**Name of Journal:** *World Journal of Clinical Cases*

**Manuscript NO:** 83529

**Manuscript Type:** CASE REPORT

**Hypothetical hypoxia-driven rapid disease progression in hepatocellular carcinoma post transarterial chemoembolization: A case report**

Yeo *et al*. Rapid disease progression in hepatocellular carcinoma

Kai-Fuan Yeo, Amy Ker, Pei-En Kao, Chi-Chih Wang

**Kai-Fuan Yeo, Chi-Chih Wang,** Division of Gastroenterology and Hepatology, Department of Internal Medicine, Chung Shan Medical University Hospital, Taichung 402306, Taiwan

**Amy Ker, Pei-En Kao, Chi-Chih Wang,** School of Medicine, Chung Shan Medical University, Taichung 402306, Taiwan

**Author contributions:** YeoKF, Ker A, and Kao PE reviewed the literature and contributed to manuscript drafting; Wang CC were responsible for the revision and final approval of the manuscript; This article supported by Yeo KF, Yeo KF, and Ker A contributed equally to the manuscript as the first authors.

**Corresponding author: Chi-Chih Wang, PhD, Associate Professor, Director,** Division of Endoscopy, Department of Internal Medicine, Chung Shan Medical University Hospital, No. 110 Sec. 1, Jianguo N.Rd., Taichung 402306, Taiwan. bananaudwang@gmail.com

**Received:** February 18, 2023

**Revised:** May 25, 2023

**Accepted:** June 6, 2023

**Published online:** July 6, 2023

**Abstract**

BACKGROUND

Transarterial chemoembolization (TACE) is widely performed for intermediate-stage or unresectable hepatocellular carcinoma (HCC), but approximately half of patients do not respond to TACE treatment. We describe a case of rapidly progressing of HCC after TACE and provide a possible hypothesis for this condition. The finding may contribute to identifying patients who obtain less benefit from TACE, thus avoiding the unnecessary waste of medical resources and treatment during the golden hour window.

CASE SUMMARY

A 61-year-old woman had been diagnosed with chronic hepatitis B infection and HCC at Barcelona Clinic Liver Cancer stage B, which had been treated by segmental hepatectomy 14 mo ago. The tumor recurred in the two months after surgery. She received an initial TACE and then underwent systemic therapy with lenvatinib 8 mg daily due to an increased level of alpha-fetoprotein (AFP) after the first TACE. However, the tumor continued to progress with an increased level of AFP, and she underwent a second TACE, after which the tumor volume did not obviously decrease on the contrast-enhanced computed tomography image. One month later, she had a third TACE to control the residual HCC tumors. Two weeks after that, the HCC had increased dramatically with tea-colored urine and yellowish skin turgor. Eventually, the patient refused further treatment and went into hospice care.

CONCLUSION

Intense hypoxia induced by TACE can trigger rapid disease progression in infiltrative HCC patients with a large tumor burden

**Key Words:** Carcinoma; hepatocellular; Transarterial chemoembolization; Tumor hypoxia; Disease progression; Tumor burden; Case report

**©The** **Author(s) 2023.** Published by Baishideng Publishing Group Inc. All rights reserved.

**Citation**: Yeo KF, Ker A, Kao PE, Wang CC. Hypothetical hypoxia-driven rapid disease progression in hepatocellular carcinoma post transarterial chemoembolization: A case report. *World J Clin Cases* 2023; 11(19): 4664-4669

**URL**: https://www.wjgnet.com/2307-8960/full/v11/i19/4664.htm

**DOI**: https://dx.doi.org/10.12998/wjcc.v11.i19.4664

**Core Tip:** We report an hepatocellular carcinoma (HCC) case with a large tumor burden and infiltrative tumor pattern who exhibited rapidly increased tumor volume within two weeks after undergoing a third trans-arterial chemoembolization (TACE). Although the Barcelona Clinic Liver Cancer staging system classifies multinodular HCC without portal invasion or extrahepatic spread in stage B, it appears that TACE is not suitable for those with a large tumor burden or infiltrative tumor pattern. In addition, hypoxia is an important factor for tumor development, metastasis, and drug resistance. Our case suggests that intense hypoxia induced by TACE may lead to the rapid progression of HCC.

