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

**Manuscript NO:** 56585

**Manuscript Type:** CASE REPORT

**Primary chondrosarcoma of the liver: a case report**

Liu Z Y *et al.* Primary liver chondrosarcoma

Zhuo-Yan Liu, Xue-Mei Jin, Guang-Hai Yan, Guang-Yu Jin

**Zhuo-Yan Liu, Guang-Yu Jin,** Department of Radiology, Affiliated Hospital of Yanbian University, Yanji 133000, Jilin Province, China

**Xue-Mei Jin,** Department of Pathology, Affiliated Hospital of Yanbian University, Yanji 33000, Jilin Province, China

**Guang-Hai Yan,** Department of Anatomy, Basic Medical College, Yanbian University, Yanji 133002, Jilin Province, China

**Author contributions:** Liu ZY collected the data, performed the literature research and drafted the manuscript; Jin XM and Yan GH collected the data; Jin GY conceived the study design, interpreted all data and revised the manuscript in depth; All authors have read and approved final version of this manuscript for publication.

**Supported by** National Natural Science Foundation of China, No. 81860304 and No. 81860043; "Thirteenth Five-Year Plan" Science and Technology Project of Education Department of Jilin Province, No. JKH20191141KJ; and Natural Science Research Foundation of Jilin Province for Sciences and Technology, No. 20200201511JC.

**Corresponding author: Guang-Yu Jin, MD, Professor,** Department of Radiology, Affiliated Hospital of Yanbian University, No. 1327 Juzi Street, Yanji 133000, Jilin Province, China. kimguangyu@163.com

**Received:** May 6, 2020

**Revised:** June 16, 2020

**Accepted:** September 2, 2020

**Published online:** October 26, 2020

**Abstract**

BACKGROUND

Primary chondrosarcoma of the liver are extremely rare. Moreover, there are few reports focusing on typical clinical symptoms and imaging characteristics. Therefore, the diagnosis of chondrosarcoma of the liver remains a challenge.

CASE SUMMARY

A 59-year-old male was admitted due to a lesion occupying the right liver lobe that was found by physical examination. Magnetic resonance imaging showed a lobular mass with high T2 weighted image and low T1 weighted image with enhanced internal separation and edge in the right liver. He was diagnosed with liver cystadenoma by using magnetic resonance imaging. At 3 mo later, the magnetic resonance scan showed that the mass was enlarged. Laparoscopic liver tumor resection was performed with a pathological diagnosis of liver chondrosarcoma. Then he received a surgical resection for the recurrent lesion. However, intrahepatic and abdominal metastases were found again at 8 mo after the second operation. The patient then received conservative management and is now under follow-up.

CONCLUSION

Primary liver chondrosarcoma generally is presented as lobulated and heterogeneous density/signal, cystic, solid masses without calcification with enhanced edge, internal septa and solid part. The imaging features are closely related to pathology, which may be helpful for clinical diagnosis.

**Key Words:** Extraskeletal chondrosarcoma; Liver tumor; Differential diagnosis; Case report

**Citation:** Liu ZY, Jin XM, Yan GH, Jin GY. Primary chondrosarcoma of the liver: A case report. *World J Clin Cases* 2020; 8(20): 5042-5048

**URL:** https://www.wjgnet.com/2307-8960/full/v8/i20/5042.htm

**DOI:** https://dx.doi.org/10.12998/wjcc.v8.i20.5042

**Core Tip:** Primary chondrosarcoma of the liver is extremely rare. Moreover, there are few studies focusing on typical clinical symptoms and imaging characteristics. Therefore, the diagnosis of chondrosarcoma of the liver remains a challenge. This study analyzed the imaging characteristics of a case of primary liver chondrosarcoma to improve the understanding and diagnosis of this disease, which may help to improve the understanding of the disease.

