Orthotopic Liver Transplantation for Primary Intrahepatic Adenosquamous Carcinoma Using A Marginal Graft: the First Case rRport

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Case report

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Abstract

Background.

Primary intrahepatic adenosquamous carcinoma (iASC) is a very rare subtype of cholangiocarcinoma (CCA), with a worse prognosis than adenocarcinoma. Hepatectomy is still the first choice for iASC, and liver transplantation (LT) for iASC has not been previously reported.

Case presentation

The young male patient with an unresectable primary iASC underwent a LT using a marginal fatty liver graft. The overall survival of this patient was 16 months, with a recurrence-free survival of 8 months. The patient had a good quality of life with normal liver function until 3 months before death.

Conclusions.

This is the first case report of LT as treatment of unresectable iASC using a marginal graft. Although the recipient may benefit from LT, LT for iASC should proceed with caution owing to its more aggressive features.

Background

Primary intrahepatic adenosquamous carcinoma (iASC) is defined by the coexistence of both squamous and glandular elements in the same lesion histologically. iASC is clinically more aggressive with a worse prognosis, which accounts for only 2–3% of cholangiocarcinoma (CCA), and a majority of patients died within 1 year¹. We herein describe a patient with primary iASC who underwent orthotopic liver transplantation (LT). To our knowledge, this is the first reported case for iASC treated by LT.

Case Presentation

A 28-year-old man was admitted to our hospital complaining of a 10-day history of fatigue, upper abdominal discomfort, and a 5-day history of jaundice. He admitted a history of ulcerative colitis for 2 years and congenital cyst of biliary track. No abnormalities but jaundice was found on systemic examinations. The relevant laboratory results were as follows: WBC 13.11×10^9 /L, alanine aminotransferase (ALT) 299 IU/L, total bilirubin (TBIL) 586.0 μ mol/L, albumin 26 g/L, carbohydrate antigen (CA) 19 – 9 453.5 U/mL, CA 125 21.60 U/mL, carcinoembryonic antigen (CEA) 1.73 ng/mL, alpha-fetoprotein (AFP) 4.69 ng/mL, Hepatitis B and hepatitis C were both negative. Abdominal enhanced magnetic resonance imaging (MRI) and magnetic resonance cholangiopancreatography (MRCP) revealed an irregular mass (approximately 48mm × 44 mm in its dimensions) mainly located at segment VIII (S8) of the liver involving the hepatic hilum with mild delayed enhancement, a small metastatic lesion at segment VII (S7), and several small sporadic cysts in the liver (Fig. 1A-C). Positron emission tomography / computed tomography (PET/CT) clearly demonstrated hotspots (SUV max: 7.7 for the primary mass

and 4.2 for the metastatic lesion) at the same locations as the tumors detected by MRI, but no hot spots for lymph nodes and distant metastases were observed (Fig. 1D). Taking together, the clinical preoperative diagnosis of intrahepatic perihilar CCA with S7 metastasis was made and it was unresectable.

After the ethical approval and consent were clearly and fully obtained from the Ruijin Hospital's Ethical Committee, this patient was enrolled in the national waiting list of LT. A marginal fatty liver graft from a donor of brain death was allocated via the China Organ Transplant Response System two months after admission. After the percutaneous transhepatic cholangiodrainage (PTCD), LT with hepatoduodenal ligament lymph node skeletal dissection was performed successfully. The postoperative initial immunosuppression consisted of tacrolimus and mycophenolate without hormone. Histopathologically, the lesions were consisted of malignant squamous cells and a glandular component, accompanied by invasion of the perihilar bile ducts and metastases of the lymph nodes (Fig. 2A-C). Immunohistochemical study showed p40 and p63 protein were strongly positive (Fig. 3A and B). Meanwhile, cytokeratin 7 (CK-7) and mucin 1 (MUC1) labelling adenocarcinoma were strongly positive where p40 and p63 were negative (Fig. 3C and D).

The patient recovered well and was discharged 14 days postoperatively. On postoperative four weeks, the immunosuppressive regimen was converted to sirolimus-based immunosuppressive therapy. The patient received the first cycle of capecitabine chemotherapy 2 months after surgery and totally 4 cycles. Despite of intensive treatment strategies, this patient showed an increase in CA19-9 eight months after surgery without any discomforts (Fig. 4). Chest computed tomography (CT) revealed tumor metastases of the thoracic vertebrae. The patient then received a reduction of immunosuppressive drugs, systemic capecitabine chemotherapy, and infusion of zoledronic acid which has a significant clinical benefit in bone metastases from multiple tumor types². The gene mutations related to CCA in peripheral circulating tumor DNA (ctDNA) were detected by next-generation sequencing in order to identify if there were effective targeted drugs. Unfortunately, the results showed no mutations of typical genes, like EGFR, IDH1, IDH2³.

This patient with iASC died from multiple metastases of tumor eventually. The overall survival and recurrence-free survival are 16 months and 8 months, correspondingly (Fig. 5). Of note, the patient had a good quality of life with normal liver function until 3 months before death.

