A Simple Method for Diagnosing Gallbladder Malignant Tumors with Subserosa Invasion by Endoscopic Ultrasonography

Mitsuru Sugimoto (kita335@fmu.ac.jp)  
Fukushima Medical University, School of Medicine  
https://orcid.org/0000-0002-4223-613X

Hiroki Irie  
Fukushima Medical University

Mika Takasumi  
Fukushima Medical University

Minami Hashimoto  
Fukushima Medical University

Yuka Oka  
Fukushima Medical University

Tadayuki Takagi  
Fukushima Medical University

Naoki Konno  
Fukushima Medical University

Rei Suzuki  
Fukushima Medical University

Hiroyuki Asama  
Fukushima Medical University

Yuki Sato  
Fukushima Medical University

Jun Nakamura  
Fukushima Medical University

Tsunetaka Kato  
Fukushima Medical University

Ryoichiro Kobashi  
Fukushima Medical University

Yuko Hashimoto  
Fukushima Medical University

Sigeru Marubashi  
Fukushima Medical University

Takuto Hikichi
Technical advance

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Abstract

**Background:** If the depth of gallbladder malignant tumor (GBMT) invasion is deeper than the subserosa (ss), cholecystectomy is insufficient. In past reports that used endoscopic ultrasonography (EUS) to diagnose the depth of tumor invasion, it was difficult to diagnose GBMT invasion in the ss without a narrow or disrupted lateral hyperechoic layer (LHEL). Therefore, we developed a simple preoperative method to diagnose GBMTs with ss invasion.

**Methods:** Forty-nine GBMT patients who underwent both EUS and surgery were enrolled: 15 patients whose tumors invaded the mucosa (m) or muscularis propria (mp) were classified as the “shallow group”, and 34 patients whose tumors invaded the ss were classified as the “deep group”. The EUS findings were compared between the two groups.

**Results:** An irregular (narrow or thickened) LHEL was significantly more frequently observed on EUS in the deep group than in the shallow group. The diagnosis of ss invasion based on an irregular LHEL had the highest sensitivity and accuracy among the EUS imaging parameters (sensitivity 97.1% (33/34), specificity 86.7% (13/15), accuracy 93.8% (46/49)). When the deep group was limited to patients with a tumor depth of ss, the results were similar. When an irregular LHEL was used, the diagnosability of GBMTs with ss invasion was not significantly different between EUS specialists and beginners.

**Conclusions:** An irregular (thickened or narrow) LHEL observed on EUS could be a reliable and simple method of diagnosing GBMTs with ss invasion and could contribute to choosing an appropriate surgical method.

**Trial Registration:** not applicable

Background

Surgery is the only radical treatment for gallbladder malignant tumors (GBMTs). If the depth of tumor invasion is the mucosa (m) or muscularis propria (mp), cholecystectomy is sufficient. However, if the depth of tumor invasion reaches the subserosa (ss), approximately 50% of patients develop lymph node metastases [1–8]. In such cases, cholecystectomy is insufficient. Therefore, a preoperative diagnosis of GBMTs with ss invasion is important for the selection of the surgical method.

There are several modalities that can be used to closely investigate GBMTs. Transabdominal ultrasonography (US) and computed tomography (CT) are used first. However, the detectability on US is influenced by the physical status of the patients (e.g., subcutaneous fat), and it is difficult to observe gallbladder (GB) lesions in detail on CT. On the other hand, pancreatobiliary lesions can be visualized clearly from the inside of the upper gastrointestinal tract on endoscopic ultrasonography (EUS). Therefore, EUS is preferred over US or CT for the diagnosis of pancreaticobiliary diseases [9, 10]. In fact, many reports on the efficacy of EUS for diagnosing GBMTs have been published [11–23].
Compared to studies that aimed to improve GB cancer diagnosability by EUS, there have been few studies that reported on the diagnosability of the depth of GB invasion by EUS [17, 24–28]. In past reports, tumor depth was predicted by the shape of the wall of the GB on EUS imaging. The GB wall is depicted as a two-layer structure on EUS. These layers are the internal hypoechoic layer and lateral hyperechoic layer (LHEL). The internal hypoechoic layer involves the mucosal muscular layers and the superficial layer of the ss. The LHEL involves the serosal and ss layers [29, 30]. However, the classification of tumor depth based on the combination of tumor shape and the characteristics of the GB wall is slightly complicated, and it is difficult to judge whether the GBMT has invaded the superficial layer of the ss. Some GBMTs that have invaded the superficial layer of the ss do not show a narrowing of the LHEL. Therefore, in this study, we aimed to develop a simple and comprehensible method of diagnosing whether a GBMT has invaded the ss using EUS.

