.9–6.5)	5.1 (3.7–6.9)	0.603
RBC ($\times 10^{12}$ cell/L)	4.2 (3.7–4.8)	4.3 (3.9–4.7)	0.461
ANC (%)	48.1 (3.0–64.5)	52.7 (3.2–63.4)	0.650
Hb (g/L)	128.0 (112.0–145.0)	132.0 (118.0–141.0)	0.544
PLT ($\times 10^9$ cell/L)	143.0 (76.0–210.0)	147.5 (93.8–221.5)	0.539

Table 1 (Continued)

Table 1 (Continued)

Parameters	DEB-TACE group (N=171)	cTACE group (N=164)	P value
Kidney function			
BCr ($\mu\text{mol/L}$)	72.5 (61.0–85.4)	73.0 (63.0–83.0)	0.696
BUN (mmol/L)	4.7 (3.9–5.8)	4.8 (3.9–6.0)	0.398
Tumor markers			
AFP ($\mu\text{g/L}$)	203.0 (8.6–1,210.0)	82.6 (5.6–1,000.0)	0.058
CEA ($\mu\text{g/L}$)	1.8 (1.0–2.9)	2.1 (1.2–3.2)	0.335
CA199 (k $\mu\text{g/L}$)	21.7 (7.9–35.2)	20.1 (8.2–34.2)	0.668
Previous treatments (n/%)			
cTACE	62 (36.3)	40 (24.4)	0.018
Surgery	22 (12.9)	31 (18.9)	0.130
Systematic chemotherapy	3 (1.8)	3 (1.8)	0.959
Radiofrequency ablation	7 (4.1)	7 (4.3)	0.936
Targeted therapy	7 (4.1)	2 (1.2)	0.104

Data were presented as mean \pm standard deviation, median (25th–75th quantiles) or count (%). Comparison between 2 groups was determined by *t* test, Wilcoxon rank sum test or Chi-square test. P value <0.05 was considered significant, and the significant results were shown in boldface. HCC, hepatocellular carcinoma; DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemo-embolization; HB, hepatitis b; HC, hepatitis c; ECOG, Eastern Cooperative Oncology Group; BCLC, Barcelona Clinic Liver Cancer; WBC, white blood cell; RBC, red blood cell; ANC, absolute neutrophil count; Hb, hemoglobin; PLT, platelet; BCr, blood creatinine; BUN, blood urea nitrogen; AFP, alpha fetoprotein; CEA, carcino-embryonic antigen; CA199, carbohydrate antigen 199.

percentage of patients with TBIL \geq 1 upper limit of normal (ULN) in the DEB-TACE group was higher than that in the cTACE group (P=0.003) (Table 2). In addition, the median TP level in the DEB-TACE group was higher than that in the cTACE group (P=0.024). Moreover, the median AST (P=0.066) and ALP (P=0.083) levels in the DEB-TACE group were higher than those in the cTACE group.

At W1 after treatment, increased TBIL levels were observed in the DEB-TACE group compared with the cTACE group (P<0.001), and more patients presented with TBIL \geq 1 ULN (P=0.004) as well as TBIL \geq 2 ULN (P=0.007) in the DEB-TACE group than in the cTACE group (Table 3). The median ALB level in the DEB-TACE group showed a trend of being higher than that in the cTACE group (P=0.099).

At M1 after treatment, the median ALT level in the DEB-TACE and cTACE groups was similar (P=0.105), while the percentage of patients with ALT \geq 1 ULN was higher in the DEB-TACE group than in the cTACE group (P=0.025) (Table 4). The median ALP value in the DEB-TACE group was higher than that the cTACE group

(P=0.005), and the percentage of patients with ALP \geq 1 ULN in the DEB-TACE group was greater than that the cTACE group (P=0.030). Additionally, the median TBIL level was increased in the DEB-TACE group compared to the cTACE group (P=0.031). These results suggest that the liver function indexes in the DEB-TACE group were worse than those in the cTACE group at baseline, W1 posttreatment and M1 posttreatment.