Discussion And Conclusion

Primary iASC is a very rare histological type of CCA, consisting of adenocarcinoma and squamous cell carcinoma components in the same tumor⁴. Since 1971, when iASC was first reported by Pianzola and Durt⁵, including our case, about 90 cases have been published. Due to its rarity, the clinicopathological features and biological behaviors have been incompletely elucidated. The main initial symptoms of iASC are fever, weight loss and epigastric pain⁶, which are not characteristic. Jaundice may occur in biliary obstruction due to hilar bile duct invasion or biliary lithiasis.

The common tumor markers for iASC are CA 19 – 9 and CEA. However, CA 19 – 9 can also be very high in obstructive jaundice⁷. In this case, tumor was sensitive to CA 19 – 9, and an increase level of CA 19 – 9 eventually suggested tumor recurrence.

Enhanced CT and MRI with MRCP are first choice to assess the tumor and vascular involvement. ERCP can place biliary stent to drain bile in patients with biliary obstruction while invasiveness and low biopsy positive rate limit its clinical use. PET/CT can be a reliable tool for detection of CCA and metastatic lymph nodes with an accuracy of 80% approximately⁸. Thus, laparoscopic lymph node biopsy closer to the time of transplantation are recommended in order to rule out nodal metastases regardless of imaging appearance.

Pathological features of iASC need to be elucidated. Cell keratinization, intercellular bridges, keratin pearl formation and/or dyskeratosis are considered to be squamous cell carcinoma components while various-size of gland formations and intracellular and/or intraluminal mucin give evidence to glandular differentiation. Moreover, immunohistochemistry (IHC) is crucial in diagnosis, like p40/63 and CK5/6 indicating squamous differentiation⁹, while CK-7, MUC1 are always positive in adenocarcinoma¹⁰. In accordance with the IHC findings, we successfully made a pathological diagnosis of iASC.

The pathogenesis of iASC is not well understood. Chronic inflammation associated with bile duct infection or biliary lithiasis and various congenital cysts of the biliary tract, are two main reasons of metaplastic changes in the biliary epithelium, ultimately ending with malignant neoplasia¹¹. In our case, persistent irritations from stones and congenital cysts might contribute to the development of iASC. Recent study show inflammatory bowel disease is also a risk factor for CCA¹², the chronic colitis in our patient may lead to iASC. Some authors^{13, 14} advocated the squamous cell carcinoma component might arise from squamous metaplasia of adenocarcinoma cells in the same lesion, this hypothesis has been confirmed by an experimental animal model, showing squamous cell carcinoma is a result of histopathology alteration from adenocarcinoma as a transitional form¹⁵. In this case, all tumors contained squamous cell carcinoma and adenocarcinoma without normal bile duct epithelium as hypothesis.

The first-line treatment of choice for iASC is hepatectomy. Unfortunately, most patients with iASC are diagnosed with either unresectable locally-advanced or metastatic disease at presentation. In such unresectable cases, LT is theoretically an attractive option, as it offers the maximum resection margin regardless of the status of the liver function and the volume of residual liver. The prognosis of iASC has been reported to be extremely poor, even with resection. Kobayashi⁶ analyzed that the mean survival of 34 patients with iASC was only 8.74 months.

Historically, early experience with LT for intrahepatic CCA (iCCA) was discouraging with essentially poor overall survival rates and increased recurrence¹⁶. Due to the scarce resources of grafts, the main concern about offering a liver donor to these patients is the influence on other patients on the waiting list. Therefore, LT to treat patients with CCA is confined to those with an unresectable hilar CCA especially

meeting the “Mayo clinic protocol” or with “very early” (single tumor < 2 cm) iCCA in most liver transplant centers¹⁶. Obviously, the patient in our case was beyond these rigorous selection standards for CCA. Considering this young patient’s age and no extrahepatic metastases, LT using a marginal fatty liver donor which was rejected in patients on waiting list was eventually approved by the organ transplantation ethic committee of our hospital to prolong his life maximumly. Due to the different organ allocation policies in various countries, in some with relatively sufficient donors, the use of marginal liver grafts as exploratory therapy may give some patients with advanced liver tumors a chance to cure.

In view of the patient’s poor liver function, preoperative adjuvant therapy was not performed, instead he received oral capecitabine chemotherapy 2 months after LT. Calcineurin inhibitors (CNIs) combined corticosteroid are regarded as the first line immunosuppressive therapy for LT, but some clinical studies have revealed a CNIs dose-dependent increase in the post-transplant risk of hepatocellular carcinoma (HCC) relapse¹⁷. Corticosteroid-free immune-suppression which is performed as a routine immunosuppressive regimen for HCC in our center has been reported to be safe for patients with HCC and showed an advantage in reducing the recurrence of HCC¹⁸. Mammalian target of rapamycin inhibitors, like sirolimus and everolimus, have shown an antineoplastic effect in vitro and clinical studies. In a meta-analysis, sirolimus was associated with a lower incidence of HCC recurrence after LT, compared with CNIs¹⁹. Thus, in our case, the patient received an initial steroid-free tacrolimus-based immunosuppressive regimen after LT and was converted tacrolimus to rapamycin as the central immunosuppressive drug 4 weeks after operation. The incision had healed 4 weeks after operation, so the switch to rapamycin would not add the incidence of its side effects that have been reported previously, like delayed wound healing, incisional hernias and arterial thrombosis²⁰.

