Rapid aneurysm growth after transarterial chemoembolization

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Chemotherapy has been anecdotally related to aneurysm growth, but no correlation has been noted to date for localized transarterial chemoembolization. We present the case of a 64-year-old man with clearly documented accelerated aortic and iliac artery aneurysmal dilation after two rounds of transarterial chemoembolization for hepatocellular carcinoma.

Given the large size with rapid growth of his aneurysms and inability to be listed for transplant consideration before repair, he was offered endovascular repair and was successfully treated. (J Vasc Surg Cases 2015;1:65-7.)

CASE REPORT

A 64-year-old Caucasian man was referred to our clinic for evaluation of aortoiliac aneurysmal disease identified during workup for a liver transplant. The patientʼs medical history was significant for hepatocellular carcinoma (HCC) secondary to hepatitis C, portal hypertension, chronic obstructive pulmonary disease, and a resolved hepatitis B infection. He received treatment with ribavirin and interferon for hepatitis C, resulting in an undetectable viral load for 11 months leading up to his presentation. Of note, he had no prior or family history of aneurysmal diseases or vasculitides.

On initial magnetic resonance imaging during the HCC workup, a noninflammatory infrarenal aneurysm measuring 3.9 cm was identified in November 2013, with a right common iliac artery (RCIA) aneurysm measuring 3.3 cm and a left common iliac artery (LCIA) measuring 2.8 cm (Fig 1).

Three rounds of TACE were performed over the course of 5 months while the patient was being evaluated for liver transplantation. During round one (March 2014), doxorubicin (50 mg) and lipiodol (1 mL) were used to the right hepatic lobe. Computed tomography angiography imaging in June 2014 revealed progressive aneurysmal dilatation, with measurements of the infrarenal aorta at 4.8 cm, RCIA at 4.3 cm, and the LCIA at 3.3 cm (Fig 3).

The patient was offered endovascular repair of the aneurysm given the significant growth, the size of the aneurysm, and the transplant committeeʼs decision to exclude the patient for liver transplantation secondary to the aneurysms. After his third TACE procedure using lipiodol (10 mL) as the sole agent, the patient elected to undergo an endovascular repair.

Off-label techniques, as previously described, were used to perform successful exclusion of the aneurysm, with bilateral hypogastric artery preservation, and he was discharged home on postoperative day 2 without incident.3,4

DISCUSSION

This is the first documented case of accelerated CIA aneurysm growth after TACE treatment. TACE is an accepted treatment for unresectable HCC.5-7 The procedure involves gaining access to the hepatic arteries for targeted chemotherapy, theoretically allowing much lower doses of chemotherapeutic agents to be used with little to no systemic influence of the drug. Hepatic artery aneurysms developing after TACE have been reported, hypothesized to form secondary to trauma to vessels during the procedure or as a side effect of the chemotherapy itself, but no known literature has documented iliac aneurysmal growth rates after TACE.8,9

Although the growth rate of CIA aneurysms is debated, the maximum reported growth rate in the literature is defined as 0.32 cm/yr.10,11 Our patient presented with significantly greater growth. The RCIA and LCIA grew 1.0 cm during a period of just 7 months, and 0.7 cm and
0.5 cm, respectively, in just 1 month after two rounds of TACE. Aneurysmal disease as a result of chemotherapy is debated, with a recent report demonstrating no association between aortic aneurysm growth and chemotherapy.12

CONCLUSIONS

Despite this study, there remains a broadly held concern regarding the effects of these cytotoxic agents

Fig 1. Magnetic resonance imaging of the abdomen and pelvis, November 2013.

Fig 2. Magnetic resonance imaging abdomen and pelvis, April 2014.
often prompting aneurysm repair at an earlier time than would otherwise be considered. There is no prevailing explanation for aneurysmal growth with chemotherapy use; however, some have questioned altered generation of DNA, collagen and elastin, release of inflammatory markers, and disturbance in smooth muscle proliferation.12 In theory, TACE is a localized treatment and should not have systemic effects, as are believed to be seen with other cytotoxic treatments, but this first documented incidence indicates a need for further review and awareness.

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