Korean J Gastroenterol Vol. 74 No. 6, 356-361 https://doi.org/10.4166/kjg.2019.74.6.356 pISSN 1598-9992 eISSN 2233-6869

CASE REPORT



담도선섬유종의 자기공명영상 소견

이선영 1,2 , 김경원 2 , 정우경 3 , 유은실 4 , 장기택 5

연세대학교 의과대학 세브란스병원 영상의학과¹, 울산대학교 의과대학 서울아산병원 영상의학과², 성균관대학교 의과대학 삼성서울병원 영상의학과³, 울산대학교 의과대학 서울아산병원 병리과⁴, 성균관대학교 의과대학 삼성서울병원 병리과⁵

Magnetic Resonance Imaging Findings of Biliary Adenofibroma

Sunyoung Lee^{1,2}, Kyoung Won Kim², Woo Kyoung Jeong³, Eunsil Yu⁴ and Kee-Taek Jang⁵

Department of Radiology and Research Institute of Radiological Science, Severance Hospital, Yonsei University College of Medicine¹; Department of Radiology and Research Institute of Radiology, Asan Medical Center, University of Ulsan College of Medicine²; Department of Radiology and Center for Imaging Science, Samsung Medical Center, Sungkyunkwan University School of Medicine³; Department of Pathology, Asan Medical Center, University of Ulsan College of Medicine⁴; Department of Pathology, Samsung Medical Center, Sungkyunkwan University School of Medicine⁵, Seoul, Korea

Biliary adenofibroma is a rare tumor with a bile duct origin characterized by a complex tubulocystic non-mucin secreting biliary epithelium with abundant fibrous stroma. The MRI features of biliary adenofibroma are not well established. The authors encountered two patients with biliary adenofibroma and reviewed the literature focusing on the MRI findings. A well-circumscribed multicystic tumor with septal enhancement and no intrahepatic bile duct communication may be the characteristic MRI findings of biliary adenofibroma. (Korean J Gastroenterol 2019;74:356-361)

Key Words: Bile ducts; Adenofibroma; Magnetic resonance imaging

INTRODUCTION

Biliary adenofibroma is an extremely rare biliary epithelial tumor that is characterized by tubulocystic bile duct proliferation lined with non-mucin secreting biliary epithelium and supported by abundant fibroblastic stroma. The MRI findings of biliary adenofibroma are not well established owing to its rarity. This paper presents histopathologically confirmed cases of biliary adenofibroma found in two patients who underwent a preoperative MRI, along with a review of the relevant literature focusing on the MRI findings.

CASE REPORT

Case 1

A 63-year-old man with chronic hepatitis B and hepaticellular carcinoma treated previously with locoregional therapy was referred to the authors' medical institution for an evaluation of a solitary hepatic lesion increasing in size, as shown on the regular follow-up CT. The patient denied abdominal pain, weight loss, nausea, fever, or jaundice. The laboratory tests revealed a normal liver function. The tumor markers, including alpha-fetoprotein, protein induced by vitamin K

Received July 24, 2019. Revised September 10, 2019. Accepted September 30, 2019.

① This is an open access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/by-nc/4.0) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited. Copyright © 2019. Korean Society of Gastroenterology.

교신저자: 김경원, 05505, 서울시 송파구 올림픽로43길 88, 울산대학교 의과대학 서울아산병원 영상의학과

Correspondence to: Kyoung Won Kim, Department of Radiology and Research Institute of Radiology, Asan Medical Center, University of Ulsan College of Medicine, 88 Olympic-ro 43-gil, Songpa-gu, Seoul 05505, Korea. Tel: +82-2-3010-4400, Fax: +82-2-476-4719, E-mail: kimkw@amc.seoul.kr, ORCID: https://orcid.org/0000-0001-6471-6727

Financial support: This research was supported by the Basic Science Research Program through the National Research Foundation (NRF) of Korea, funded by the Ministry of Science, ICT, and Future Planning (no. 2017R1E1A1A03070961).