**INTRODUCTION**

Primary liver cancer was the third-leading cause of cancer death globally in 2020, with an age-standardized incidence rate of 19.3 cases per 100000 people and an age-standardized mortality rate of 17.7 per 100000 people[1]. Hepatocellular carcinoma (HCC) is the most common type of liver cancer, accounting for approximately 75%–85% of cases. Clinically, trans-arterial chemoembolization (TACE) is used as the first-line treatment for patients with Barcelona Clinic Liver Cancer (BCLC) stage B and some unresectable HCC[2,3]. TACE injects cytotoxic agents into the arteries, followed by the embolization of tumor blood vessels, which induces strong cytotoxic and ischemic effects to destroy tumor cells. However, the response rate of intermediate-stage HCC patients receiving TACE is low for unclear reasons, with a pooled objective response rate of about 52%[4]. To our best knowledge, only a few studies have investigated the baseline characteristics of those not benefitting from TACE, such as large tumor volume, high tumor number, and poor performance status[5,6]. In our case, an intermediate-stage HCC patient underwent a third TACE and experienced rapid disease progression within two weeks. This case may provide information on the failure of and resistance to TACE in HCC treatment.

**CASE PRESENTATION**

***Chief complaints***

Progressive yellowish skin turgor and firm sensation over the epigastric area.

***History of present illness***

A 61-year-old woman had been diagnosed with HCC at BCLC stage B as well as chronic hepatitis B infection 14 mo earlier. Because the typical image of contrast-enhanced abdominal computed tomography (CT) showed three heterogeneous arterial enhancing lesions with delayed phase wash out in segments S4 and S8, with the largest measuring 7.3 cm, she received segmental hepatectomy and cholecystectomy on October 5, 2021 based on the extensive criteria of the University of California San Francisco[7]. Tumor recurrence emerged two months later, and we performed a TACE followed by systemic therapy of lenvatinib 8 mg daily due to an elevated alpha-fetoprotein (AFP) level after the TACE. The subsequent CT confirmed the progression of HCC volume, so we arranged a second course of TACE after the lenvatinib treatment.

***History of past illness***

The patient had a history of major depression with good medication control.

***Personal and family history***

The patient denied smoking, alcohol use, and betel nut use. She also denied any family history of malignancy. There was no family history of hepatitis B or hepatitis C.

***Physical examination***

The patient’s height and weight were 162 cm and 60 kg. Her vital signs were stable, with a body temperature of 37.5°C, pulse rate of 90 bpm, respiration rate of 17 breaths per minute, and blood pressure of 136/76 mmHg. A physical examination revealed icteric sclera on the day of admission. A palpable firm mass lesion was observed at the epigastric area.

***Laboratory examinations***

The laboratory data revealed mild anemia, including a low red blood cell count of 392 × 106 cells/μL and a high mean corpuscular hemoglobin of 32.7 pg. The hemoglobin, hematocrit, mean corpuscular volume, and mean corpuscular hemoglobin concentrations were within normal values. Coagulation tests showed a low platelet count of 126000 platelets/μL and a prothrombin time/international normalized ratio within the normal value. Alanine aminotransferase and aspartate aminotransferase were elevated at 273 U/L and 272 U/L, respectively, while total bilirubin was 4.7 mg/dL. The renal function tests and electrolyte tests were within normal ranges.

***Imaging examinations***

An abdominal contrast CT showed increased residual HCC volume (Figure 1A) after the second TACE course. The coronal view showed a patent portal vein and normal bile ducts (Figure 1B). Contrast-enhanced magnetic resonance imaging (MRI) was arranged taken two weeks after a third TACE course because of an episode of jaundice. It revealed rapidly progressing of HCC volume in a short time, with left intra-hepatic duct (IHD) and portal vein tumor invasions (Figure 2).

**FINAL DIAGNOSIS**

HCC with rapid, extensive progression and left IHD invasion after TACE.

**TREATMENT**

The abdominal enhanced CT still showed increased residual HCC volume and a high AFP level after the second TACE. Therefore, a third TACE was done one month later, after which a firm abdomen over the epigastric area, tea-colored urine, and yellowish skin turgor appeared within two weeks. Abdominal sonography showed dilated left IHDs and a greatly increased liver tumor burden, which were confirmed by a contrast-enhanced MRI. Although we hypothesized that mutations in hypoxia-related genes may contribute to the disease progression after TACE, we did not perform liver biopsies to confirm these mutations due to laboratory limitation in our hospital. We described an endoscopic retrograde cholangiopancreatography intervention for bile duct drainage and immunotherapy to the patient and her husband, which had previously been described in the outpatient clinic before the series of TACE, but she refused and opted for hospice care.

