Synchronous gallbladder metastasis originating from residual gastric cancer: a case report and the review of literature

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Abstract

Background: Gastric cancer rarely metastasizes to the gallbladder. Furthermore, there has never been a case report of simultaneous gallbladder metastasis from residual gastric cancer. Here, we report a case of synchronous gallbladder metastasis originating from a residual gastric cancer.

Case presentation: A 67-year-old man underwent a follow-up upper endoscopy 18 months after a partial gastrectomy for gastric cancer; an ulcerative lesion was found in the remnant stomach at the gastrojejunal anastomosis. A biopsy revealed gastric signet-ring cell carcinoma (SRCC). A full-body examination revealed no abnormalities other than gallstones in the gallbladder. With a diagnosis of residual gastric cancer (cT2N0M0 cStage I), the patient underwent open total gastrectomy and cholecystectomy. Macroscopic findings of the resected specimen revealed thickening of the gallbladder wall; however, no obvious neoplastic lesions were found on the mucosal surface of the gallbladder. The pathological findings showed that the SRCC had invaded the submucosa of the gastrojejunalostomy site with a high degree of lymphatic invasion and lymph node metastases. SRCCs were also found in the lymphatic vessels of the gallbladder wall. These findings suggested the possibility of gallbladder metastasis through lymphatic vessels. The patient and his family members refused postoperative chemotherapy. Ten months after the operation, the patient experienced respiratory failure due to lymphangitis carcinomatosa and died.

Conclusions: At present, it is difficult to determine whether resection of the gallbladder contributes to an improved prognosis of gastric cancer patients. However, reports in such cases demonstrate that gallbladder metastasis could be a poor predictor of prognosis for gastric cancer.

Keywords: Residual gastric cancer, Metastasis, Signet-ring cell carcinoma, Gallbladder

Background

Gallbladder metastasis from gastric cancer is extremely rare [1]. A previous report showed that the occurrence rate of gallbladder metastasis from gastric cancer is only 0.06% [2]. In addition, the preoperative diagnosis of gallbladder metastasis from gastric cancer is difficult due to few reported cases and a lack of specific imaging findings. Furthermore, imaging findings may also overlap with features of other gallbladder diseases, such as cholecystitis, gallbladder polyps, and adenomyomatosis of the gallbladder [3].

The prognosis of gallbladder metastasis is poor. Yoon et al. [4] reported that the median survival time for metastatic gallbladder tumors 8.7 months. Here, we present a case of synchronous gallbladder metastasis originating from a residual gastric SRCC in addition to a review of the literature.

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Case presentation
A 67-year-old man was diagnosed with gastric cancer via an upper endoscopy for the evaluation of esophageal varices while commuting to our hospital for alcoholic cirrhosis (Child–Pugh classification: grade B). The patient tested positive for *Helicobacter pylori* infection. He had a history of drinking >108 g of alcohol per day for 40 years (stopped 5 years ago) and smoking 40 cigarettes per day for 40 years (stopped 8 years ago). Due to impaired liver function and the presence of esophageal varices, the surgical risk was high, and partial gastrectomy (D0 dissection, with antecolic Roux-en-Y reconstruction) was performed with preservation of the left gastric artery. The postoperative pathological diagnosis was LMD, Gre, Type 3, 90 × 55 mm, poorly differentiated adenocarcinoma, non-solid type, pT2 (MP), INFc, Ly0, V0, pPM0 (15 mm), pDM0 (3 mm), and pN0 (#3:0/9, #4d:0/2) according to the Japanese Classification of Gastric Carcinoma [5] and T2N0 Stage IB in accordance with the 8th edition of the Union for International Cancer Control classification[6], with no signet-ring cell (Fig. 1).