**Methods**

**Study Design and Ethics**

This study was a retrospective study that was performed to develop a simple method for diagnosing ss invasion by GBMTs. This study was approved by the Institutional Review Board of Fukushima Medical University.

**Patients**

Forty-nine GBMT patients who underwent both EUS and surgery between May 2005 and September 2019 at Fukushima Medical University were enrolled in this study. Among them, 15 patients were ultimately diagnosed with a tumor invading the m or mp (m: nine patients, mp: six patients); these patients were classified as the “shallow group”. The remaining 34 patients were ultimately diagnosed with a tumor invading the ss; these patients were classified as the “deep group”. The final diagnosis was confirmed by investigation of surgical specimens.

**Preoperative Diagnosis of GBMTs**

GBMT patients first underwent US or contrast-enhanced CT (CECT). Then, the patients underwent EUS. After sedation with a transvenous administration of midazolam, an echoendoscope was gently inserted into the patients. GBMTs were observed through the antrum of the stomach or the duodenal bulb.

The echoendoscopes used in this study were GF-UMP 230, GF-UM2000, GF-UC240P-AL, GF-UCT240-AL5, GF-UE260-AL5, and GF-UCT260 (Olympus Tokyo, Japan). The EUS systems used in this study were EU-M2000, EU-M30, EU-ME1 and EU-ME2 (Olympus Tokyo, Japan).

All EUS observations were performed by specialists who had performed more than 1000 pancreaticobiliary EUS procedures. No adverse events related to EUS were observed.

**Examination Parameters**
Patient characteristics (age, sex), serum tumor markers (CEA, CA19-9), imaging findings, and histopathological findings (tubular adenocarcinoma or not) were compared between the shallow and deep groups. The following imaging findings were assessed: tumor enhancement on CECT, maximum tumor diameter or height (measured on EUS), form (protruded or wide), internal echo (heterogeneous or homogeneous), and the LHEL observed with EUS. An irregular LHEL was defined as a thickened or narrow LHEL observed on EUS (Fig. 1). When a narrow LHEL is observer, ss invasion is obvious [24, 25, 27]; we also included a thickened LHEL in the examination parameters to achieve better diagnosability of ss invasion. CECT and EUS images were retrospectively reviewed by more than two specialists who were blinded to the histologic depth of the GBMT.

After the most reliable method to diagnose ss invasion was determined, the usability of the method was compared between three pancreaticobiliary EUS beginners and specialists. The pancreaticobiliary EUS specialists were defined as described above. The three EUS beginners were endoscopists who did not fit the definition of a specialist. EUS beginners diagnosed GBMTs with ss invasion by using the most efficient method.

**Statistical Analyses**

Continuous variables that did not follow a normal distribution were analyzed with the Mann-Whitney U test. Nominal variables were analyzed with Fisher’s exact test. P < 0.05 was defined as statistically significant. Bonferroni correction and Holm correction were used for multiple comparisons. All statistical analyses were performed using EzR (Saitama Medical Centre, Jichi Medical University, Saitama, Japan).

**Results**

Age, sex, serum tumor marker levels, and tumor size were not significantly different between the shallow and deep groups (Table 1). Among the imaging findings, enhancement on CECT and internal echo (heterogeneous or homogeneous) were not significantly different between the shallow and deep groups. Wide-shaped tumors, an irregular LHEL, a thickened LHEL, and a narrow LHEL were significantly more common in the deep group than in the shallow group (Table 2). If patients in the deep group were limited to those with a tumor depth of the ss, these four parameters were significantly more frequently observed in the deep group (only the ss) than in the shallow group (Table 3).
Table 1
Comparison of patient characteristics and tumor size

|                         | Shallow group (N = 15) | Deep group (N = 34) | P value |
|-------------------------|------------------------|---------------------|---------|
| Age, years              | 71.0 (55–87)           | 78.5 (51–87)        | 0.071   |
| Sex, male/female        | 6/9                    | 16/18               | 0.76    |
| Serum CEA, ng/ml        | 2.6 (1.1–8.0)          | 2.6 (0.8–889)       | 0.38    |
| Serum CA19-9, U/ml      | 22.1 (2.8–139.4)       | 9.35 (0.1–864.9)    | 0.24    |
| Tumor size, mm          |                        |                     |         |
| Maximum tumor diameter, mm | 20.0 (4.5–40.0)     | 22.0 (10.0–80.0)    | 0.54    |
| Tumor height, mm        | 15.0 (3.0–40.0)        | 15.0 (5.0–50.0)     | 1.0     |