Changes in liver function indexes from W0 to W1 posttreatment and M0 to M1 posttreatment

Considering that the disease condition was more serious in the DEB-TACE group than in the cTACE group, the direct comparison of liver function indexes between the two groups might not accurately reflect the advantages of the treatments. Therefore, we further compared the changes in liver function indexes between the two groups, and the results show less change in the ALB (P<0.05) and TP (P<0.05) levels from W1 to W0 in the DEB-TACE group than in the cTACE group, while there were no differences

Table 2 Baseline laboratory testing of liver function

Parameters	DEB-TACE group (N=171)	cTACE group (N=164)	P value
ALT (μ /L)	37.0 (23.2–59.0)	35.0 (22.0–52.2)	0.453
≥1 ULN (n/%)	72/170 (42.4)	65/162 (40.1)	0.680
≥2 ULN (n/%)	21/170 (12.4)	18/162 (11.1)	0.725
≥3 ULN (n/%)	7/170 (4.1)	9/162 (5.6)	0.541
AST (μ /L)	52.1 (35.0–79.4)	44.3 (30.3–75.0)	0.066
≥1 ULN (n/%)	112/170 (65.9)	97/162 (59.9)	0.257
≥2 ULN (n/%)	42/170 (24.7)	36/162 (22.2)	0.594
≥3 ULN (n/%)	23/170 (13.5)	16/162 (9.9)	0.301
ALP (μ /L)	133.0 (87.5–179.0)	112.0 (81.0–163.0)	0.083
≥1 ULN (n/%)	67/133 (50.4)	56/137 (40.9)	0.117
≥2 ULN (n/%)	18/133 (13.5)	9/137 (6.6)	0.057
≥3 ULN (n/%)	4/133 (3.0)	2/137 (1.5)	0.653
TBIL (μ mol/L)	18.1 (12.7–24.6)	14.9 (10.9–20.4)	0.001
≥1 ULN (n/%)	78/170 (45.9)	48/161 (29.8)	0.003
≥2 ULN (n/%)	14/170 (8.2)	10/161 (6.2)	0.478
≥3 ULN (n/%)	4/170 (2.4)	3/161 (1.9)	0.757
ALB (g/L)	36.8 (36.4–40.1)	36.5 (32.9–39.6)	0.917
≥1 ULN (n/%)	0/170 (0.0)	2/161 (1.2)	0.145
≥2 ULN (n/%)	0/170 (0.0)	1/161 (0.6)	0.303
≥3 ULN (n/%)	0/170 (0.0)	1/161 (0.6)	0.486
TP (g/L)	67.4 (63.6–71.3)	65.4 (61.2–70.1)	0.024
≥1 ULN (n/%)	10/170 (5.9)	4/159 (2.5)	0.131
≥2 ULN (n/%)	0/170 (0.0)	1/159 (0.6)	0.300
≥3 ULN (n/%)	0/170 (0.0)	0/159 (0.0)	–
TBA (l/L)	10.5 (5.1–24.6)	9.3 (4.4–20.3)	0.235
≥1 ULN (n/%)	84/167 (50.3)	70/152 (46.1)	0.448
≥2 ULN (n/%)	50/167 (29.9)	38/152 (25.0)	0.324
≥3 ULN (n/%)	32/167 (19.2)	24/152 (15.8)	0.429

Data were presented as median (25th–75th quantiles) or count (%). Comparison between 2 groups was determined by Wilcoxon rank sum test or Chi-square test. P value <0.05 was considered significant, and the significant results were shown in boldface. DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemo-embolization; ALT, alanine aminotransferase; ULN, upper limit of normal; AST, aspartate aminotransferase; ALP, alkaline phosphatase; TBIL, total bilirubin; ALB, albumin; TP, total protein; TBA, total bile acid.

