Fairly rare clear cell adenocarcinoma mimicking liver cancer: a case report

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Abstract

Background: Clear cell carcinoma commonly occurs in the ovary and kidney, and clear cell cholangiocarcinoma was rarely reported. Differential diagnosis which the origin of the tumor located on the liver surface is intrahepatic or extrahepatic was difficult. Herein, we report a case of clear cell adenocarcinoma mimicking liver cancer.

Case presentation: This was a 55-year-old female who had the tumor with cystic component in the liver. She was performed hepatectomy and diagnosed as clear cell adenocarcinoma. Histopathological evaluation revealed intracystic clear cell adenocarcinoma. The tumor has ductal structure including mucin and atypical nuclear with clear cytoplasm. The tumor was separated from the liver and the diaphragm. The expression of Pax8 was positive, but the expression CK7 and HNF1β was positive and that of CD10 and ER was negative, which indicate that the tumor has the feature of clear cell carcinoma of ovary, not renal cell carcinoma nor cholangiocarcinoma.

Conclusions: Our experience with this patient suggests that this tumor may originate from the endometriosis onto the diaphragm from the detailed results of immunohistochemical staining.

Keywords: Liver, Ovary, Clear cell carcinoma, Hepatectomy

Background

Clear cell carcinoma is generally thought to originate from ovary and kidney [1]. Clear cell carcinoma of the ovary is composed of glycogen-containing clear cells and hobnail cells [1, 2]. Clear cell carcinoma of the ovary also shares many similarities with renal clear cell carcinoma [3]. Primary renal cell carcinoma is sometimes implanted to the ovary or peritoneum [4]. Additionally, clear cell cholangiocarcinoma [5] and peritoneal clear cell carcinoma [6, 7] were also rarely reported. Although recent genomics research will reveal the difference of these carcinomas, differential diagnosis of the primary site is difficult.

Herein, we report a case of clear cell adenocarcinoma mimicking liver cancer.

Case presentation

A 55-year-old woman regularly visited our hospital as an outpatient because of hepatitis B occult infection. A liver tumor was pointed out by CT. CT revealed a protruding liver tumor located at segment 8 3 cm in size, which include cystic lesion (Fig. 1a). US and MRI reveal the same feature (Fig. 1b). There was no distant metastasis. The patient had no past or family history including gynecological illness. 18F-FDG PET revealed the accumulation of 18F-FDG, and maximum standard uptake value was 2.3. Laboratory results included a white blood cell count of 3200/μL and platelet count of 189,000/μL. Prothrombin time international normalized ratio was 1.02. Total serum bilirubin was 0.9 mg/dL, direct bilirubin 0.03 mg/dL, albumin 4.5 g/dL, aspartate aminotransferase 22 U/L, alanine aminotransferase 17 U/L, alkaline phosphatase 187 U/L, and gamma-glutamyltranspeptidase 49 U/L. Tumor markers such as CEA, CA19-9, AFP, and DCP were normal. HBs-antigen and HBc-antibody were positive, and HBs-antibody and HCV-antibody were negative. The Child–Pugh score was 5, grade A. She was diagnosed
as intrahepatic cystadenocarcinoma and received extended posterior segmentectomy including diaphragm. Macroscopic findings revealed the tumor buried to the liver with the intracystic hemorrhage (Fig. 1c). The protruded compartment was closely touched to the diaphragm. Microscopic findings revealed the tumor and hemorrhage within the cyst (Fig. 1d). Tumor was located between the liver and diaphragm.

Histopathological evaluation revealed intracystic clear cell adenocarcinoma. The tumor has ductal structure including mucin and atypical nuclear with clear cytoplasm (Fig. 2a). The tumor was separated from the liver and the diaphragm (Fig. 2b). There is no traffic with the bile duct and ovarian stroma. PAS staining was positive. There was lack of ovarian clear cell carcinoma’s features such as hobnail appearance. The expression of Pax8 (Fig. 2c) was positive, but the expression CK7 and HNF1β(Fig. 2d) was positive and that of CD10 and ER was negative, which indicate that the tumor has the feature of clear cell carcinoma of the ovary, not renal cell carcinoma nor cholangiocarcinoma. The patient was discharged 13 days after surgery with no complication. According to the microscopic findings, gynecological interview and examinations after hepatic resection were performed, but there were no sign of menstrual irregularity and genital bleeding, and no endometriosis.

**Conclusions**

Clear cell carcinoma is characterized by clear cells containing glycogen arranged in tubular, papillary, and solid patterns, which commonly occurs in the ovary and kidney [1, 2]. On the other hand, clear cell cholangiocarcinoma was rarely reported [5]. Differential diagnosis which the origin of the tumor is intrahepatic or extrahepatic was difficult in this case. Microscopic findings revealed membrane of cyst was separated from the liver and diaphragm. In this case, there was no malignant tumor in the ovary or kidney and there was no evidence of endometriosis at the peritoneum. Fujiki et al. reported a case of diaphragmatic clear cell carcinoma in a patient with a medical history of ovarian endometriosis [8]. The clear cells and hobnail cells were observed in the diaphragmatic tumor. The hobnail cells were not in this case, but the immunohistochemical stain such as PAX8 and HNF1β. Pax8 is a crucial transcription factor for organogenesis of the thyroid gland, kidney, and müllerian system and is highly found in highly ovarian and renal clear cell carcinoma.
cell carcinomas [9]. HNF1β is also a transcription factor involved in glucose homeostasis and anti-apoptosis, and recent immunohistochemical studies have shown that it is frequently and highly expressed by ovarian clear cell carcinoma [10]. The great majority of ovarian clear cell carcinomas reported as a HNF-1β-positive and ER-negative immunoprofile [11]. CK7 was the subtype of cytokeratin, and CD10 was a cell surface enzyme with neutral metalloendopeptidase activity. These were used for the differential diagnosis of renal clear cell carcinoma [12]. In renal clear cell carcinoma, CK7 is negative and CD10 is positive. In ovarian clear cell carcinoma, CK is positive and CD10 is negative, which is similar to this case.