Values are shown as the median (range) or n.
Table 2
Comparison of imaging and histopathological findings

|                         | Shallow group (N = 15) | Deep group (N = 34) | P value |
|-------------------------|------------------------|---------------------|---------|
| Imaging findings        |                        |                     |         |
| Enhancement in CECT     | 14* (100)              | 31* (96.9)          | 1.0     |
| EUS findings            |                        |                     |         |
| Form, protruded/wide    | 8/7                    | 5/29                | 0.011   |
| Internal echo, heterogeneous/homogeneous | 14/1                  | 34/0                | 0.31    |
| Irregular (thickened or narrow) LHEL | 2 (13.3)              | 33 (97.1)          | < 0.01  |
| Thickened LHEL          | 0 (0)                  | 18 (52.9)           | < 0.01  |
| Narrow LHEL             | 2 (13.3)               | 28 (82.3)           | < 0.01  |
| Histopathological findings |                      |                     |         |
| Tumor type              |                        |                     | 0.063   |
| Tubular adenocarcinoma  | 5                      | 22                  |         |
| Other                   | 10                     | 12                  |         |
| Tumor depth             |                        |                     |         |
| m                       | 9                      |                     |         |
| mp                      | 6                      |                     |         |
| Deeper than the ss      |                        | 34                  |         |

Values are shown as the median (range) or n (%) or n.

*Some patients lacked CECT images.

CECT, contrast-enhanced CT; EUS, endoscopic ultrasonography; LHEL, lateral hyperechoic layer; m, mucosa; mp, muscularis propria; ss, subserosa.
Table 3
Comparison of EUS findings in patients whose tumors invaded the m-ss

| EUS findings                        | Shallow group (N = 15) | Deep group (only the ss) (N = 19) | P value |
|-------------------------------------|------------------------|-----------------------------------|---------|
| Form, protruded/wide                | 8/7                    | 3/16                              | 0.03    |
| Irregular (thickened or narrow) LHEL| 2 (13.3)               | 18 (94.7)                         | < 0.01  |
| Thickened LHEL                      | 0 (0)                  | 8 (42.1)                          | < 0.01  |
| Narrow LHEL                         | 2 (13.3)               | 15 (78.9)                         | < 0.01  |

Values are shown as n (%) or n.

m, mucosa; ss, subserosa; EUS, endoscopic ultrasonography; LHEL, lateral hyperechoic layer.

Among the four EUS imaging parameters that were significantly different between the deep and shallow groups, the sensitivity and accuracy for diagnosing ss invasion of an irregular LHEL was the highest (sensitivity 97.1% (33/34), specificity 86.7% (13/15), accuracy 93.8% (46/49)) (Fig. 2a). Regarding the specificity for diagnosing ss invasion, a thickened LHEL was the most specific among the four EUS imaging parameters (100% (15/15)). However, the sensitivity for diagnosing ss invasion of a thickened LHEL was the lowest among the four EUS imaging parameters (52.9% (18/34)). If patients in the deep group were limited to those with a tumor depth of the ss, the sensitivity, specificity, and accuracy were similar (Fig. 2b).

In this study, a thickened LHEL was observed only in the deep group. In patients whose tumors had slightly invaded the ss, edematous thickness of the ss was histologically observed (Fig. 3).

An irregular LHEL was then used as the diagnostic parameter for the ss invasion of GBMTs, and the diagnosability was compared between pancreaticobiliary EUS specialists and beginners. The diagnosability of ss invasion was not significantly different between the specialists and three beginners (endoscopists A, B, and C) (Fig. 4).