**INTRODUCTION**

Chondrosarcoma is a malignancy characterized by cartilage matrix production by tumor cells, which mainly occurs in cartilage tissue or connective tissue. It more commonly involves the cartilage tissue or connective tissue of long bones of the extremities and pelvis and rarely occurs in soft tissue or other organs, such as the lung, heart, liver, pancreas and kidney, which are known as extraskeletal chondrosarcomas. Extraskeletal chondrosarcomas were first reported by Stout *et al*[1] and account for 3% of all soft tissue sarcomas. It has been shown that extraskeletal chondrosarcomas have a histopathological characteristic of chondrocyte-like differentiation but no obvious cartilage areas, no connection with bone structure or punctate calcification. Extraskeletal chondrosarcomas are classified as “uncertain differentiated tumors” in the 2013 World Health Organization Soft Tissue Sarcomas[2]. Extraskeletal chondrosarcomas originating in the liver are rarely observed. Therefore, imaging features of liver chondrosarcoma remain insufficient. In this study, we analyzed the imaging characteristics of a case with primary liver chondrosarcoma to improve the understanding and diagnosis of this disease.

**CASE PRESENTATION**

***Chief complaints***

A 59-year-old male was admitted after a physical examination discovered a lesion occupying the right liver lobe.

***History of present illness***

He had a 40-year history of hepatitis B and received oral “Entecavir” antiviral therapy.

***Laboratory examinations***

Laboratory tests and tests on tumor markers showed no abnormalities. Ultrasound showed an irregular hypoechoic region in the right posterior lobe of the liver with a poorly smoothed capsule. The uneven echo distribution and a 4.3 cm × 3.8 cm strip-shaped blood flow signal were observed within the hypoechoic region.

***Imaging examinations***

Magnetic resonance imaging (MRI) showed masses in the posterior lower lobe of the right lobe with a high T2 weighted image (T2WI) signal, low signal rings and separated low signals. The lesion had a low signal in the T1WI positive and negative phases with patches and other signals in the center, had a high diffusion weighted imaging signal with an intralesion patchy low signal and a significantly high signal apparent diffusion coefficient with a low intralesion signal. The lesion had clear boundaries and defoliated edges with a size of 3.1 cm × 4.3 cm × 3.7 cm. No dilatation of the internal bile duct was observed (Figure 1A-C). The edge of the lesion and its internal separating arteries were slightly enhanced during the enhanced scan and continuously enhanced during portal vein and the delayed phase. No lymphadenopathy was seen in the abdominal cavity (Figure 1D-F).

Diagnostic results of MRI: occupied lesion in the right lobe of the liver, suggesting cystadenomas. He did not receive further treatment. After 3 mo, he received an MRI scan again, which showed a larger mass with obvious lobulated margin in the right lobe with a size of 4.3 cm × 5.3 cm × 4.9 cm (Figures 2A and 2B). The relationship between lesions and colonic liver curvature were unclear. Therefore, he was admitted for surgical resection. Intraoperatively, a hard, cauliflower-like mass with size of 5 cm × 6 cm × 5 cm in the lower posterior segment of the right lobe of the moderate sclerosis liver was observed, which adhered to the colonic hepatic mesangium and the right renal capsule.

Pathological gross appearance showed that the cut surface of the tumor was gray-blue, nodular, with a sense of mucus, hard and brittle, and the tumor showed swelling to the surrounding liver tissue. Microscopic pathological examination showed a lobulated tumor with sparsely distributed tumor cells and cartilage pit formation (Figures 3A and 3B). Tumor cell nuclei were heterogeneous. The cells were mostly mononuclear but double nuclei were also observed. Mitotic cells were commonly observed. There was a transparent cartilage matrix among the cells. The tumor was presented as infiltrating growth to the surrounding liver tissue. The surrounding liver tissue showed nodular cirrhosis. Immunohistochemistry showed vimentin (+), S-100 (+) and Ki67 (60% +) (Figures 3C-E). The pathological diagnosis was chondrosarcoma of the liver.

Emission computed tomography (CT) was performed after surgery and found no space-occupying lesions. Therefore, he was diagnosed as primary liver chondrosarcoma. He had an uneventful postoperative course and received no other treatments. A relapse was found at 8 mo following surgery. He received laparoscopic tumor resection. At 8 mo following the second surgery, he was admitted due to melanosis, fatigue and intermittent midupper abdominal pain.

***Physical examination***

Physical examination showed a 10 cm × 15 cm mass in the right abdomen. CT scan showed multiple metastases within the liver and abdomen, presenting as multiple cystic masses with marginal lobes and enhanced posterior marginal enhancement, which were like the primary imaging findings (Figures 4A and 4B). Then, the patient received conservative management and is being followed up until now.

**FINAL DIAGNOSIS**

Primary liver chondrosarcoma after surgery, multiple metastases within the liver and abdomen.