Primary peritoneal clear cell carcinoma (PPCC) is very rare. It was firstly described in 1990 [6], and so far, only 11 cases have been reported in the English literature [7]. The differential diagnosis between PPCC and ovarian clear cell carcinoma was difficult.

We previously reported that rare metastases which appear to be buried in the liver parenchyma on US or CT were closely touched under the diaphragm [13]. In this case, the tumor was not invaded to both liver and diaphragm mimicking liver cancer. That may be implanted from ovarian endometriosis onto diaphragm.

In conclusion, we presented a case of clear cell carcinoma mimicking liver cancer. We suspected that this tumor may originate from the endometriosis onto the diaphragm from the detailed results of immunohistochemical staining.

**Abbreviations**

18F-FDG-PET: 2-Deoxy-2-[fluorine-18]fluoro-D-glucose integrated with computed tomography; AFP: α-fetoprotein; CK: Cytokeratin; CT: Computed tomography; DCP: Des-γ-carboxy prothrombin; ER: Estrogen receptor; MRI: Magnetic resonance imaging; PAX8: Paired-box gene 8; US: Ultrasound sonography

**Authors’ contributions**

NH reported the case and wrote the manuscript. KA, KH, TY, NI, TI, AW, and NK performed the surgery and perioperative management of the patient and helped draft the manuscript. HI and SA examined the immunohistochemical staining. TM, MS, HK, and KS participated in revising the manuscript critically. All authors read and approved the final manuscript.

**Ethics approval and consent to participate**

This case report was performed in accordance with the Declaration of Helsinki, and we got the consent from the patient to publish according to our institutional consent form.

**Consent for publication**

We had consent from the patient to publish.

**Competing interests**

The authors declare that they have no competing interests.

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References
1. Rivkine E, Jakubowicz D, Marciano L, Polland C, Poncelet C, Ziol M, Barrat C. Hepatic endometrioma: a case report and review of the literature: report of a case. Surg Today. 2013;43(10):1188–93.
2. Terada T, Kawaguchi M. Primary clear cell adenocarcinoma of peritoneum. Tohoku J Exp Med. 2005;206:271–5.
3. Ji JX, Wang YK, Cochrane DR, Huntsman DG. Clear cell carcinomas of the ovary and kidney: clarity through genomics. J Pathol. 2018;244(5):550–64.
4. Guneysu S, Guney N, Ozcan D, Sayilgan T, Ozakin E. Ovarian metastasis of a primary renal cell carcinoma: case report and review of literature. Eur J Gynaecol Oncol. 2010;31(3):339–41.
5. Haas S, Güttgemann I, Wolff M, Fischer HP. Intrahepatic clear cell cholangiocarcinoma: immunohistochemical aspects in a very rare type of cholangiocarcinoma. Am J Surg Pathol. 2007;31(6):902–6.
6. Evans H, Yates WA, Palmer WE, Cartwright RL, Antemann RW. Clear cell carcinoma of the sigmoid mesocolon: a tumor of the secondary Müllerian system. Am J Obstet Gynecol. 1990;162:161–3.
7. Insabato L, Natella V, Somma A, Persico M, Camera L, Losito NS, Masone S. Primary peritoneal clear cell carcinoma versus ovarian carcinoma versus malignant transformation of endometriosis: a vexing issue. Int J Surg Pathol. 2015;23(3):211–6.
8. Fujii K, Miyamoto H, Hashimoto S, Suzuki N, Takano Y, Teranishi Y, Sakuma H, Suzuki H. A case of diaphragmatic clear cell carcinoma in a patient with a medical history of ovarian endometriosis. Int J Clin Oncol. 2010;15(5):489–92.
9. Bowen NJ, Logani S, Dickerson EB, et al. Emerging roles for PAX8 in ovarian cancer and endosalpingeal development. Gynecol Oncol. 2007;104:331–7.
10. Hoang LN, Han G, McConechy M, Lau S, Chow C, Gilks CB, Huntsman DG, Köbel M, Lee CH. Immunohistochemical characterization of prototypical endometrial clear cell carcinoma—diagnostic utility of HNF-1B and oestrogen receptor. Histopathology. 2014;64(4):585–96.
11. Fadare O, Liang SX. Diagnostic utility of hepatocyte nuclear factor 1-beta immunoreactivity in endometrial carcinomas: lack of specificity for endometrial clear cell carcinoma. Appl Immunohistochem Mol Morphol. 2012;20(6):580–7.
12. Ohta Y, Suzuki T, Shikawa A, Mitsuwa T, Ota H. Expression of CD10 and cytokeratin in ovarian and renal clear cell carcinoma. Int J Gynecol Pathol. 2005;24(3):339–45.
13. Harimoto N, Shimada M, Shirabe K, Tanaka S, Taketomi A, Maehara S, Itoh S, Kitagawa D, Honda H, Maehara Y. Hepatobiliary and pancreatic peritoneal dissemination mimicking liver metastases. J Gastroenterol Hepatol. 2004;19(8):938.