**OUTCOME AND FOLLOW-UP**

For personal and religious reasons, the patient decided to pursue hospice care and was referred for home hospice care.

**DISCUSSION**

TACE is considered a first-line treatment for unresectable, multinodular, or intermediate-stage HCC, according to the European Association for the Study of the Liver and the American Association for the Study of Liver Diseases[2,3]. A systematic review of TACE therapy’s efficacy for HCC found an objective response rate of 52.5% (95% confidence interval: 43.6-61.5), and overall survival was 70.3% at one year and 32.4% at five years[4]. However, approximately half of intermediate-stage HCC patients had a poor response to TACE, and the best window for other anticancer therapies was missed due to TACE. In this report, our patient with extensive multinodular HCC had a poor response to the second TACE and rapidly progressed to BCLC stage C HCC within two weeks after the third. Despite the probability of the natural course of the disease, it appeared that TACE stimulated the tumors and rapidly led to this upward stage of transition. In the past decade, several studies have revealed that hypoxia can cause tumor development, tumor angiogenesis, and drug resistance and even promote metastasis, which are mediated by hypoxia-markers including hypoxia-inducible factors (HIF), COX-2, AMP-activated protein kinase and glucose transporter[8-14]. It was reasonably hypothesized that the strong tumor hypoxia induced by TACE may cause drug resistance, rapid growth, invasion, angiogenesis, and metastasis of tumors instead of killing tumor cells. Therefore, these hypoxia-markers might serve as indicators of an unfavourable prognosis in patients who undergoing TACE. While we speculated that mutations in hypoxia-related gene could potentially contribute to an increased tumor burden and disease progression, the clinical use of tests to assess these hypoxia markers is still limited in most hospitals. This may tend to occur in multinodular, large HCC tumors because of multiple regions of intra-tumoral hypoxia and increased expression of HIF[12]. In the prospective study of Tsai *et al*[5], 746 newly diagnosed HCC patients were enrolled, including 624 who received TACE as the primary therapy and 122 who received the best supportive care. Of 624 patients, 102 had a poor response at three months[5]. Among them, 44 died within three months, and 58 patients with rapid disease progressions had contraindications a subsequent TACE for residual tumors, including a high serum bilirubin level, distant metastases, cachexia, main portal vein thrombosis, and regional extrahepatic invasion[5]. The patients with poor responses were of a higher proportion of performance status ≥ 1, albumin ≤ 3.8 g/dL, Child-Turcotte-Pugh class B, AFP > 40 ng/mL, total tumor volume > 65 cm3, and vascular invasion, than those without poor responses[5]. Furthermore, a retrospective study found that, among intermediate-stage HCC patients who received repeated TACE as the primary therapy, a multiple tumor number of ≥ 4 and a large tumor size of ≥ 5 cm were independent risks for a shorter time of progression from BCLC stage B to stage C than that of their counterparts[6]. Both studies showed that TACE may be ineffective for intermediate-stage HCC patients with a large tumor burden and may even lead to rapid disease progression, such as portal invasion, extrahepatic spread, and worse liver function. However, the mechanisms are still poorly understood and should be explored in future studies to improve the treatment outcomes of HCC.

Due to the nature of the case report, causality cannot be established and external validity is limited. However, it was unusual for tumors to grow so rapidly in two weeks, indicating that they were likely caused by the third TACE. Large cohort studies and clinical trials are required to explore this relationship. A strength of this study is that our case identified a potential causal relationship in which TACE stimulates tumors to grow rapidly with portal invasion, bile duct invasion, or extrahepatic spread—that is, the upward stage of transition—among specific intermediate-stage patients with a large tumor burden. This case may raise global awareness of the current limitations of the BCLC staging system and contribute to reducing the incidence of ineffective and even harmful TACE treatment in specific patients.

**CONCLUSION**

Several guidelines recommend TACE as the first-line treatment for intermediate-stage HCC patients. However, even without portal invasion or extrahepatic spread, cases with a large tumor burden and multi-foci tumor infiltration tends not to respond to TACE, and HCC may even increase dramatically.

**ACKNOWLEDGEMENTS**

We extend our gratitude to the reviewers and editor for their valuable feedback.