**TREATMENT**

Abdominal drainage and anti-inflammatory and liver-protecting symptomatic treatment

**OUTCOME AND FOLLOW-UP**

The patient was discharged after treatment in good and stable condition, and the general condition of the patient was fair after follow-up.

**DISCUSSION**

Extraskeletal chondrosarcoma is a rare ectopic tumor. Over 90% of patients have a characteristic translocation that involves the *NR4A3* gene on chromosome 9[3]. Extraskeletal chondrosarcoma, which can be divided into myxoid, mesenchymal and differentiated types[4], is commonly observed in the deep soft tissues of the proximal limbs with the highest incidence at the age of 50-60 years and more common in males[5,6].

The incidence of chondrosarcoma in the liver is extremely low. There is no relevant literature abroad. Only two case reports focusing on primary hepatic chondrosarcoma in China were found[7,8]. Both cases were middle-aged/elderly and male. The main clinical symptom was upper abdominal discomfort but without obvious pain, and both lesions occurred in the right lobe of the liver. Their imaging manifestations were very similar to those of liver cystic tumors, and both were misdiagnosed as hepatic cystadenoma or cystadenocarcinoma before surgery. In this study, the MRI manifestations of liver chondrosarcoma were lobular masses with clear boundaries, obvious high T2WI and diffusion weighted imaging signals with uneven intratumor low signal separation. The lesion had a low T1WI signal without bleeding. There was a capsule around the lesion with a low T2WI signal. After enhancement, the edge of the lesion and its internal separation continually enhanced. CT showed a liquid density mass with lobulated margin and no calcification and same enhancement as that of the MRI.

Chondrosarcoma of the liver should be differentiated from hepatic cystadenoma or cystadenocarcinoma, hepatic hydatid cyst, secondary infection of liver cyst, liver abscess and cystic liver metastases. Hepatic cystadenoma or cystadenocarcinoma is a rare liver epithelial tumor that occurs in middle-aged and elderly females without history of hepatitis and cirrhosis[9]. The typical CT image characteristics of hepatic cystadenoma or cystadenocarcinoma are single or multiple circular cystic masses with smooth capsule wall, intracapsule separation and calcification[10]. In the enhanced phrase, the posterior capsule wall, the capsule nodules and intracapsule separation are enhanced lightly and moderately but not the capsule fluid, which is very similar to the imaging of the case in our study.

Most patients with hepatic hydatid cyst have a history of nomadic life. The hepatic hydatid cyst is usually presented as single or multiple cystic lesions of the liver with a thick cyst wall and intracyst separation, cyst wall calcification, floating ascus and scolices in the cyst[11]. For cases with secondary infection of liver cysts, the cyst wall is thickened uniformly and without wall nodules, small air bubbles can appear in the cyst, and a circular edema band is seen around the cyst wall[12]. The cyst wall is evenly enhanced following the enhancement phase. Clinical manifestations of patients with liver abscess are fever, elevated white blood cell levels and pain in the liver area[13]. CT of a liver abscess is manifested as circular or quasicircular low-density shadows in the liver parenchyma with even or uneven density[14]. CT values are higher than water but lower than liver parenchyma. Enhanced scanning shows lesions with a “double ring sign” and a “petal sign,” which is decreased in the delayed scan phase[15].

Primary lesions of cystic liver metastases are found in leiomyosarcoma of the digestive tract and colorectal cancer. Most lesions present as multiple lesions with a round shape, while a few are presented as lobulated shape[16]. When cystic lesions or cystic and parenchymal lesions coexist, the cyst wall can be thin and thick, wall nodules are sometimes seen, and the lesion may have incomplete separation and fluid-fluid level[17]. The capsule wall, wall nodules and separation can be enhanced during the enhanced phase.

Primary liver chondrosarcoma is extremely rare in the clinic, which also has no specified clinical manifestations. The diagnosis of liver chondrosarcoma mainly depends on pathological evaluation. Imaging characteristics can facilitate the early diagnosis of liver chondrosarcoma. The local recurrence rate of extraskeletal chondrosarcomas is 37%-48%[5], with a metastasis rate of 50%[5]. Extraskeletal chondrosarcoma is not sensitive to chemotherapy and radiotherapy. Currently, early surgical resection is considered the most effective treatment method for extraskeletal chondrosarcomas[18,19]. In this study, the patient was misdiagnosed as a hepatic cystadenoma due to lack of understanding of the imaging manifestations of liver chondrosarcoma. The lesion was increasingly enlarged, and surgical resection was performed after 3 mo. Thereafter, he received a surgical resection for recurrent lesion. However, intrahepatic and abdominal metastases were found again after the second operation. Thereafter, conservative treatments and regular follow-up was administrated.