Table 3 Liver function testing at W1 post-treatment

Parameters	DEB-TACE group (N=171)	cTACE group (N=164)	P value
ALT (μ /L)	73.7 (39.4–131.0)	60.0 (32.8–128.8)	0.353
≥1 ULN (n/%)	117/158 (74.1)	102/144 (70.8)	0.532
≥2 ULN (n/%)	70/158 (44.3)	61/144 (42.4)	0.734
≥3 ULN (n/%)	47/158 (29.7)	40/144 (27.8)	0.706
AST (μ /L)	82.0 (44.4–191.0)	81.3 (41.6–155.5)	0.392
≥1 ULN (n/%)	129/158 (81.6)	110/144 (76.4)	0.261
≥2 ULN (n/%)	80/158 (50.6)	74/144 (51.4)	0.896
≥3 ULN (n/%)	63/158 (39.9)	55/144 (38.2)	0.765
ALP (μ /L)	131.0 (95.0–189.0)	120.0 (86.0–175.0)	0.218
≥1 ULN (n/%)	58/115 (50.4)	48/113 (42.5)	0.228
≥2 ULN (n/%)	18/115 (15.7)	14/113 (12.4)	0.478
≥3 ULN (n/%)	9/115 (7.8)	4/113 (3.5)	0.163
TBIL (μ mol/L)	27.3 (18.5–40.7)	21.5 (15.4–30.8)	<0.001
≥1 ULN (n/%)	116/158 (73.4)	83/144 (57.6)	0.004
≥2 ULN (n/%)	46/158 (29.1)	23/144 (16.0)	0.007
≥3 ULN (n/%)	19/158 (12.0)	9/144 (6.3)	0.084
ALB (g/L)	32.8 (29.7–37.0)	34.2 (30.3–37.4)	0.099
≥1 ULN (n/%)	2/158 (1.3)	2/144 (1.4)	0.926
≥2 ULN (n/%)	2/158 (1.3)	2/144 (1.4)	0.926
≥3 ULN (n/%)	2/158 (1.3)	1/144 (0.7)	0.617
TP (g/L)	61.9 (57.9–66.8)	62.4 (58.6–66.3)	0.786
≥1 ULN (n/%)	0/157 (0.0)	1/140 (0.7)	0.289
≥2 ULN (n/%)	0/157 (0.0)	1/140 (0.7)	0.289
≥3 ULN (n/%)	0/157 (0.0)	1/140 (0.7)	0.289
TBA (l/L)	11.8 (5.2–20.4)	9.8 (5.2–21.0)	0.525
≥1 ULN (n/%)	82/142 (57.7)	61/128 (47.7)	0.097
≥2 ULN (n/%)	38/142 (26.8)	33/128 (25.8)	0.855
≥3 ULN (n/%)	25/142 (17.6)	19/128 (14.8)	0.540

Data were presented as median (25th–75th quantiles) or count (%). Comparison between 2 groups was determined by Wilcoxon rank sum test or Chi-square test. P value <0.05 was considered significant, and the significant results were shown in boldface. W1, 1 week; DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemo-embolization; ALT, alanine aminotransferase; ULN, upper limit of normal; AST, aspartate aminotransferase; ALP, alkaline phosphatase; TBIL, total bilirubin; ALB, albumin; TP, total protein; TBA, total bile acid.