Conflict of interest: None.

absence or antagonist-II, CA 19-9, and CEA, were not elevated.

The CT scans obtained at an outside hospital revealed a tumor size of 1.1×0.7 cm, 2.3×1.4 cm, and 3.6×3.0 cm 5, 3, and 1 year ago, respectively, presenting a slowly growing nature. MRI demonstrated a well-circumscribed multiseptated mass within segments IV and VIII of the liver, measuring 4.7×4.5 cm. The T2-weighted image revealed a multicystic tumor with bright signal intensity (Fig. 1A, B). The precontrast T1-weighted image of the tumor showed low signal intensity (Fig. 1C). After gadolinium administration, thin septa within the tumor were enhanced (Fig. 1D). Communication with the intrahepatic bile ducts was not evident on MRI. No lymphadenopathy was observed.

The patient underwent a central bisegmentectomy. The gross pathological examination showed a well-circumscribed

multicystic tumor, measuring 4.5×4.3 cm with thin fibrous septa (Fig. 2A). The tumor was comprised of tubulo-glandular and microcystic structures embedded in the fibrous stroma (Fig. 2B). The tubules and cysts were lined with cuboidal or low columnar epithelial cells. The epithelial component did not contain atypia. The background stroma contained scattered spindle cells. The stroma was fibrous, but ovarian-like stroma was not detected (Fig. 2C). No communication was observed between the lesion and intrahepatic bile ducts, either grossly or microscopically. Immunohistochemically, the epithelial component was positive for cytokeratins 7 and 19, suggesting a bile duct origin. P53 was focally positive. Ki-67/index proliferation was low (<2%). The stromal cells were stained with α -smooth muscle actin (Fig. 2D). The background liver showed septal fibrosis and features of chronic

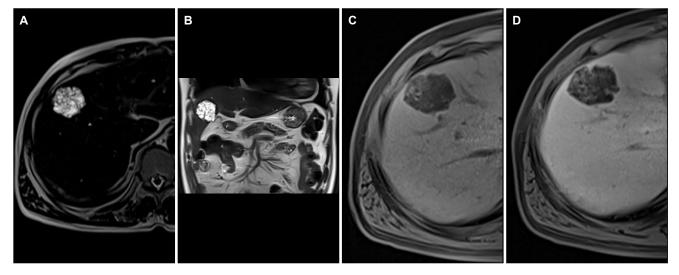


Fig. 1. A 63-year-old man with biliary adenofibroma (case 1). (A) Axial and (B) coronal T2-weighted images showed a bright signal intensity tumor with hypointense septa. (C) Axial unenhanced T1-weighted image showed a low signal intensity tumor with a lobulated border. After gadolinium administration, (D) axial delayed phase T1-weighted image showed septal enhancement.

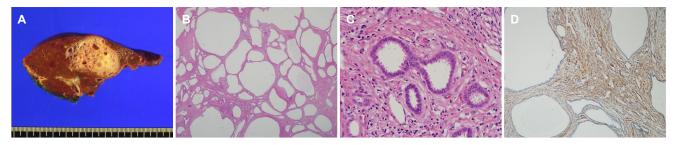


Fig. 2. A 63-year-old man with a biliary adenofibroma (case 1). Gross specimen (A) showed a well-circumscribed multicystic tumor with thin fibrous septa delineating small cysts. (B) The tumor was composed of tubulo-glandular and microcystic structures embedded in the fibrous stroma (H&E, ×40). (C) The tubules and cysts were lined with cuboidal or low columnar biliary epithelial cells. The fibrous stroma contained spindle cells. No ovarian-like stroma was observed (H&E, ×400). (D) Positive immunostaining for α-smooth muscle actin (×100) was observed in the fibrous stroma.