**Discussion**

In this study, we developed a simple method that can be used to diagnose the ss invasion of GBMTs on EUS. In past reports, a narrow LHEL was recognized as a sign of ss invasion. In this study, a thickened LHEL was also indicative of ss invasion. As a result, an irregular (thickened or narrow) LHEL was determined to be a simple and reliable EUS finding that could be used to diagnose GBMTs with ss
invasion. An irregular LHEL was also reliable if the patients were limited to those with a tumor depth of m-ss. By using this method, pancreaticobiliary EUS beginners can diagnose GBMTs with ss invasion.

As described in the beginning of this paper, the depth of GB invasion was determined by the combination of tumor form and the structure of the GB wall under the tumor. Regarding the form, the depth of pedunculated tumor invasion is usually the m. However, when wide-shaped lesions are involved, any depth of invasion can be observed [25, 27]. In this study, the specificity for diagnosing ss invasion was low when the form of the tumor was used. Therefore, the form of the tumor was removed from the simple method developed to diagnose ss invasion, while EUS findings of the GB wall were retained.

Some decades ago, it was believed that the internal hypoechoic layer involved the mucosal muscular layers and that the LHEL involved the serosal and subserosal layers [24]. Currently, it is known that the superficial layer of the ss constitutes a part of the internal hypoechoic layer [29, 30]. As described in the introduction, there have been few reports on the efficacy of EUS for diagnosing the depth of GBMT invasion. In these reports, an LHEL that was thinned or disrupted by the tumor indicated tumor invasion deeper than the ss [24, 25, 27]. However, a problem existed in these reports: ss invasion in patients in whom the LHEL was not narrow or disrupted was not examined. In this report, a thickened LHEL was included as a type of irregular LHEL. Our ability to diagnose the ss infiltration of GBMTs was superior to that described in previous reports. In fact, when only a thickened LHEL was observed, the ss that was slightly invaded by the GBMT was observed to be edematous and thickened in pathological specimens (Fig. 3).

It is difficult to diagnose GBMT with invasion into the ss without a narrow or disrupted LHEL, although several attempts have been made. In 2002, Kimura et al. [26] performed EUS and angiography to diagnose ss invasion. When angiography was performed on patients to assess the LHEL, those with no abnormal findings at the cystic artery or its branches were diagnosed as having m or mp GBMTs. On the other hand, patients with abnormal findings at the second or third branches of the cystic artery were diagnosed as having GBMTs with ss invasion. Good diagnosability of GBMTs with ss invasion was reported when these methods were used (sensitivity 81.8%, specificity 90.6%, accuracy 88.4%) [26]. In 2019, Sakamoto et al. [28] reported a scoring system that could be used to diagnose GBMTs with ss invasion: \(-3.954 + 0.555 \text{CEA} + 0.094 \times \text{diameter of the GBMT}\). The sensitivity and specificity using this score for diagnosing ss invasion were 85.0% and 87.1%, respectively (cut-off value – 0.584). Although these methods are efficient for diagnosing GBMTs with ss invasion, angiography is invasive, and the diameter of a wide GBMT is difficult to measure. Therefore, a simple and widely applicable method is desirable. In this report, we developed a simple method using EUS findings of a narrow or thickened LHEL.

This study has some limitations. First, the study design is retrospective, and it was performed at a single institution. Second, EUS is influenced by the technique of the endoscopist. However, EUS was performed by pancreaticobiliary specialists in this study. Therefore, the quality of the EUS image should have been maintained. Third, the evaluation of EUS images is subjective to a certain degree. To overcome this
limitation, the diagnostic method used in this study was also evaluated by pancreaticobiliary EUS beginners. In fact, the efficacy of this method was retained in the beginners.

Conclusions

EUS findings of a narrow or thickened LHEL were observed mainly in patients with GBMTs with ss invasion. This reliable and simple method can be used to diagnose GBMT invasion that is deeper than the ss and may contribute to the appropriate selection of the operative method.

Abbreviations

GBMT: Gallbladder malignant tumor; ss: Subserosa; EUS: Endoscopic ultrasonography; LHEL: Lateral high echoic layer; m: Mucosa; mp: Muscularis propria; US: Ultrasonography; CT: Computed tomography; GB: Gallbladder; CECT: Contrast-enhanced CT.

Declarations

Ethics Approval and Consent to Participate

The study protocol was reviewed and approved by the Institutional Review Board of Fukushima Medical University (Number 2399). The analysis uses anonymous clinical data obtained after all the participants agreed to treatment by written consent. Therefore, patients were not required to give informed consent to the study. The details of the study are published on the homepage of Fukushima Medical University.