**REFERENCES**

1 **Sung H**, Ferlay J, Siegel RL, Laversanne M, Soerjomataram I, Jemal A, Bray F. Global Cancer Statistics 2020: GLOBOCAN Estimates of Incidence and Mortality Worldwide for 36 Cancers in 185 Countries. *CA Cancer J Clin* 2021; **71**: 209-249 [PMID: 33538338 DOI: 10.3322/caac.21660]

2 **Heimbach JK**, Kulik LM, Finn RS, Sirlin CB, Abecassis MM, Roberts LR, Zhu AX, Murad MH, Marrero JA. AASLD guidelines for the treatment of hepatocellular carcinoma. *Hepatology* 2018; **67**: 358-380 [PMID: 28130846 DOI: 10.1002/hep.29086]

3 **European Association for the Study of the Liver.** European Association for the Study of the Liver. EASL Clinical Practice Guidelines: Management of hepatocellular carcinoma. *J Hepatol* 2018; **69**: 182-236 [PMID: 29628281 DOI: 10.1016/j.jhep.2018.03.019]

4 **Lencioni R**, de Baere T, Soulen MC, Rilling WS, Geschwind JF. Lipiodol transarterial chemoembolization for hepatocellular carcinoma: A systematic review of efficacy and safety data. *Hepatology* 2016; **64**: 106-116 [PMID: 26765068 DOI: 10.1002/hep.28453]

5 **Tsai YJ**, Hsu CY, Huang YH, Su CW, Lin HC, Lee RC, Chiang JH, Huo TI, Lee SD. Early identification of poor responders to transarterial chemoembolization for hepatocellular carcinoma. *Hepatol Int* 2011; **5**: 975-984 [PMID: 21533669 DOI: 10.1007/s12072-011-9276-9]

6 **Kim HY**, Park JW, Joo J, Jung SJ, An S, Woo SM, Kim HB, Koh YH, Lee WJ, Kim CM. Severity and timing of progression predict refractoriness to transarterial chemoembolization in hepatocellular carcinoma. *J Gastroenterol Hepatol* 2012; **27**: 1051-1056 [PMID: 22098152 DOI: 10.1111/j.1440-1746.2011.06963.x]

7 **Decaens T**, Roudot-Thoraval F, Hadni-Bresson S, Meyer C, Gugenheim J, Durand F, Bernard PH, Boillot O, Sulpice L, Calmus Y, Hardwigsen J, Ducerf C, Pageaux GP, Dharancy S, Chazouilleres O, Cherqui D, Duvoux C. Impact of UCSF criteria according to pre- and post-OLT tumor features: analysis of 479 patients listed for HCC with a short waiting time. *Liver Transpl* 2006; **12**: 1761-1769 [PMID: 16964590 DOI: 10.1002/Lt.20884]

8 **Godet I**, Shin YJ, Ju JA, Ye IC, Wang G, Gilkes DM. Fate-mapping post-hypoxic tumor cells reveals a ROS-resistant phenotype that promotes metastasis. *Nat Commun* 2019; **10**: 4862 [PMID: 31649238 DOI: 10.1038/s41467-019-12412-1]

9 **Gilkes DM**, Semenza GL, Wirtz D. Hypoxia and the extracellular matrix: drivers of tumour metastasis. *Nat Rev Cancer* 2014; **14**: 430-439 [PMID: 24827502 DOI: 10.1038/nrc3726]

10 **Erin N**, Grahovac J, Brozovic A, Efferth T. Tumor microenvironment and epithelial mesenchymal transition as targets to overcome tumor multidrug resistance. *Drug Resist Updat* 2020; **53**: 100715 [PMID: 32679188 DOI: 10.1016/j.drup.2020.100715]

11 **Lai JP**, Conley A, Knudsen BS, Guindi M. Hypoxia after transarterial chemoembolization may trigger a progenitor cell phenotype in hepatocellular carcinoma. *Histopathology* 2015; **67**: 442-450 [PMID: 25425262 DOI: 10.1111/his.12623]

12 **Semenza GL**. Intratumoral Hypoxia and Mechanisms of Immune Evasion Mediated by Hypoxia-Inducible Factors. *Physiology (Bethesda)* 2021; **36**: 73-83 [PMID: 33595388 DOI: 10.1152/physiol.00034.2020]

13 **Huang M**, Wang L, Chen J, Bai M, Zhou C, Liu S, Lin Q. Regulation of COX-2 expression and epithelial-to-mesenchymal transition by hypoxia-inducible factor-1α is associated with poor prognosis in hepatocellular carcinoma patients post TACE surgery. *Int J Oncol* 2016; **48**: 2144-2154 [PMID: 26984380 DOI: 10.3892/ijo.2016.3421]