Table 4 Liver function testing at M1 post-treatment

Parameters	DEB-TACE group (N=171)	cTACE group (N=164)	P value
ALT (μ /L)	37.1 (21.7–57.0)	31.0 (21.1–50.8)	0.105
≥1 ULN (n/%)	60/129 (46.5)	42/128 (32.8)	0.025
≥2 ULN (n/%)	19/129 (14.7)	11/128 (8.6)	0.126
≥3 ULN (n/%)	6/129 (4.7)	4/128 (3.1)	0.527
AST (μ /L)	53.5 (33.5–81.4)	44.6 (30.9–70.3)	0.110
≥1 ULN (n/%)	84/128 (65.6)	73/128 (57.0)	0.158
≥2 ULN (n/%)	33/128 (25.8)	23/128 (18.0)	0.131
≥3 ULN (n/%)	16/128 (12.5)	13/128 (10.2)	0.554
ALP (μ /L)	144.0 (108.1–193.0)	120.0 (84.0–160.0)	0.005
≥1 ULN (n/%)	59/101 (58.4)	52/119 (43.7)	0.030
≥2 ULN (n/%)	13/101 (12.9)	12/119 (10.1)	0.516
≥3 ULN (n/%)	3/101 (3.0)	1/119 (0.8)	0.502
TBIL (μ mol/L)	17.4 (13.5–24.1)	15.2 (11.0–22.2)	0.031
≥1 ULN (n/%)	56/129 (43.4)	45/129 (34.9)	0.161
≥2 ULN (n/%)	13/129 (10.1)	8/129 (6.2)	0.255
≥3 ULN (n/%)	6/129 (4.7)	6/129 (4.7)	1.000
ALB (g/L)	35.1 (30.0–38.6)	35.8 (31.4–39.6)	0.287
≥1 ULN (n/%)	1/129 (0.8)	0/129 (0.0)	0.316
≥2 ULN (n/%)	0/129 (0.0)	0/129 (0.0)	–
≥3 ULN (n/%)	0/129 (0.0)	0/129 (0.0)	–
TP (g/L)	69.1 (63.4–73.5)	67.4 (63.1–73.0)	0.591
≥1 ULN (n/%)	6/129 (4.7)	8/128 (6.3)	0.572
≥2 ULN (n/%)	0/129 (0.0)	2/128 (1.6)	0.154
≥3 ULN (n/%)	0/129 (0.0)	1/128 (0.8)	0.498
TBA (l/L)	10.7 (5.8–27.0)	9.5 (5.2–22.6)	0.474
≥1 ULN (n/%)	66/125 (52.8)	59/122 (48.4)	0.485
≥2 ULN (n/%)	41/125 (32.8)	35/122 (28.7)	0.484
≥3 ULN (n/%)	27/125 (21.6)	26/122 (21.3)	0.956

Data were presented as median (25th–75th quantiles) or count (%). Comparison between 2 groups was determined by Wilcoxon rank sum test or Chi-square test. P value <0.05 was considered significant, and the significant results were shown in boldface. “–” indicated that the data were unable to compare due to lack of events. M1, 1 month; DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemo-embolization; ALT, alanine aminotransferase; ULN, upper limit of normal; AST, aspartate aminotransferase; ALP, alkaline phosphatase; TBIL, total bilirubin; ALB, albumin; TP, total protein; TBA, total bile acid.

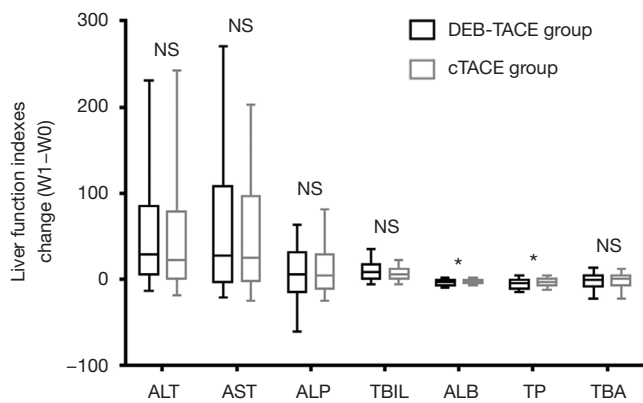


Figure 1 Changes of liver function indexes from W1 to M0. *, P<0.05. W1, 1 week; W0, before treatment; TACE, transarterial chemoembolization; cTACE, conventional cTACE; NS, no significance; ALT, alanine aminotransferase; AST, aspartate aminotransferase; ALP, alkaline phosphatase; TBIL, total bilirubin; ALB, albumin; TP, total protein; TBA, total bile acid.