The patient underwent a follow-up upper endoscopy 18 months after gastric cancer surgery. A 0–IIc lesion was found at the gastrojejunal anastomosis; a biopsy of the same site revealed SRCC (Fig. 2). Blood chemistry tests showed that liver function improved with abstinence (Child–Pugh classification: grade A), and other blood count and biochemistry parameter levels were within normal ranges. Levels of tumor markers, carcinoembryonic antigen (2.2 ng/ml), and carbohydrate antigen 19–9 (11.6 U/ml) were within normal limits.

Abdominal computed tomography (CT) showed difficulty in identifying the primary lesion, and there were no findings suggestive of lymph node (LN) metastasis or distant metastasis. There were no abnormal findings in the gallbladder wall, except for the presence of gallbladder stones, with no changes observed for 2 years (Fig. 3a, b). Fluoro-deoxyglucose (FDG)-positron emission tomography (PET)/CT revealed negative uptake in the gastric lesion (Fig. 3c).

Surgical findings showed no distant metastasis, such as peritoneal dissemination or liver metastasis, or obvious LN metastasis. In addition, the intraoperative washing cytology was negative. There was also no obvious exposure of the serous surface to residual gastric cancer. In the gallbladder, the wall of the body appeared thickened; however, there were no obvious neoplastic lesions. Therefore, we performed a total gastrectomy with LN dissection and cholecystectomy to prevent cholecystitis.
As for the macroscopic findings, a thickening of the mucosa on the gastric side of the previous gastrojejunostomy anastomosis from the previous operation was observed, and the main lesion was \(75 \times 55\) mm (Fig. 4a). Histopathologically, the stomach showed an SRCC and a non-seminomatous poorly differentiated adenocarcinoma in the mucosa (Fig. 4b, c). The cancer cells showed strong lymphatic invasion and had invaded the layer of tissue under the serosa. The LNs showed metastasis in No. 1, 3, and 4sa (11/14).

The gallbladder showed partial wall thickening, suggestive of adenomyosis (Fig. 5a). Histological examination of the gallbladder revealed no mucosal epithelial lesions. The lymphatic vessels of the gallbladder wall were extensively filled with tumor cells, especially in the grossly thickened areas (Fig. 5b–d). Since the histological findings of the gastric and gallbladder lesions were identical, the gallbladder lesion was diagnosed as gallbladder metastasis of the gastric cancer. The final pathological stage was T3N3aM1 (GAL) stage IV.

The patient was discharged on the 11th postoperative day in a good condition. The patient and his family members refused postoperative chemotherapy. There was no metastasis or recurrence other than lymphangitis carcinomatosa, and the patient died of respiratory failure 10 months postoperatively.
Discussion
In this case, gallbladder metastasis was incidentally discovered 18 months after residual gastric cancer surgery. Regarding the gastric lesions, the initial lesions had no lymphatic invasion (Ly0) or lymph node metastasis (#3:0/9, #4d:0/2); thus, lymphatic invasion was different from that of the recurrent lesions. SRCCs tend to spread laterally across the normal mucosal surface [7] and gastric cancer is highly heterogeneous [8]. In this case, it is possible that a new type of SRCC may have developed; however, considering that the extent of the tumor was difficult to identify visually, and that the proximal margin of the first surgery was short (15 mm), the possibility that cancer cells may have remained histologically in an area that could not be confirmed macroscopically and that remaining tumor cells with high lymphatic invasion potential had recurred and proliferated was considered.

Gallbladder metastases are rare [1]. Chan et al. [2] reported 36 cases of gallbladder metastasis (0.46%) out of 7,910 excised gallbladders in patients with malignant tumors, of which only five cases (0.06%) originated from gastric cancer. To the best of our knowledge, only 18 cases of gallbladder metastasis from gastric cancer have been reported from 1990 to 2020 in Japan, in addition to our own cases. Furthermore, gallbladder metastasis occurring in residual gastric cancer has not been reported in the literature and is considered extremely rare. Seventeen cases of the available pathological type had poorly differentiated histology, such as por or sig (70%). Eleven of 16 cases (69%) with available pathological data showed lymphatic invasion (Ly ≥ 2) and LN metastasis (N ≥ 2). In addition, macroscopic findings of the gallbladder often include submucosal masses and thickened walls. In only one of 17 cases (6%), gallbladder metastasis was clinically diagnosed preoperatively, indicating that preoperative diagnosis is difficult. In addition, 10 of 19 patients (53%) died within one year of surgery (Table 1).