14 **Qu K**, Yan Z, Wu Y, Chen Y, Qu P, Xu X, Yuan P, Huang X, Xing J, Zhang H, Liu C, Zhang J. Transarterial chemoembolization aggravated peritumoral fibrosis *via* hypoxia-inducible factor-1α dependent pathway in hepatocellular carcinoma. *J Gastroenterol Hepatol* 2015; **30**: 925-932 [PMID: 25641377 DOI: 10.1111/jgh.12873]

**Footnotes**

**Informed consent statement:** Informed written consent was obtained from the patient for publication of this report and any accompanying images.

**Conflict-of-interest statement:** The authors declare that they have no conflict of interest to disclose.

**CARE Checklist (2016) statement:** The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised in accordance with it.

**Open-Access:** This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: https://creativecommons.org/Licenses/by-nc/4.0/

**Provenance and peer review:** Unsolicited article; Externally peer reviewed.

**Peer-review model:** Single blind

**Peer-review started:** February 18, 2023

**First decision:** May 16, 2023

**Article in press:** June 6, 2023

**Specialty type:** Gastroenterology and hepatology

**Country/Territory of origin:** Taiwan

**Peer-review report’s scientific quality classification**

Grade A (Excellent): 0

Grade B (Very good): 0

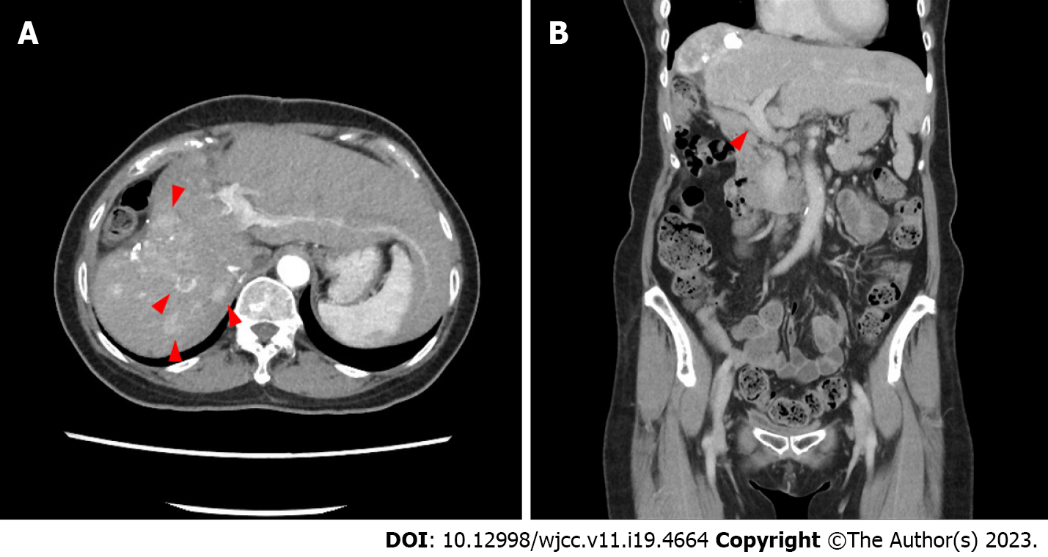
Grade C (Good): C

Grade D (Fair): D

Grade E (Poor): 0

**P-Reviewer:** koganti SB, United States; Radhakrishnan K, South Korea **S-Editor:** Liu JH **L-Editor:** A **P-Editor:** Liu JH

**Figure Legends**



**Figure 1 Abdominal computed tomography scans following the second transarterial chemoembolization.** A: Transverse plane showing multiple hepatocellular carcinoma nodules in the right lobe of the liver (red arrows); B: Coronal plane showing an unobstructed portal vein and typical bile ducts (red arrow).



**Figure 2 Contrast-enhanced magnetic resonance imaging showed rapid progression of hepatocellular carcinoma volume with involvement of the left intra-hepatic duct and portal vein tumor invasions (red arrow) two weeks after the third transarterial chemoembolization.**



Published by **Baishideng Publishing Group Inc**

7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

**Telephone:** +1-925-3991568

**E-mail:** bpgoffice@wjgnet.com

**Help Desk:** https://www.f6publishing.com/helpdesk

https://www.wjgnet.com



**© 2023 Baishideng Publishing Group Inc. All rights reserved.**