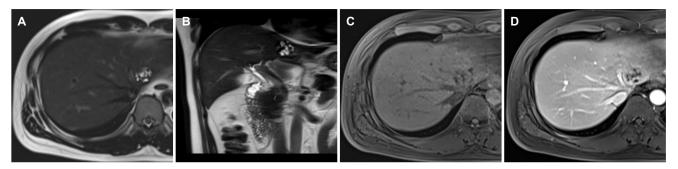


Fig. 3. 38-year-old man with biliary adenofibroma (case 2). (A) Axial and (B) coronal T2-weighted images showed a multilobulated and multiseptated cystic tumor. (C) Axial unenhanced T1-weighted image showed a tumor with low signal intensity. (D) Axial portal venous phase T1-weighted image showed septal and wall enhancement.

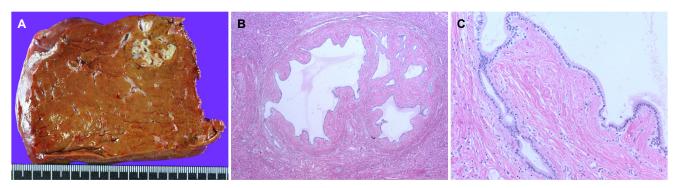


Fig. 4. 38-year-old man with biliary adenofibroma (case 2). Gross specimen (A) showed an unencapsulated tumor with a solid and cystic appearance. (B) Tubulo-glandular structures with fibrous stroma characterized this tumor (H&E, ×40). (C) The tubules were lined with cuboidal or low columnar non-mucin secreting biliary epithelial cells (H&E, ×200).

hepatitis consistent with a HBV infection. The patient has been well with no recurrence after a 41-month follow-up period.

Case 2

A 38-year-old man was referred to the authors' medical institution for an evaluation of a hepatic tumor detected incidentally on a CT scan. A physical examination and laboratory, data including tumor markers, were within the normal range. He denied any personal or family history of hepatobiliary disease.

Abdominal CT revealed a well-circumscribed low attenuation mass within the left lateral section of the liver, measuring 2.7×2.4 cm. MRI revealed a multiseptated multicystic mass with hypointensity on the T1-weighted image and bright signal intensity on the T2-weighted image (Fig. 3A-C). After gadolinium administration, septa and wall enhancement were noted (Fig. 3D). No communication was observed between the lesion and the intrahepatic bile ducts seen on MRI. An

enlarged lymph node was not demonstrated.

The patient underwent a left lateral section ectomy. The gross examination revealed an unencapsulated tumor, measuring 2.5×2.2 cm (Fig. 4A). Microscopically, the tubules were lined with cuboidal or low columnar non-mucin secreting epithelial cells, embedded in a fibrous stroma (Fig. 4B, C). No epithelial atypia was observed. Pathology analysis did not reveal evident communication with the bile ducts. The background liver showed macrovesicular and microvesicular fatty changes. The patient did not develop any recurrence during the 39-month follow-up.

DISCUSSION

Biliary adenofibroma is a rare primary intrahepatic tumor with an unknown etiology. Tsui et al. 1 first described the disease in 1993, and it is now recognized as a distinct tumor of a bile duct origin according to the 2010 World Health Organization (WHO) classification.² Biliary adenofibroma is

Table 1. Clinical and MRI Features of Biliary Adenofibromas Reported in the Literature