Consent for Publication

Not applicable

Availability of Data and Materials

The datasets generated and/or analyzed during the current study are available from the corresponding author upon reasonable request.

Competing Interests

The authors declare that they have no competing interests.

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Department of Gastroenterology, Fukushima Medical University
Authors' Contributions

MS wrote the paper and designed and performed the research; HI, MT and MH designed and oversaw the research; TT, RS, NK, HA, TH, JN, YS, TK and RK provided clinical advice; SM supervised the report; YO, YH performed the pathological diagnoses; and H.O. supervised the report and the writing of the paper. All authors have read and approved the final manuscript.

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Authors' Information

1 Department of Gastroenterology, School of Medicine, Fukushima Medical University, Fukushima, Japan
2 Department of Endoscopy, Fukushima Medical University Hospital, Fukushima, Japan
3 Department of Diagnostic Pathology, School of Medicine, Fukushima Medical University, Fukushima, Japan
4 Department of Hepato-Biliary-Pancreatic and Transplant Surgery, School of Medicine, Fukushima Medical University, Fukushima, Japan

References

1. Chijiiwa K, Nakano K, Ueda J, Noshiro H, Nagai E, Yamaguchi K, et al. Surgical treatment of patients with T2 gallbladder carcinoma invading the subserosal layer. J Am Coll Surg. 2001;192:600-7.
2. Choi SB, Han HJ, Kim CY, Kim WB, Song TJ, Suh SO, et al. Surgical outcomes and prognostic factors for T2 gallbladder cancer following surgical resection. J Gastrointest Surg. 2010;14:668-78.
3. Shimizu Y, Ohtsuka M, Ito H, Kimura F, Shimizu H, Togawa A, et al. Should the extrahepatic bile duct be resected for locally advanced gallbladder cancer? Surgery. 2004;136:1012-7; discussion 8.
4. Shindoh J, de Aretxabala X, Aloia TA, Roa JC, Roa I, Zimmitti G, et al. Tumor location is a strong predictor of tumor progression and survival in T2 gallbladder cancer: an international multicenter study. Ann Surg. 2015;261:733-9.
5. Shirai Y, Yoshida K, Tsukada K, Muto T, Watanabe H. Early carcinoma of the gallbladder. Eur J Surg. 1992;158:545-8.
6. Steinert R, Nestler G, Sagynaliev E, Muller J, Lippert H, Reymond MA. Laparoscopic cholecystectomy and gallbladder cancer. J Surg Oncol. 2006;93:682-9.

7. Suzuki S, Yokoi Y, Kurachi K, Inaba K, Ota S, Azuma M, et al. Appraisal of surgical treatment for pT2 gallbladder carcinomas. World J Surg. 2004;28:160-5.

8. Wakai T, Shirai Y, Yokoyama N, Ajioka Y, Watanabe H, Hatakeyama K. Depth of subserosal invasion predicts long-term survival after resection in patients with T2 gallbladder carcinoma. Ann Surg Oncol. 2003;10:447-54.

9. Tamm EP, Loyer EM, Faria SC, Evans DB, Wolff RA, Charnsangavej C. Retrospective analysis of dual-phase MDCT and follow-up EUS/EUS-FNA in the diagnosis of pancreatic cancer. Abdom Imaging. 2007;32:660-7.

10. DeWitt J, Devereaux B, Chriswell M, McGreevy K, Howard T, Imperiale TF, et al. Comparison of endoscopic ultrasonography and multidetector computed tomography for detecting and staging pancreatic cancer. Ann Intern Med. 2004;141:753-63.

11. Sugiyama M, Xie XY, Atomi Y, Saito M. Differential diagnosis of small polypoid lesions of the gallbladder: the value of endoscopic ultrasonography. Ann Surg. 1999;229:498-504.

12. Sugiyama M, Atomi Y, Yamato T. Endoscopic ultrasonography for differential diagnosis of polypoid gallbladder lesions: analysis in surgical and follow up series. Gut. 2000;46:250-4.

13. Azuma T, Yoshikawa T, Araida T, Takasaki K. Differential diagnosis of polypoid lesions of the gallbladder by endoscopic ultrasonography. Am J Surg. 2001;181:65-70.