Choi et al. [3] classified the CT findings of metastatic gallbladder tumors into wall thickening and protruding lesions into the lumen, and identified specific growth patterns based on the type of primary tumor. They stated that metastatic gallbladder adenocarcinoma was often observed to have invasive wall thickening lesions, while the polyp type was mainly associated with non-adenocarcinomatous histological types. However, these findings have also been observed in primary gallbladder tumors. In addition, the findings of gallbladder metastasis are diverse, and the findings on CT may overlap with those of other gallbladder diseases [9], thus preoperative diagnosis is difficult.

Regarding the route of metastasis of gastric cancer to the gallbladder, according to previous reports, the presence of advanced LN metastasis and lymphatic vessel invasion of tumor cells caused obstruction of various lymphatic vessels and the development of abnormal lymphatic pathways, leading to lymphatic metastasis to the
Table 1  Reported cases of gallbladder metastasis from gastric cancer in Japan

| Case | Year | Age | Sex | Gastric cancer | Histological type | Preoperative diagnosis | Macroscopic findings | Metastatic pathway | Cause of death | Duration/Prognosis |
|------|------|-----|-----|----------------|-------------------|------------------------|---------------------|------------------|----------------|--------------------|
| 1    | 1990 | 44  | M   | Por            | 2                 | GB polyps             | SMT                 | L                | Carcinomatous meningitis | 8 months/Death |
| 2    | 1996 | 54  | M   | ND             | ND                | ND                     | ND                  | ND               | ND              | ND                 |
| 3    | 1996 | 63  | F   | ND             | ND                | ND                     | ND                  | ND               | ND              | ND                 |
| 4    | 1999 | 62  | M   | Sig            | 2                 | GB stones             | –                   | ND               | DIC            | 4.5 months/Death |
| 5    | 2002 | 86  | F   | Adenosquamous  | 3                 | WT                     | WT                  | L                | Acute renal failure | 30 days/Death  |
| 6    | 2006 | 88  | M   | Endocrine      | 3                 | NA                     | –                   | L or H           | 8 months/Alive  |
| 7    | 2007 | 54  | M   | Por            | 2                 | GB stones             | WT                  | L                | Obstructive jaundice Carcinomatous peritonitis | 5 months/Death |
| 8    | 2008 | 71  | M   | Tub2           | 1                 | GB stones             | –                   | L                | Carcinomatous peritonitis Pneumonia | 53 days/Death  |
| 9    | 2008 | 76  | M   | Por            | 2                 | NA                     | –                   | H                | Bone metastasis Weakness | 43 days/Death  |
| 10   | 2009 | 54  | M   | Por            | 2                 | GB stones             | SMT                 | L                | 3 years/Alive   |
| 11   | 2011 | 66  | F   | Por            | 1                 | NA                     | –                   | L                | 13 months/Alive |
| 12   | 2013 | 50  | F   | Tub2           | 3                 | Primary GB tumor      | WT                  | L                | 33 months/Alive |
| 13   | 2014 | 68  | F   | Sig            | 3                 | Primary GB tumor      | WT                  | L                | 24 months/Alive |
| 14   | 2014 | 62  | F   | Por            | 3                 | NA                     | SMT                 | L                | Recurrence of cancer | 9 months/Death |
| 15   | 2014 | 55  | M   | Tub2           | 2                 | GB metastasis         | SMT                 | L or H           | ND              |
| 16   | 2019 | 66  | F   | Por+tub2       | 1                 | GB stones Adenomyomatosis | SMT                 | L                | Lymphangitis carcino- matosis | 7 months/Death |
| 17   | 2020 | 71  | M   | Sig            | 0                 | GB stones             | –                   | L                | Death of cancer | 199 days/Death  |
| 18   | 2021 | 84  | M   | Por            | ND                | Acute cholecystitis   | WT                  | Peritoneal dissemina- tion | ND/Alive    |
| Present case | 2020 | 67  | M   | Sig            | 1                 | GB stones             | WT                 | L                | Lymphangitis carcino- matosis | 10 months/Death |