% %	Case	Sex/age	Sex/age Tumor size (cm) Tumor location	Tumor location	Symptom	MRI features	Surgery	Follow-up
₽	Varnholt et al. (2003) ⁷	F/47	16	Segment IV, V, VIIII	Right upper quadrant abdominal pain, weight gain	Mass with prominent septa	Subtotal (80%) surgical enucleation	No significant growth of the residual tumor or metastasis after 3 years follow-up
7	Nguyen et al. (2012) ¹⁰	F/53	9.9	Segment IV	No symptom	Multiseptated cystic mass with septal enhancement	Segmental resection of segments III and IV	No recurrence or metastasis after 12 months follow-up
m	Tsutsui et al. $(2014)^{11}$	F/69	3.5	Segment VI	No symptom	T1 hypointense with septa and wall enhancement and T2 hyperintense lesion	Partial liver resection	No recurrence or metastasis after 4 years follow-up
4	Thompson et al. $(2016)^{15}$	M/71	11	Left lobe	No symptom	Multilobulated multiseptated cystic mass	Left hepatectomy	No recurrence or metastasis after 9 years follow-up
Ŋ	Thompson et al. (2016) ¹⁵	M/71	6.3	Caudate lobe	No symptom	with peripheral enhancement	Caudate lobectomy	No recurrence or metastasis after 1 month follow-up
9	Kaminsky et al. (2017) ¹⁶	F/37	4.5	Segment V	Nausea, vomiting, epigastric pain	T1 hypointense and heterogeneously T2 hyperintense lesion	Wedge resection	No recurrence or metastasis after 4 months follow-up
_	Present case 1 (2019)	M/63	4.7	Segment IV and VIII	No symptom	Well-circumscribed multiseptated multicystic tumor T1 hypointensity with septal enhancement	Bisegmentectomy	No recurrence or metastasis after 41 months follow-up
∞	Present case 2 (2019)	M/38	2.7	Segment II	No symptom	and T2 bright signal intensity No bile duct communication	Left lateral sectionectomy	No recurrence or metastasis after 39 months follow-up

MRI, magnetic resonance imaging.

characterized by a complex tubulocystic biliary epithelial tumor without mucin production and with abundant fibroblastic stromal components. To the best of the authors' knowledge, only 21 cases of biliary adenofibroma have been reported thus far in the medical literature. 1,3-19 Previously reported cases focused on the clinical and pathological features of biliary adenofibroma, while the imaging findings were recorded only sporadically. Currently, MRI is the imaging modality of choice for evaluating a range of intrahepatic tumors. MRI has advantages regarding the characterization and diagnosis of these lesions owing to its high contrast resolution. Consequently, the literature was reviewed to characterize the MRI features of biliary adenofibroma.

Table 1 provides brief clinical information and lists the MRI features of the tumor. Biliary adenofibroma occurs in both men and women. The MRI features usually reveal a well-circumscribed multiseptated multicystic tumor that varies in diameter from 2.7 cm to 16 cm. The tumor exhibits hypointensity on the precontrast T1-weighted image and bright signal intensity on the T2-weighted image. After contrast administration, septa and wall enhancement are noted. The present cases also showed no communication with the bile ducts on MRI.

The differential diagnosis for biliary adenofibroma includes intraductal papillary neoplasms of the bile duct (IPNB) and mucinous cystic neoplasm (MCN) of the liver. The tumors grew in a papillary architecture and may be histopathologically similar to biliary adenofibroma. 16 On the other hand, mucin secretion was absent in biliary adenofibroma, whereas IPNB and MCN form mucin-containing cysts. The presence of communication between the lesion and the bile duct system is essential for a diagnosis of IPNB. In contrast to IPNB, biliary adenofibroma has no bile duct communication. MCN requires the presence of ovarian-like stroma and occurs almost exclusively in women. Compared to MCN, biliary adenofibroma does not contain ovarian-like stroma. The MRI findings of IPNB and MCN of the liver are well established. The characteristic MRI features of IPNB are visible intraductal masses within the dilated intrahepatic or extrahepatic bile ducts, and the downstream bile duct can also be dilated.^{20,21} Multifocal papillary tumors are well visualized on the T2-weighted image because of the high contrast between the tumor and background bile.²² The imaging patterns of IPNB can be classified into four subtypes according to the presence of intraductal lesions

and the degree of mucin production: intraductal mass with proximal ductal dilatation, intraductal mass with proximal and distal ductal dilatation, disproportional duct dilatation without a visible mass, and cystic dilatation with a mass.²² Among them, cystic dilatation with a mass manifests as a focal aneurysmal dilatation of the involved duct or a cystic lesion, showing similar imaging features to biliary adenofibroma. A previous study reported that IPNB could manifest the "thread sign", which is defined as intraductal linear or curvilinear hypointense striations on MRI. ²³ On MRI, MCN of the liver typically manifests as a multilocular and septated cystic tumor with an irregular thick wall.²⁴ MRI characterizes intralesional locules with various signal intensities on the T1- and T2-weighted images depending on the cyst fluid protein concentration.²⁵ MCN of the liver does not show dilatation of the downstream bile duct, whereas a ductal dilatation upstream to a cystic lesion can be seen.26

