CASE REPORT

Signet-ring cell carcinoma of ampulla of Vater: Contrast-enhanced ultrasound findings

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INTRODUCTION

Signet-ring cell carcinoma (SRCC) usually occurs in the gastrointestinal tract. The World Health Organization (WHO) defines it as a special type or a variant of gastrointestinal adenocarcinoma. SRCCs may exist alone or coexist with any other types of malignant gastrointestinal tumors. SRCC is very rarely found among carcinomas of the ampulla of Vater. Here, we describe one patient with SRCC in the ampulla of Vater, which was found by contrast-enhanced ultrasound (CEUS). This is the first case reported in literature, which was successfully diagnosed with CEUS.

CASE REPORT

A 38-year-old woman was hospitalized because of pruritus for 13 d, and dermatic and scleral jaundice with urine the color of bean oil for 5 d. The stool s had a silver color. The patient had nausea but without vomiting, fever and abdominal pain. She lost weight of about 3 kg in 1 mo. She had a history of surgery for left breast adenoma at another institution several years ago.

Physical examination revealed mucocutaneous jaundice without tenderness in the epigastrium. The laboratory test results showed that white blood cells and hemoglobin were normal. Biochemical tests demonstrated the presence of glutamate-pyruvate transaminase at 446.5 IU/L (normal range, 0-40), glutamic-oxal (o) acetic transaminase at 277.3 IU/L (normal range, 5-34), alkaline phosphatase at 744.1 IU/L (normal range, 40-150), \(\gamma\)-glutamyltransferase at 1687.2 IU/L (normal range, 9-64), total bilirubin at 186.6 mg/dL (normal range, 3.4-20.5), direct bilirubin at 154.2 mg/dL (normal range, 0-8.6), and indirect bilirubin at 32.4 mg/dL (normal range, 3.4-11.9). The tumor markers of carcinoembryonic antigen were 4.74 ng/mL (normal range, 0-5), alpha fetoprotein 3.87 ng/mL (normal range, 0-9), and carbohydrate antigen 19-9 143.13 ng/mL (normal range, 0-37).

Endoscopic ultrasound (Figure 1) showed a heterogenic, hypoechoic mass with ill-defined margins at the junction of the common bile duct (CBD) and the main pancreatic duct (PMD)-ampulla of Vater.

Conventional gray-scale ultrasound using a Logiq 9 scanner (GE, USA) equipped with a C2-4 transducer with a central frequency of 3.5 MHz revealed that the
intrahepatic bile duct was dilated with a diameter of 0.8 cm, and the initial and intermediate portion of the CBD were dilated with a maximal diameter of 2.0 cm. The end part of the CBD suddenly became narrow, with a diameter of 0.7 cm, and there was no exact mass at the end of the CBD (Figure 2). CEUS was performed with low acoustic power, providing real-time imaging using low-mechanical index modes. Contrast-specific CEUS mode of contrast pulse sequencing was applied. The contrast agent, SonoVue (Brocca, Milan, Italy) (2.4 mL) was administered. The wall of the CBD began to enhance at 12 s after contrast agent was administered (Figure 3), while there was no obvious hyper-enhanced or hypo-enhanced lesion in the ampulla of Vater. A hypo-enhanced lesion about 1.7 cm × 1.6 cm with blurred borders in the ampulla of Vater was found from 20 s to 180 s, compared with the adjacent pancreas (Figure 4). At delayed phase (120 s after contrast agent administration), we scanned the whole liver and no abnormal enhanced lesions were found, indicating that there was no metastases in the liver.

The patient underwent a pancreato-duodenectomy with an extended lymphadenectomy and gastrectomy of 1/4 of the normal stomach. The mass was located in the ampulla of Vater with a size about 2.0 cm × 2.0 cm, brittle and protruding to the cavity of the duodenum. It had infiltrated to the periphery pancreatic tissue and adhered to the inferior vena cava. Lymph nodes of No. 16 were tumescent. The final pathological examination (Figure 5) showed that the cancer cells were widespread and polygonal, and the nuclei of the cells were located on one side, which are prominent signet-ring features. Final pathology confirmed an SRCC of the ampulla of Vater, and the pancreas and the whole wall of the duodenum were infiltrated with carcinoma. No distal or nodal metastases were identified. There was no evidence of lymphatic and vascular invasion. The ampullary cancer was
The properties of SonoVue and the high sensitivity of recent ultrasound equipment to the presence of microbubbles have shown that CEUS is potentially very useful in revealing many organs and vascular structures. It has been a rapidly evolving technique for clinical application. CEUS allows the assessment of the macrovasculature and microvasculature in different parenchymas, and the identification and characterization of lesions in organs. It has been reported that CEUS produced results very similar to those obtained with contrast-enhanced CT and magnetic resonance imaging in the characterization of various liver lesions. The causes of obstructive jaundice can be divided into two categories: tumorous and non-malignant stenosis. For non-malignant stenosis, such as acute or chronic inflammation of the papilla, fibroid stenosis at the end of the CBD can be irritated by cholesterol calculi or sludge at the end of the CBD; blood clots at the end of the CBD may cause obstructive jaundice too. However, the main cause of obstructive jaundice is the tumors arising from the ampulla of Vater, and mostly are malignant. In the diagnosis of obstructive jaundice, the emphasis should be laid on excluding the non-malignant reasons: non-shadowing stones, blood clots and sludge. This may influence the selection of therapy. The non-shadowing stones, blood clots and sludge may appear non-enhanced by CEUS because of an absence of blood supply. In the present case, the carcinoma showed iso-enhancement at an early stage after contrast agent administration, and obvious hypo-enhancement at the delayed phase, because it had intraluminal tumor tissue with blood supply; microbubbles are distributed within the blood and appear wherever there is a blood supply. Our case showed that CEUS may provide an efficient means of diagnosis of ampullary carcinomas. CEUS could offer real-time imaging of the microcirculation in the lesions. By CEUS, the lesion may be displayed much clearer than by conventional gray-scale ultrasound. It can also offer a good method in the discrimination of ampullary carcinoma from non-malignant lesions.

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