ND no data, por poorly differentiated adenocarcinoma, sig signet-ring cell carcinoma, tub2 moderately differentiated tubular adenocarcinoma, NA no abnormality, GB gallbladder, WT wall thickness, SMT submucosal tumor, L lymphogenous, H hematogenous
stroma within the gallbladder wall [10]. The lymphatic flow from residual gastric cancer, such as in our case, develops a different flow from that of the normal stomach depending on the initial surgical technique, degree of LN dissection, and degree of adhesions after surgery [11]. In residual gastric cancer after pyloric gastrectomy without dissection of the left gastric artery, lymphatic flow in the upper part of the gastric body is reported to follow the following pathways: from the cardia, right side of the gastric fornix, and lesser curvature of the gastric body to the left gastric artery; from the left side of the gastric fornix and greater curvature of the gastric body to the left gastroepiploic artery and short gastric artery to the splenic artery; from the gastric fornix and posterior wall of the gastric body to the splenic artery via the posterior gastric artery; and from the cardia to the left diaphragmatic artery [11]. The lymphatic flow at the anastomosis depends on the reconstruction method, and it is thought that the adhesion of the remaining stomach to the surrounding tissues leads to the formation of new lymphatic channels, which can easily metastasize to mesenteric LNs adjacent to the anastomosis and the root of the superior mesenteric vein (No. 14) in cases of residual gastric jejunal anastomosis such as Roux-en-Y, which is high in cancer [12].

In our case, although the gastric lesion showed invasion into the subserosa, the possibility of disseminated metastasis or direct invasion to the gallbladder was considered to be low. This was because there was no continuity between the gastric and gallbladder lesions, no change in the serous surface of the gallbladder, and the intraoperative washing cytology was negative. In addition, in our case, the gastric lesion was negative for venous invasion (V0) and showed severe lymphatic invasion (Ly1) and LN metastasis (N3). The gallbladder lesion also showed a high degree of lymphatic invasion, and the histology was identical to that of the gastric lesion. Gastrojejunostomy cases are said to easily metastasize to the periaortic LN due to the aforementioned characteristics of lymphatic flow [12]. Since there were no grossly obvious tumor lesions in the falciform ligament or other parts of the liver, spleen, and pancreas in this case, extended LN dissection was not performed; therefore, the actual extent of LN metastasis is unknown. However, since metastasis was found in dissected LN 1, 3, and 4sa, it was inferred that cancer cells flowed from the left gastric artery to the celiac artery, and from the left gastroepiploic artery and posterior gastric artery to the celiac artery through the splenic artery. It was speculated that lymphatic vessel obstruction or narrowing in

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**Fig. 6** The speculated pathway of metastasis from gastric cancer to gallbladder. The black circled area is blockage or stenosis site in the lymphatic vessels. Imaginary abnormal lymphatic backflow shown with the dotted pathway.
the downstream region caused lymphatic reflux into the common hepatic artery and right hepatic artery, which metastasized lymphatically to the gallbladder (Fig. 6).

The prognosis of patients with gallbladder metastasis from gastric cancer, including our patient, is extremely poor, with a median survival time of 7 months. To establish a treatment strategy in such cases, a detailed case analysis is required.

Conclusions
In this study, a review of the literature revealed that gallbladder metastasis from gastric cancer has poorly differentiated histology (por/sig) in 70% of cases, lymphatic invasion with Ly ≥ 2, and LN metastasis with N ≥ 2 in 69% of cases. Preoperative diagnosis was difficult; only one of the 17 reported patients had a correct diagnosis before surgery. In addition, 10 of 16 patients died within 1 year after surgery. At present, it is difficult to determine whether resection of the gallbladder contributes to improved gastric cancer prognosis; however, the diagnosis of gallbladder metastasis may be a poor predictor of prognosis for gastric cancer.