Another differential diagnosis of biliary adenofibroma includes a hepatic echinococcal cyst. Unlike biliary adenofibroma, a typical echinococcal cyst contains multiple daughter cysts. A mother cyst and peripherally located daughter cysts appear as a multilocular cystic lesion with internal septa.²⁷ On MRI, echinococcal cyst shows mixed hypointensity on the T1-weighted image depending on the amount of proteinaceous debris. After contrast administration, the cvst walls and internal septa are enhanced.²⁸

Although biliary adenofibroma is classified as a benign biliary tumor and precursor in the 2019 WHO tumor classification system,²⁹ histopathological and biological spectra of biliary adenofibroma are unclear owing to its rarity. To date, biliary adenofibromas with benign behavior, 1,6-8,13 carcinoma in situ, 10 and malignant transformation 4,5,12,14-16,18 have been reported.

In conclusion, biliary adenofibroma is recognized as a distinct entity of biliary tumor in the WHO classification. The disease is characterized by a tubulocystic bile duct proliferation lined with non-mucin secreting biliary epithelium and supported by abundant fibroblastic stroma. A well-circumscribed multicystic tumor with septal enhancement and no intrahepatic bile duct communication may be the characteristic MRI findings of biliary adenofibroma.

REFERENCES

- 1. Tsui WM, Loo KT, Chow LT, Tse CC. Biliary adenofibroma. A heretofore unrecognized benign biliary tumor of the liver. Am J Surg Pathol 1993;17:186-192.
- 2. Bosman FT, Carneiro F, Hruban RH, Theise ND. WHO classification of tumours of the digestive system. 4th ed. Lyon: IARC
- 3. Parada LA, Bardi G, Hallén M, et al. Monosomy 22 in a case of biliary adenofibroma. Cancer Genet Cytogenet 1997;93:
- 4. Haberal AN, Bilezikci B, Demirhan B, Karakayali H, Haberal M. Malignant transformation of biliary adenofibroma: a case report. Turk J Gastroenterol 2001;12:149-153.
- 5. Akin O, Coskun M. Biliary adenofibroma with malignant transformation and pulmonary metastases: CT findings. AJR Am J Roentgenol 2002;179:280-281.
- 6. Garduño-López AL, Mondragón-Sánchez R, Bernal-Maldonado R, Hinojosa-Becerril CA, Meneses-García A. A case of biliary adenofibroma of the liver causing elevated serum CA 19-9 levels. Rev Oncol 2002;4:271-273.
- 7. Varnholt H, Vauthey JN, Dal Cin P, et al. Biliary adenofibroma: a rare neoplasm of bile duct origin with an indolent behavior. Am J Surg Pathol 2003;27:693-698.
- 8. Gurrera A, Alaggio R, Leone G, Aprile G, Magro G. Biliary adenofibroma of the liver: report of a case and review of the literature. Patholog Res Int 2010;2010:504584.
- 9. Kai K, Yakabe T, Kohya N, et al. A case of unclassified multicystic biliary tumor with biliary adenofibroma features. Pathol Int 2012:62:506-510.
- 10. Nguyen NT, Harring TR, Holley L, Goss JA, O'Mahony CA. Biliary adenofibroma with carcinoma in situ: a rare case report. Case Reports Hepatol 2012;2012:793963.
- 11. Tsutsui A, Bando Y, Sato Y, et al. Biliary adenofibroma with ominous features of imminent malignant changes. Clin J Gastroenterol 2014;7:441-448.
- 12. Thai E, Dalla Valle R, Evaristi F, Silini EM. A case of biliary adenofibroma with malignant transformation. Pathol Res Pract 2016;212:468-470.
- 13. Elpek GÖ, Ünal B, Başsorgun Cİ, Ayik E. A problematic case of unclassified multicystic biliary tumor with adenofibroma features. Turk Patoloji Derg 2016;32:60-62.
- 14. Godambe A, Brunt EM, Fulling KH, Reza Kermanshahi T. Biliary adenofibroma with invasive carcinoma: case report and review of the literature. Case Rep Pathol 2016;2016:8068513.
- 15. Thompson SM, Zendejas-Mummert B, Hartgers ML, et al.