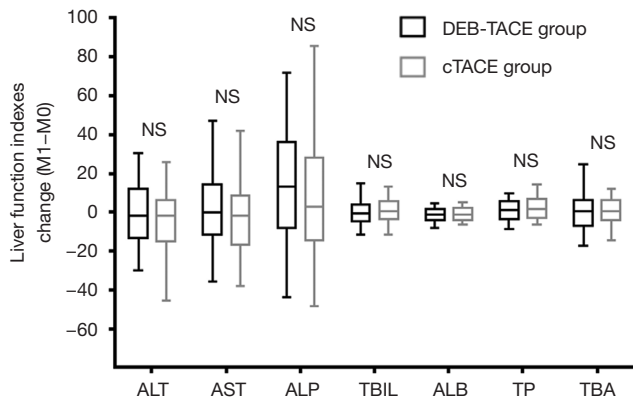


Figure 2 Changes of liver function indexes from M1 to M0. M1, 1 month; M0, before treatment; TACE, transarterial chemoembolization; cTACE, conventional cTACE; NS, no significance; ALT, alanine aminotransferase; AST, aspartate aminotransferase; ALP, alkaline phosphatase; TBIL, total bilirubin; ALB, albumin; TP, total protein; TBA, total bile acid.

in the change (W1-W0) in the ALT, AST, ALP, TBIL and TBA levels between the two groups (all P>0.05) (Figure 1).

Additionally, no differences were discovered in the change in liver function indexes (including ALT, AST, ALP, TBIL, ALB, TP and TBA) from M0 to M1 after treatment between the DEB-TACE and cTACE groups (all P>0.05) (Figure 2). These results indicate that DEB-TACE was not inferior to cTACE in terms of liver function protection.

Table 5 Adverse events occurred during treatment and hospitalization

Parameters	DEB-TACE group (N=171)	cTACE group (N=164)	P value
During treatment			
Pain (n/%)	33 (19.3)	18 (11.0)	0.034
Pain grade (NRS) (n/%)			0.040
Mild pain	23 (69.7)	17 (94.4)	
Moderate pain	8 (24.2)	1 (5.6)	
Severe pain	2 (6.1)	0 (0.0)	
Nausea/vomiting (n/%)	14 (8.2)	12 (7.3)	0.766
Rise in blood pressure (n/%)	4 (2.3)	1 (0.6)	0.192
During hospitalization			
Pain (n/%)	40 (23.4)	29 (17.7)	0.196
Pain grade (NRS) (n/%)			0.449
Mild pain	33 (82.5)	26 (89.7)	
Moderate pain	7 (17.5)	2 (6.9)	
Severe pain	0 (0.0)	1 (3.4)	
Fever (n/%)	31 (18.1)	15 (9.1)	0.017
Nausea/vomiting (n/%)	17 (9.9)	12 (7.3)	0.393

Data were presented as count (%). Comparison between 2 groups was determined by Chi-square test or Wilcoxon rank sum test. P value <0.05 was considered significant, and the significant results were shown in boldface. DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemo-embolization; NRS, numeric rating scale.

Comparison of AEs between the DEB-TACE and cTACE groups during treatment and hospitalization

During treatment, a greater proportion of patients presented with pain (P=0.034) and a high grade of pain (P=0.040) in the DEB-TACE group than in the cTACE group (Table 5). During hospitalization, a greater proportion of patients developed fever in the DEB-TACE group than in the cTACE group (P=0.017).

Logistic regression was used to analyze factors affecting AE occurrence (Table 6). Univariate logistic regression showed that a history of drinking (P=0.038), a largest nodule size ≥7 cm (P<0.001), portal vein invasion (P<0.001), hepatic vein invasion (P<0.001), a higher BCLC stage (P<0.001), previous cTACE treatment (P<0.001), previous surgery (P=0.044) and ALP ≥1 ULN (P=0.040) were

Table 6 Factors affecting adverse events occurrence by logistic regression model analysis