**CONCLUSION**

In conclusion, primary liver chondrosarcoma is very rare, and it is difficult to accurately diagnose only by imaging findings. The diagnosis is confirmed mainly based on pathological evaluation and immunohistochemical phenotypes. The imaging characteristics of the patient with primary liver chondrosarcoma in this study may help to improve the understanding of the disease.

**REFERENCES**

1 **Stout AP**, Verner EW. Chondrosarcoma of the extraskeletal soft tissues. *Cancer* 1953; **6**: 581-590 [PMID: 13042781 DOI: 10.1002/1097-0142(195305)6]

2 **Balanzá R**, Arrangoiz R, Cordera F, Muñoz M, Luque-de-León E, Moreno E, Molinar L, Somerville N. Pulmonary extraskeletal myxoid chondrosarcoma: A case report and literature review. *Int J Surg Case Rep* 2016; **27**: 96-101 [PMID: 27591381 DOI: 10.1016/j.ijscr.2016.08.025]

3 **Paoluzzi L**, Ghesani M. Extraskeletal myxoid chondrosarcoma with massive pulmonary metastases. *Clin Sarcoma Res* 2018; **8**: 20 [PMID: 30534357 DOI: 10.1186/s13569-018-8]

4 **Kim H**, Nakaichi M, Itamoto K, Taura Y. Primary chondrosarcoma in the skull of a dog. *J Vet Sci* 2007; **8**: 99-101 [PMID: 17322781 DOI: 10.4142/jvs.2007.8.1.99]

5 **Zhang L**, Wang R, Xu R, Qin G, Yang L. Extraskeletal Myxoid Chondrosarcoma: A Comparative Study of Imaging and Pathology. *Biomed Res Int* 2018; **2018**: 9684268 [PMID: 29977924 DOI: 10.1155/2018/9684268]

6 **Drilon AD**, Popat S, Bhuchar G, D'Adamo DR, Keohan ML, Fisher C, Antonescu CR, Singer S, Brennan MF, Judson I, Maki RG. Extraskeletal myxoid chondrosarcoma: a retrospective review from 2 referral centers emphasizing long-term outcomes with surgery and chemotherapy. *Cancer* 2008; **113**: 3364-3371 [PMID: 18951519 DOI: 10.1002/cncr.23978]

7 **Zheng Q,** Xiao F, Luo C. Chondrosarcoma of liver: case report. *J Clin Exp Pathol* 2011; **27**: 794-795 [DOI: 10.13315/j.cnki.cjcep.2011.07.040]

8 **Zhou W,** Guo J. Chondrosarcoma of the liver: 1 case report. *Shandong Yiyao* 2011; **51**: 115 [doi: 10.3969/j.issn.1002-266X.2011.02.085]

9 **Martel G**, Alsharif J, Aubin JM, Marginean C, Mimeault R, Fairfull-Smith RJ, Mohammad WM, Balaa FK. The management of hepatobiliary cystadenomas: lessons learned. *HPB (Oxford)* 2013; **15**: 617-622 [PMID: 23458638 DOI: 10.1111/hpb.12026]

10 **Tholomier C**, Wang Y, Aleynikova O, Vanounou T, Pelletier JS. Biliary mucinous cystic neoplasm mimicking a hydatid cyst: a case report and literature review. *BMC Gastroenterol* 2019; **19**: 103 [PMID: 31234803 DOI: 10.1186/s12876-019-1001-5]

11 **Mihmanli M**, Idiz UO, Kaya C, Demir U, Bostanci O, Omeroglu S, Bozkurt E. Current status of diagnosis and treatment of hepatic echinococcosis. *World J Hepatol* 2016; **8**: 1169-1181 [PMID: 27729953 DOI: 10.4254/wjh.v8.i28.1169]