- Malignant transformation of biliary adenofibroma: a rare biliary cystic tumor. J Gastrointest Oncol 2016;7:E107-E112.
- 16. Kaminsky P, Preiss J, Sasatomi E, Gerber DA. Biliary adenofibroma: a rare hepatic lesion with malignant features. Hepatology 2017;65:380-383.
- 17. Arnason T, Borger DR, Corless C, et al. Biliary adenofibroma of liver: morphology, tumor genetics, and outcomes in 6 cases. Am J Surg Pathol 2017;41:499-505.
- 18. Chua D, Chiow AKH, Ang TL, Wang LM. Malignant transformation arising within unusual and rare hepatic lesions: fibropolycystic disease form of ductal plate malformation and biliary adenofibroma. Int J Surg Pathol 2018;26:542-550.
- 19. Esteban M, Amin J, Hertl M, Jakate S, Singh A. Double trouble: a rare case of concurrent biliary adenofibroma and hepatobiliary mucinous cystic neoplasm. ACG Case Rep J 2018;5:e72.
- 20. Joo I, Lee JM, Yoon JH. Imaging diagnosis of intrahepatic and perihilar cholangiocarcinoma: recent advances and challenges. Radiology 2018;288:7-13.
- 21. Lee S, Kim MJ, Kim S, Choi D, Jang KT, Park YN. Intraductal papillary neoplasm of the bile duct: assessment of invasive carcinoma and long-term outcomes using MRI. J Hepatol 2019;70: 692-699.
- 22. Park HJ, Kim SY, Kim HJ, et al. Intraductal papillary neoplasm of the bile duct: clinical, imaging, and pathologic features. AJR Am J Roentgenol 2018;211:67-75.
- 23. Hong GS, Byun JH, Kim JH, et al. Thread sign in biliary intraductal papillary mucinous neoplasm: a novel specific finding for MRI. Eur Radiol 2016;26:3112-3120.
- 24. Soares KC, Arnaoutakis DJ, Kamel I, et al. Cystic neoplasms of the liver: biliary cystadenoma and cystadenocarcinoma. J Am Coll Surg 2014;218:119-128.
- 25. Lewin M, Mourra N, Honigman I, et al. Assessment of MRI and MRCP in diagnosis of biliary cystadenoma and cystadenocarcinoma. Eur Radiol 2006;16:407-413.
- 26. Kim HJ, Yu ES, Byun JH, et al. CT differentiation of mucin-producing cystic neoplasms of the liver from solitary bile duct cysts. AJR Am J Roentgenol 2014;202:83-91.
- 27. Pakala T, Molina M, Wu GY. Hepatic echinococcal cysts: a review. J Clin Transl Hepatol 2016;4:39-46.
- 28. Qian LJ, Zhu J, Zhuang ZG, Xia Q, Liu Q, Xu JR. Spectrum of multilocular cystic hepatic lesions: CT and MR imaging findings with pathologic correlation. Radiographics 2013;33:1419-1433.
- 29. WHO Classification of Tumours Editorial Board. WHO classification of tumours of the digestive system. 5th ed. Lyon: IARC Press, 2019.

Copyright of Korean Journal of Gastroenterology is the property of Korean Society of Gastroenterology and its content may not be copied or emailed to multiple sites or posted to a listsery without the copyright holder's express written permission. However, users may print, download, or email articles for individual use.