Parameters	Univariate logistic regression				Multivariate logistic regression			
	P value	OR	95% CI		P value	OR	95% CI	
			Lower	Higher			Lower	Higher
DEB-TACE vs. cTACE	0.507	1.167	0.739	1.843	0.012	3.740	1.344	10.403
Age \geq 60 years	0.371	0.804	0.499	1.297	0.570	0.744	0.269	2.063
Male	0.877	0.948	0.480	1.871	0.919	1.092	0.201	5.921
History of drink	0.038	0.581	0.348	0.970	0.513	0.701	0.241	2.035
History of HB	0.887	0.966	0.599	1.557	0.826	1.129	0.384	3.316
History of HC	0.497	0.577	0.118	2.823	1.000	–	0.000	–
History of cirrhosis	0.067	1.541	0.970	2.446	0.416	0.637	0.215	1.890
Multifocal disease	0.195	1.377	0.849	2.231	0.525	0.660	0.183	2.376
Tumor location-Bilobar	0.175	1.385	0.865	2.216	0.018	6.469	1.385	30.219
Largest nodule size \geq 7 cm	<0.001	3.312	2.036	5.387	0.066	2.896	0.932	8.996
Portal vein invasion	<0.001	0.326	0.197	0.538	0.916	0.911	0.161	5.154
Hepatic vein invasion	<0.001	0.187	0.097	0.359	0.899	1.122	0.190	6.617
Higher ECOG performance status	0.224	0.751	0.474	1.191	0.750	0.836	0.279	2.509
Higher Child-Pugh stage	0.147	1.467	0.874	2.461	0.225	0.412	0.098	1.727
Higher BCLC stage	<0.001	2.285	1.690	3.088	0.434	2.368	0.273	20.565
Previous cTACE treatment	<0.001	3.635	2.006	6.586	<0.001	24.622	5.576	108.719
Previous surgery	0.044	2.066	1.018	4.191	0.678	1.371	0.308	6.103
Previous systematic chemotherapy	0.999	–	0.000	–	0.999	0.000	0.000	–
Previous radiofrequency ablation	0.171	0.472	0.161	1.382	0.281	4.220	0.308	57.803
Previous targeted therapy	0.457	0.602	0.158	2.289	0.999	–	0.000	–
ALT \geq 1 ULN	0.276	1.293	0.815	2.053	0.876	1.103	0.322	3.774
AST \geq 1 ULN	0.062	1.591	0.977	2.592	0.621	1.407	0.364	5.441
ALP \geq 1 ULN	0.040	1.675	1.024	2.740	0.993	1.005	0.310	3.256
TBIL \geq 1 ULN	0.142	1.419	0.890	2.264	0.522	1.581	0.390	6.415
ALB \geq 1 ULN	0.999	0.000	0.000	–	0.506	0.661	0.195	2.240
TP \geq 1 ULN	0.712	0.800	0.245	2.611	0.850	0.891	0.270	2.943
TBA \geq 1 ULN	0.598	0.881	0.551	1.409	0.309	0.543	0.167	1.761
WBC abnormal	0.959	1.013	0.614	1.671	0.284	1.769	0.623	5.025
RBC abnormal	0.584	0.879	0.555	1.393	0.416	0.649	0.228	1.842
ANC abnormal	0.844	1.058	0.602	1.862	0.522	0.671	0.198	2.278
Hb abnormal	0.163	1.393	0.874	2.219	0.620	1.357	0.405	4.548

Table 6 (Continued)

Table 6 (Continued)

Parameters	Univariate logistic regression				Multivariate logistic regression			
	P value	OR	95% CI		P value	OR	95% CI	
			Lower	Higher			Lower	Higher
PLT abnormal	0.051	0.620	0.384	1.002	0.975	1.018	0.333	3.110
BCr abnormal	0.423	1.469	0.573	3.767	0.866	1.158	0.211	6.362
BUN abnormal	0.119	0.602	0.318	1.138	0.692	0.766	0.205	2.862
AFP abnormal	0.052	1.697	0.994	2.896	0.990	1.007	0.361	2.811
CEA abnormal	0.486	0.745	0.325	1.706	0.134	3.953	0.655	23.852
CA199 abnormal	0.253	0.693	0.370	1.299	0.313	0.546	0.169	1.766