In spite of the above intensive prevention measures in our case, the metastases were found about 8 months after surgery and progressed rapidly. The patient was given minimal immunosuppressive drugs, continued systemic capecitabine chemotherapy and palliative treatment without targeted therapy after metastases. Molecularly targeted therapy based on the key signaling pathways and mutated genes may be an option for unresectable or metastatic CCA patients, either alone or in combination with chemotherapy and immunotherapy²¹. CtDNA allows for the evaluation of patient-specific tumoral genetic and epigenetic alterations, and the mutations found in ctDNA which are highly representative in iCCA may bring a large number of potential therapeutic targets³. Unfortunately, the most frequently mutated genes in CCA were not detected in peripheral ctDNA, suggesting iASC might have a very different genomic profile from conventional CCA. The genotyping needs further studies to make a better understanding on this rare tumor. The overall survival of this patient was 16 months, which exceeded that of the most patients with iASC undergoing liver resection in previous literatures.

Undoubtedly, the rarity of iASC sets an obstacle for improving the diagnosis and treatment of such disease, more clinical data should be collected in the future. In this case, the use of marginal liver grafts in LT as exploratory therapy may give patients with advanced iASC a chance to cure. However, owing to the aggressive features of iASC and the lack of effective adjuvant drugs at present, LT for iASC should

proceed with caution. However, owing to the aggressive features of iASC and the lack of effective adjuvant drugs at present, LT for iASC should proceed with caution.

Abbreviations

AFP, alpha-fetoprotein

ALT, alanine aminotransferase

CA, carbohydrate antigen

CCA, cholangiocarcinoma

CEA, carcinoembryonic antigen

CK-7, cytokeratin 7

CNIs, calcineurin inhibitors

CT, computed tomography

ctDNA, circulating tumor DNA

HCC, hepatocellular carcinoma

iASC, intrahepatic adenosquamous carcinoma

iCCA, intrahepatic cholangiocarcinoma

IHC, immunohistochemistry

LT, liver transplantation

PTCD, percutaneous transhepatic cholangiodrainage

MRCP, magnetic resonance cholangiopancreatography

MRI, magnetic resonance imaging,

MUC1, mucin 1

PET/CT, positron emission tomography / computed tomography

TBIL, total bilirubin

Declarations

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Availability of data and materials

The clinical data used or analyzed in this case report are available from the corresponding author on reasonable request.

Author's Contributions

Peng Chen and Di Ma are co-first authors

All authors contributed to the writing and revision of the article. Peng C. and Di M. contributed equally to this work. Peng C., Ma D. and Tao L. contributed to the conception of study. Ruokun L., Xiaoqun Y., Tong H. and Debin Q. contributed to data collection and performance of the research.

Ethics approval and consent to participate

The ethical approval and consent were clearly and fully obtained from the Ethical Committee of Ruijin Hospital, the Affiliated Hospital of Shanghai Jiao Tong University School of Medicine. Patient was provided with written informed consent based on the Declaration of Helsinki

Consent for publication

Written informed consent was obtained from the patient for publication of this case report and accompanying images.

Competing interests

The authors declare that they have no competing interests.

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Figures

Figure 1

Selected diagnostic imaging findings of MRI, MRCP, and PET/CT. (A) Axial T2-weighted magnetic resonance image (MRI) and (B) coronal T2-weighted MRI show a hyperintense mass (black arrow) located at the perihilar area, with marked intrahepatic biliary dilatation in the adjacent liver (the metastatic lesion at segmental VII of the liver was not shown here). (C) MRCP shows a mass in the perihilar area (white arrow), with severe strictures at the hepatic perihilar biliary ducts and marked dilation of the adjacent biliary ducts. (D) PET-CT shows hotspots (SUV max: 7.7 for the big mass and 4.2 for the metastatic lesion) at the same locations as that detected by MRI (white short arrows). The tube of PTCD can be seen in the left hepatic bile duct (black arrow).

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Figure 2

H&E staining: showed the tumor was composed of two components: adenocarcinoma and squamous cell carcinoma. (H&E stain, ×200)

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Figure 3

Immunohistochemistry: for p63 and p40, the results reveal nuclear positive staining for squamous cell carcinoma (A and B), and adenocarcinoma cell staining for CK-7 and MUC1 is positive (C and D).
(immunohistochemistry staining: $\times 100$)

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Figure 4

Tumor markers and liver function: data showed significant elevated levels of CA19-9, ALT and TBIL before operation, suggesting poor liver function. Then CA19-9, ALT and TBIL quickly declined to normal levels after surgery. Noticeable increases in CA19-9 indicated tumor metastases 8 months after LT.

Figure 4

Tumor markers and liver function: data showed significant elevated levels of CA19-9, ALT and TBIL before operation, suggesting poor liver function. Then CA19-9, ALT and TBIL quickly declined to normal levels after surgery. Noticeable increases in CA19-9 indicated tumor metastases 8 months after LT.

Figure 5

Clinical timeline

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