12 **Balbo BE**, Sapienza MT, Ono CR, Jayanthi SK, Dettoni JB, Castro I, Onuchic LF. Cyst infection in hospital-admitted autosomal dominant polycystic kidney disease patients is predominantly multifocal and associated with kidney and liver volume. *Braz J Med Biol Res* 2014; **47**: 584-593 [PMID: 24919173 DOI: 10.1590/1414-431x20143584]

13 **Law ST**, Li KK. Is hepatic neoplasm-related pyogenic liver abscess a distinct clinical entity? *World J Gastroenterol* 2012; **18**: 1110-1116 [PMID: 22416187 DOI: 10.3748/wjg.v18.i10.1110]

14 **Mortelé KJ**, Segatto E, Ros PR. The infected liver: radiologic-pathologic correlation. *Radiographics* 2004; **24**: 937-955 [PMID: 15256619 DOI: 10.1148/rg.244035719]

15 **Kang S,** Zhang H, Liu Q. The Correlation of the Early Stage of Pyogenic Liver Abscess Between Dynamic Contrast Enhanced CT Scan and MR Plain Scan. *Zhongguo CT and MRI Zazhi* 2013; **11**: 53-57 [doi: 10.3969/j.issn.1672-5131.2013.04.017]

16 **Kanas GP**, Taylor A, Primrose JN, Langeberg WJ, Kelsh MA, Mowat FS, Alexander DD, Choti MA, Poston G. Survival after liver resection in metastatic colorectal cancer: review and meta-analysis of prognostic factors. *Clin Epidemiol* 2012; **4**: 283-301 [PMID: 23152705 DOI: 10.2147/CLEP.S34285]

17 **Yu R,** Fu L, Li R. CT diagnosis of malignant tumors of the liver cystic. Journal of Clinicial Radiology 1999; **18**: 158-161 [PMID: 10700831 DOI: 10.1259/bjr.72.863.10700831]

18 **Davis EJ,** Wu YM, Robinson D, Schuetze SM, Baker LH, Athanikar J, Xuhong C, Kunju LP, Chinnaiyan AM, Chugh R. Next generation sequencing of extraskeletal myxoid chondrosarcoma. *Oncotarget* 2017; **8**: 21770-21777 [PMID: 28423517 DOI: 10.18632/oncotarget.15568]

19 **Elsayed AG**, Al-Qawasmi L, Katz H, Lebowicz Y. Extraskeletal Chondrosarcoma: Long-term Follow-up of a Patient with Metastatic Disease. *Cureus* 2018; **10**: e2709 [PMID: 30065902 DOI: 10.7759/cureus.2709]