Data were presented as P value, OR and 95% CI. Factors affecting adverse events occurrence were determined by univariate and multivariate logistic regression analysis. P value <0.05 was considered significant, and the significant results were shown in boldface. “–” indicated that the value was unable to be calculated due to lack of events. Child-pugh Stage was scored as 0-A, 1-B, 2-C; BCLC stage was scored as 1-Stage A, 2-Stage B, 3-Stage C, 4-Stage D, the logistic analysis was performed based on these definitions. OR, odds ratio; CI, confidence interval; DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemoembolization; HB, hepatitis b; HC, hepatitis c; ECOG, Eastern Cooperative Oncology Group; BCLC, Barcelona Clinic Liver Cancer; ULN, upper limit of normal; ALT, alanine aminotransferase; AST, aspartate aminotransferase; ALP, alkaline phosphatase; TBIL, total bilirubin; ALB, albumin; TP, total protein; TBA, total bile acid; WBC, white blood cell; RBC, red blood cell; ANC, absolute neutrophil count; Hb, hemoglobin; PLT, platelet; BCr, blood creatinine; BUN, blood urea nitrogen; AFP, alpha fetoprotein; CEA, carcino-embryonic antigen; CA199, carbohydrate antigen 199.

correlated with the increased occurrence of AEs. Further multivariate logistic regression revealed that DEB-TACE (P=0.012), a bilobar tumor location (P=0.018) and previous cTACE treatment (P<0.001) were independent risk factors predicting the occurrence of AEs in HCC patients.

Discussion

This study revealed the following: (I) liver function was worse in the DEB-TACE group than in the cTACE group at baseline (W0), W1 and M1; (II) less change (W1–W0) in the ALB and TP levels was observed in the DEB-TACE group than in the cTACE group; (III) DEB-TACE was correlated with an increased occurrence and severity of pain and fever.

The liver is an important metabolic organ in the human body. HCC patients who are not suitable for surgical resection are generally recommended for TACE treatment. Nevertheless, TACE is an interventional therapy that, to some extent, causes liver damage as well as complications (17,18). Although the history of the clinical application of CSMs is relatively short, many studies have shown that DEB-TACE with CSMs is effective for the treatment of HCC and tolerated by Chinese patients, even HCC patients

who have previously undergone cTACE treatment (14). However, the effect of DEB-TACE with CSMs on liver function in HCC patients is still unclear. In a recent study, Wu *et al.* compared the effects of DEB-TACE with CSMs and cTACE on liver function in HCC patients, and the results showed that while liver function indexes were elevated in both groups at one month after the intervention, the ALT, AST and TBIL levels in the DEB-TACE group were greater than those in the cTACE group (19). A study by Liu *et al.* indicated not only that the ALB, TP, TBIL, and AST levels in patients treated with DEB-TACE with CSMs decreased one week after treatment and recovered after one month but also that the ALT, AST and TBIL levels in were lower after treatment with DEB-TACE with CSMs than cTACE. The above studies indicate that DEB-TACE with CSMs is better than cTACE in terms of liver function protection. However, contrary to previous studies, in the current study, we compared the effects of DEB-TACE with CSMs and cTACE on liver function, and the results show that the levels of liver function indexes were higher after DEB-TACE with CSMs than after cTACE at baseline, W1 and M1. The possible reasons are as follows: the largest nodule size was found to be larger in the DEB-TACE group, which means that the disease severity might

have been worse in the DEB-TACE group. Additionally, TACE is an interventional treatment that causes damage to the liver, and a higher percentage of patients with a history of previous cTACE treatment was found in the DEB-TACE group, which suggests that patients in the DEB-TACE group had poorer liver function than those in the cTACE group at baseline. Thus, liver function remained poorer in the DEB-TACE group than in the cTACE group even at W1 and M1 after treatment (17,18).

Owing to the difference at baseline between the two groups, i.e., the poorer liver function in the group of patients treated with DEB-TACE with CSMs at baseline, it was not suitable to directly compare the liver function indexes between the two groups at W1 or M1 after treatment. Therefore, we further compared the change in liver function indexes between the two groups from baseline to W1 posttreatment and M1 posttreatment, and we observed greater reductions in the ALB and TP levels at W1 in the DEB-TACE group than in the cTACE group and no difference in the change in liver function indexes between the two groups at M1 after treatment. These findings are similar to the trends of previous research results. These results indicate that DEB-TACE with CSMs was at least as effective in treating HCC patients as cTACE. This might be explained by the following aspects: (I) CSMs have physical properties that allow them to localize within tumor blood vessels and release the adsorbed chemotherapy drugs in a controlled and predictable manner, thus reducing drug toxicity at nontargeted sites and reducing liver damage (13); (II) the characteristics of CSMs result in good biocompatibility and non-biodegradability; thus, CSMs can be retained in tumor blood vessels for a long time to block the tumor blood supply and release drugs, thereby preventing secondary liver damage caused by multiple TACE procedures (20).

