



Title	Localized lymphoplasmacytic sclerosing cholecystitis in a patient with autoimmune pancreatitis.
Author(s)	Kawakami, Hiroshi; Eto, Kazunori; Kuwatani, Masaki; Asaka, Masahiro
Citation	Internal medicine, 49(21), 2359-2360 https://doi.org/10.2169/internalmedicine.49.4249
Issue Date	2010
Doc URL	http://hdl.handle.net/2115/47168
Type	article (author version)
File Information	IM49-21_2359-2360.pdf



[Instructions for use](#)

Title page

Localized lymphoplasmacytic sclerosing cholecystitis in a patient with autoimmune pancreatitis

Hiroshi Kawakami,¹; Kazunori Eto,¹; Masaki Kuwatani,¹; Masahiro Asaka,¹

¹ Department of Gastroenterology, Hokkaido University Graduate School of Medicine, Sapporo, Japan

Address correspondence to: Hiroshi Kawakami, MD, PhD

Department of Gastroenterology, Hokkaido University Graduate School of Medicine, Kita 15, Nishi 7, Kita-ku, Sapporo 060-8638, Japan

Tel: +81 11 716 1161 (Ext 5920); Fax: +81 11 706 7867

E-mail: hiropon@med.hokudai.ac.jp (H. Kawakami)

Keywords

Lymphoplasmacytic sclerosing cholecystitis; Autoimmune pancreatitis; Adenomyomatosis

Case presentation

A 55-year-old man was admitted to our hospital for diagnostic evaluation of gallbladder tumors, and elevated serum IgG4 (455 mg/dL). US revealed a mass at the fundus of the gallbladder. A CT scan revealed a well-enhanced gallbladder mass. ERCP showed segmental irregular narrowing of the pancreatic head and stricture of the lower bile duct. The pancreatic lesion was diagnosed as autoimmune pancreatitis (AIP). Diagnostic cholecystectomy was performed. Slicing resected specimen revealed small cysts within the nodule. Histological analysis revealed that the nodule consisted of lymphoplasmacytic infiltration, irregular fibrosis, and numerous IgG4⁺ plasma cells with adenomyomatosis.

Gallbladder lesions with AIP usually showing diffuse thickening of the gallbladder wall (1,2). Interestingly a localized mass was observed in this case. One possible explanation is that the nodule formation resulted from the combined effects of adenomyomatosis and exaggerated sclerosing inflammation around the invaginated epithelium, because IgG4-related inflammation is usually prominent in periductal connective tissue.

References

1. Kamisawa T, Tu Y, Nakajima H, Egawa N, Tsuruta K, Okamoto A, Horiguchi S. Sclerosing cholecystitis associated with autoimmune pancreatitis. *World J Gastroenterol.* 2006;12:3736-3739.
2. Wang WL, Farris AB, Lauwers GY, Deshpande V. Autoimmune pancreatitis-related cholecystitis: a morphologically and immunologically distinctive form of lymphoplasmacytic sclerosing cholecystitis. *Histopathology.* 2009;54:829-836.

Figure legends

Fig. 1 a: Ultrasonography showed a heterogeneously hypoechoic mass at the fundus of the gallbladder.

Fig. 1 b: Contrast-enhanced computed tomography scan revealed a well-enhanced gallbladder mass.

Fig. 2 a: Slicing the resected gallbladder specimen showed small cysts within a whitish nodular mass at the fundus of gallbladder.

Fig. 2 b: Histological analysis revealed that the nodule consisted of numerous IgG4⁺ plasma cells on immunostaining with adenomyomatosis (IgG4 immunostain, x200).

Acknowledgment

The authors would like to thank Dr. Yoh Zen (Institute of Liver Studies, King's College Hospital, London, UK) for his assistance.

Conflict of interest statement

No conflicts of interest exist.