**Footnotes**

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

**Conflict-of-interest statement:** All authors have no conflicts of interest.

**CARE Checklist (2016) statement:** The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).

**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: http://creativecommons.org/Licenses/by-nc/4.0/

**Manuscript source:** Unsolicited manuscript

**Peer-review started:** May 6, 2020

**First decision:** May 21, 2020

**Article in press:** September 2, 2020

**Specialty type:** Gastroenterology and hepatology

**Country/Territory of origin:** China

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

Grade A (Excellent): 0

Grade B (Very good): B

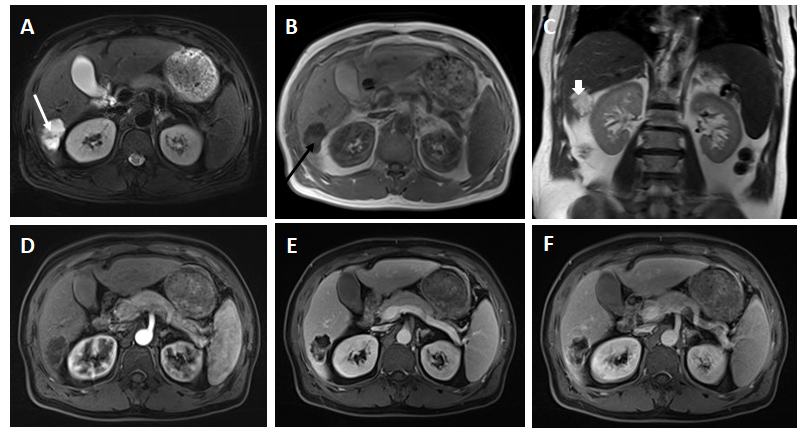
Grade C (Good): 0

Grade D (Fair): 0

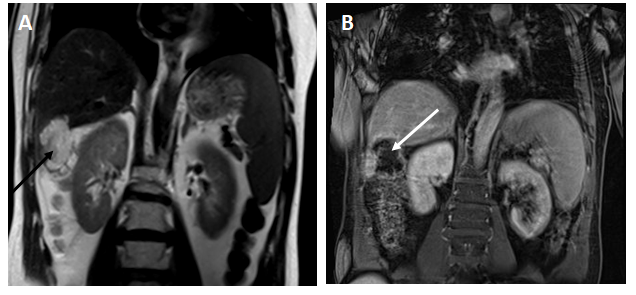
Grade E (Poor): 0

**P-Reviewer:** Ieni A **S-Editor:** Ma YJ **L-Editor:** Filipodia **P-Editor:** Liu JH

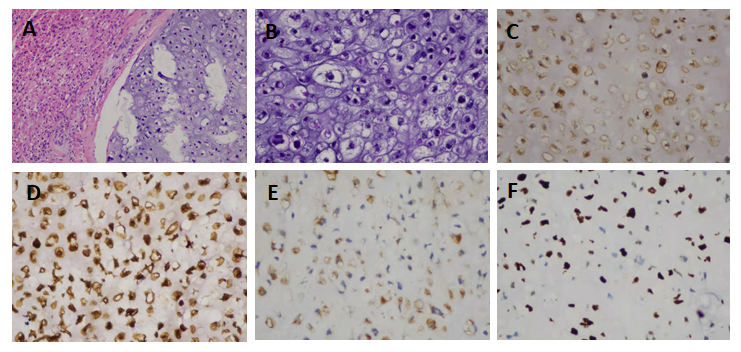
**Figure Legends**



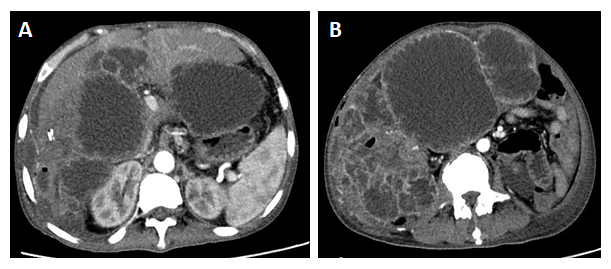
**Figure 1** **Magnetic resonance imaging at primary clinic visit.** A: a magnetic resonance (MR) T2 weighted image (T2WI) axis image showing lobular masses in the posterior lower lobe of the right lobe of the liver with clear borders, a high T2WI signal, intratumor separation with low signal and peripheral capsule with low signal(white arrow); B: an MR T1WI axis image showing the lesion was presented as low signal in T1WI with signals of patches and other (orange arrow); C: a coronal image of T2WI showing that the lesion protruded toward the lower edge of the liver (white arrowhead); D: MR enhanced arterial phase showed that the edge of the lesion and its internal separation are slightly enhanced; E: MR enhanced portal venous phase showed that the edge of the lesion and its internal separation are enhanced; F: MR enhanced delayed stage showed that the edge of the lesion and its internal separation are continually enhanced.



**Figure 2** **Magnetic resonance imaging after 3 mo.** A: a magnetic resonance T2 weighted image coronal image showed that the lesion was larger than before, and the border between the lesion and colonic liver curvature was unclear (orange arrow); B: a magnetic resonance T1 weight image coronal image (white arrow).



**Figure 3** **Histopathological evaluation.** A: Hematoxylin and eosin staining showed that tumor cells were unevenly distributed with cartilage pits, and the tumor was infiltrating and growing into surrounding liver tissue (magnifications 100 ×); B: Hematoxylin and eosin staining showed tumor cells with abnormal nuclei that were mostly mononuclear though occasionally binuclear (magnifications 200 ×); C: tumor CK protein positive; D: tumor vimentin protein positive; E: tumor S-100 protein positive; F: tumor Ki-67 immunostaining with 60% Ki-67 positive cells. (C-F immunohistochemistry, magnification 200 ×).



**Figure 4** **Magnetic resonance imaging at month 8 after the second surgery.** A: Computed tomography enhanced arterial phase image showing intrahepatic and abdominal metastases; B: Computed tomography enhanced arterial phase image showing metastatic lesions presenting as multiple cystic masses with enhanced edges.