14. Kimura K, Fujita N, Noda Y, Kobayashi G, Ito K. Differential diagnosis of large-sized pedunculated polypoid lesions of the gallbladder by endoscopic ultrasonography: a prospective study. J Gastroenterol. 2001;36:619-22.

15. Sadamoto Y, Oda S, Tanaka M, Harada N, Kubo H, Eguchi T, et al. A useful approach to the differential diagnosis of small polypoid lesions of the gallbladder, utilizing an endoscopic ultrasound scoring system. Endoscopy. 2002;34:959-65.

16. Cheon YK, Cho WY, Lee TH, Cho YD, Moon JH, Lee JS, et al. Endoscopic ultrasonography does not differentiate neoplastic from non-neoplastic small gallbladder polyps. World J Gastroenterol. 2009;15:2361-6.

17. Jang JY, Kim SW, Lee SE, Hwang DW, Kim EJ, Lee JY, et al. Differential diagnostic and staging accuracies of high resolution ultrasonography, endoscopic ultrasonography, and multidetector computed tomography for gallbladder polypoid lesions and gallbladder cancer. Ann Surg. 2009;250:943-9.

18. Cha BH, Hwang JH, Lee SH, Kim JE, Cho JY, Kim H, et al. Pre-operative factors that can predict neoplastic polypoid lesions of the gallbladder. World J Gastroenterol. 2011;17:2216-22.

19. Choi JH, Seo DW, Choi JH, Park do H, Lee SS, Lee SK, et al. Utility of contrast-enhanced harmonic EUS in the diagnosis of malignant gallbladder polyps (with videos). Gastrointest Endosc. 2013;78:484-93.
20. Imazu H, Mori N, Kanazawa K, Chiba M, Toyoizumi H, Torisu Y, et al. Contrast-enhanced harmonic endoscopic ultrasonography in the differential diagnosis of gallbladder wall thickening. Dig Dis Sci. 2014;59:1909-16.

21. Sugimoto M, Takagi T, Konno N, Suzuki R, Asama H, Hikichi T, et al. The efficacy of contrast-enhanced harmonic endoscopic ultrasonography in diagnosing gallbladder cancer. Sci Rep. 2016;6:25848.

22. Kamata K, Takenaka M, Kitano M, Omoto S, Miyata T, Minaga K, et al. Contrast-enhanced harmonic endoscopic ultrasonography for differential diagnosis of localized gallbladder lesions. Dig Endosc. 2018;30:98-106.

23. Leem G, Chung MJ, Park JY, Bang S, Song SY, Chung JB, et al. Clinical value of contrast-enhanced harmonic endoscopic ultrasonography in the differential diagnosis of pancreatic and gallbladder masses. Clin Endosc. 2018;51:80-8.

24. Mitake M, Nakazawa S, Naitoh Y, Kimoto E, Tsukamoto Y, Asai T, et al. Endoscopic ultrasonography in diagnosis of the extent of gallbladder carcinoma. Gastrointest Endosc. 1990;36:562-6.

25. Fujita N, Noda Y, Kobayashi G, Kimura K, Yago A. Diagnosis of the depth of invasion of gallbladder carcinoma by EUS. Gastrointest Endosc. 1999;50:659-63.

26. Kimura K, Fujita N, Noda Y, Kobayashi G, Ito K. Diagnosis of pT2 gallbladder cancer by serial examinations with endoscopic ultrasound and angiography. J Gastroenterol. 2002;37:200-3.

27. Sadamoto Y, Kubo H, Harada N, Tanaka M, Eguchi T, Nawata H. Preoperative diagnosis and staging of gallbladder carcinoma by EUS. Gastrointest Endosc. 2003;58:536-41.

28. Sakamoto K, Takai A, Ueno Y, Inoue H, Ogawa K, Takada Y. Scoring system to predict pt2 in gallbladder cancer based on carcinoembryonic antigen and tumor diameter. Scand J Surg. 2019;104:346-52.

29. Fujita N, Noda Y, Kobayashi G, Kimura K, Yago A, Mochizuki F. Analysis of the layer structure of the gallbladder wall delineated by endoscopic ultrasound using the pinning method. Dig Endosc. 1995;7:353-6.

30. Watanabe Y, Goto H, Naitoh Y, Hirooka Y, Itoh A, Taki T, et al. Usefulness of intraductal ultrasonography in gallbladder disease. J Ultrasound Med. 1998;17:33-9.