Abbreviations
SRCC: Signet-ring cell carcinoma; LN: Lymph node; CT: Computed tomography; FDG: Fluoro-deoxy-glucose; PET: Positron emission tomography.

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Author contributions
AK collected the associated data and edited the manuscript. TN supervised the writing of the manuscript. TI, KH, and KM performed the procedure on this patient. DT, KK, HY, YK, and RM participated in the treatment. YO contributed to pathological diagnosis. All authors read and approved the final manuscript.

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Availability of data and materials
Data sharing is not applicable to this article, as no datasets were generated or analyzed during the current study.

Declarations

Ethics approval and consent to participate
This study was conducted in accordance with the ethical standards of the Japan Surgical Society.

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 no competing interests.

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References
1. Muta Y, Mochiki E, Kumakura M, Toyomasu Y, Ishiguro T, Kumagai Y, et al. A case of synchronous gallbladder metastasis originating from advanced gastric cancer. Gan To Kagaku Ryoho. 2020;47:376–8.
2. Chan KW. Review of 253 cases of significant pathology in 7,910 cholecystectomies in Hong Kong. Pathology. 1988;20:20–3. https://doi.org/10.3109/00313028809885191.
3. Choi WS, Kim SH, Lee ES, Lee KB, Yoon WJ, Shin CI, et al. CT findings of gallbladder metastases: emphasis on differences according to primary tumors. Korean J Radiol. 2014;15:334–45. https://doi.org/10.3348/kjr.2014.15.3.334.
4. Yoon WJ, Yoon YB, Kim YJ, Ryu JK, Kim YT. Metastasis to the gallbladder: a single-center experience of 20 cases in South Korea. World J Gastroenterol. 2009;15:4806–9. https://doi.org/10.3748/wjg.15.4806.
5. Japanese Gastric Cancer Association. Japanese classification of gastric carcinoma. 15th ed. Kanehara. Japanese Gastric Cancer Association; 2017.
6. Brierley J, Gospodarowicz M, Wittekind C. TNM classification of malignant tumours. 8th ed. New Jersey: Wiley-Blackwell; 2016.
7. Sugihara H, Hattori T, Fukuda M, Fujita S. Cell proliferation and differentiation in intramucosal and advanced signet ring cell carcinomas of the human stomach. Virchows Arch A Pathol Anat Histopathol. 1987;411:117–27. https://doi.org/10.1007/BF00712735.
8. Ho SWT, Tan P. Dissection of gastric cancer heterogeneity for precision oncology. Cancer Sci. 2019;110:3405–14. https://doi.org/10.1111/cas.14191.
9. Cocco G, Pozzi AD, Basilico R, Fabiani S, Taraschi AL, Pascucci L, et al. Imaging of gallbladder metastasis. Insights Imaging. 2021;12(1):1–8.
10. Matsuda I, Yamazaki M, Taniguchi M, Kawaiuchi M, Furukawa K, Terada T. A resected case of adenosquamous cell carcinoma of the stomach with gallbladder metastasis. J Jpn Surg Assoc. 2002;63:2688–92.
11. Uebara H, Yabusaki H, Matsuki A, Atsawat M, Bamba T, Matsuyama S, et al. Investigation focusing on metastatic status and dissection effect of lymph nodes in remnant gastric cancer following distal gastrectomy for gastric cancer. Jpn J Gastroenterol Surg. 2018;51:1–11. https://doi.org/10.5833/jgss.2017.0029.
12. Kato T, Ninomiya M, Choda Y, Kanazawa T, Harano M, Matsukawa H, et al. Prognostic analysis of remnant gastric cancers with metastatic lymph nodes. Jpn J Gastroenterol Surg. 2015;48:963–70. https://doi.org/10.5833/jjgs.2014.0134.

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