Although the good therapeutic efficacy of TACE has been reported, some AEs can still occur; among these, the most common complication after TACE is post-embolization syndrome (PES), with an incidence rate of 60–80% and several clinical symptoms, including fever, abdominal pain, nausea and vomiting (21). Regarding the comparison of AEs between DEB-TACE and cTACE, contradictory results have been reported (19,22–25). Most studies have found no difference in AEs between DEB-TACE and cTACE in the treatment of HCC patients (22–24); only one study has shown that AEs were less frequent in DEB-TACE than cTACE (19). In addition, a randomized controlled trial revealed that the incidence and severity of

postprocedural fever, fatigue, nausea/vomiting, hematoma, cholecystitis, spleen infarction, infection, and liver abscess did not differ between DEB-TACE and cTACE, while postprocedural pain was less frequent in HCC patients treated with DEB-TACE (25). These studies indicate that DEB-TACE with CSMs is safe for HCC patients because it does not cause serious adverse reactions. Partially consistent with the results of previous studies, in this study, the AEs that occurred in both groups were mild, including pain, nausea/vomiting, increased blood pressure and fever. However, compared with cTACE, a greater percentage of patients experienced pain and a higher pain grade during treatment with DEB-TACE with CSMs, and more patients treated with DEB-TACE with CSMs developed fever during hospitalization. The possible explanations are as follows: (I) the median value of the largest nodule size in the CSM-TACE group was greater than that in the cTACE group; thus, larger microbeads were required during the operation, and the larger microbeads cause much more pressure on nerves and thus more pain; (II) CSMs are made from a nondegradable polymer based on calibrated microspheres and produce permanent vascular embolization, which might cause prolonged blood flow stasis and thrombosis of the proximal artery, thereby resulting in more severe pain in HCC patients; (III) as the response rate was better in the CSM-TACE group than the cTACE group, there was also more tumor tissue necrosis, which may result in an increased likelihood of fever in patients. In addition, a recent study has reported that a novel application of DEB-TACE with beads of different sizes instead of only one size results in few AEs during and after the operation; this approach could be further applied in our clinical practice to reduce AEs and improve patients' outcomes (26).

The present study still existed several limitations. Firstly, the sample size of this study was relatively small, which might result in lower statistical performance. Secondly, it was not a randomized controlled study. Thirdly, the efficacy of long-term treatment with DEB-TACE and cTACE on liver function was not investigated, further study is needed.

In conclusion, DEB-TACE with CSM is non-inferior to cTACE in terms of liver function change, while DEB-TACE with CSM leads to higher incidences of pain and fever.

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Footnote

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at <http://dx.doi.org/10.21037/tcr.2019.09.15>). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. This study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by the Institutional Review Board at each participating center (KY2018012), and written informed consent was obtained from the eligible patients or their statutory guardians.

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Supplementary

Table S1 Number of patients included in this study by medical center

Medical center	DEB-TACE group	cTACE group	Total Patients
Guangxi District Cancer Hospital (n)	42	47	89
Hunan People's Hospital (n)	36	37	73
Xiangya Hospital (n)	38	21	59
Xiangya Affiliated Second Hospital (n)	20	29	49
Hubei Provincial People's Hospital (n)	13	14	27
Wuhan Union Hospital (n)	15	5	20
General Hospital of Hubei Army (n)	3	8	11
Wuhan Zhongnan Hospital (n)	4	3	7
Total (N)	171	164	335

DEB-TACE, drug-eluting bead transarterial chemoembolization; cTACE, conventional transarterial chemo-embolization.