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**Breast cancer metastasizing to the stomach mimicking primary gastric cancer: A case report**

Yim K *et al.* Breast cancer with stomach metastasis

Kwangil Yim, Sang Mi Ro, Jieun Lee

**Kwangil Yim,** Department of Hospital Pathology, Seoul St. Mary’s Hospital, College of Medicine, The Catholic University of Korea, Seoul 06591, South Korea

**Sang Mi Ro, Jieun Lee,** Division of Medical Oncology, Department of Internal Medicine, Seoul St. Mary’s Hospital, College of Medicine, The Catholic University of Korea, Seoul 06591, South Korea

**Jieun Lee,** Cancer Research Institute, the Catholic University of Korea, Seoul 06591, South Korea

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**Correspondence to: Jieun Lee, MD, PhD,** Division of Medical Oncology, Department of Internal Medicine, College of Medicine, The Catholic University of Korea, Seocho-gu, Seoul 06591, South Korea. befamiliar@catholic.ac.kr

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**Abstract**

Breast cancer with stomach metastasis rare with an incidence of 1% or less among metastatic breast cancer patients. We experienced a case of breast cancer metastasizing to the stomach in 65-year-old female patient. She experienced dyspepsia and poor oral intake before visiting the clinic. Diffuse infiltration with nodular mucosal thickening of the stomach wall was observed, suggesting advanced gastric cancer based on gross endoscopic finding. Spread of poorly cohesive tumor cells in the gastric mucosa observed upon hematoxylin and eosin stain resembled signet ring cell carcinoma, but diffuse positive staining for GATA3 in immunohistochemical stain allowed for a conclusive diagnosis of breast cancer metastasizing to the stomach. Based on the final diagnosis, systemic chemotherapy was administered instead of primary surgical resection. After 2 cycles of docetaxel administration, she showed a partial response based on abdominal computed tomography scan. This case is an unusual presentation of breast cancer metastasizing to the gastrointestinal tract.

**Key words:** Gastric cancer; Breast cancer; Metastasis; Immunohistochemical stain; GATA3; GCDFP-15

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**Core tip:** This case report describes a patient who was clinically diagnosed as advanced gastric cancer, but final pathological confirm diagnosis was to be breast cancer with gastric metastasis. Patient received systemic chemotherapy and is currently on partial response state at present.

Yim K, Ro SM, Lee J. Breast cancer metastasizing to the stomach mimicking primary gastric cancer: A case report. *World J Gastroenterol* 2017; In press

**INTRODUCTION**

Breast cancer commonly metastasizes to bone, lung, liver, and brain, but metastasis to the gastrointestinal tract is rare[1,2]. In Korea, fewer than 10 cases of breast cancer metastasizing to the gastrointestinal tract have been reported[3]. Breast cancer with gastrointestinal metastasis requires systemic chemotherapy. However, if breast cancer with gastrointestinal metastasis is misdiagnosed as a primary gastrointestinal cancer, unnecessary surgical resection may take over place. Herein, the authors present a case of breast cancer metastasizing to the stomach, initially suspected to be primary gastric cancer. This patient was successfully treated with systemic chemotherapy.

**CASE REPORT**

A 65-year-old female patient was referred to the oncology department for evaluation of indigestion and epigastric discomfort. She had been previously diagnosed with breast cancer, treated with modified radical mastectomy (invasive lobular carcinoma, pT2N3M0), adjuvant chemotherapy (cyclophosphamide, methotrexate, 5-FU) and adjuvant radiation. Two years after surgery, she experienced cancer recurrence with bone metastasis and received an aromatase inhibitor (letrozole) as treatment for another 2 years. At the time she visited the oncology department, she was currently on aromatase inhibitor (letrozole). Other than breast cancer, she had no other medical history. Her last endoscopy was performed 2 years ago, with no specific findings.

Initial white blood cell (WBC) counts, hemoglobin level and hematocrit were 4790 cell/mm3 (neutrophil count 82%, lymphocytes count 25.8%), 13.1 g/dL (normal range 13.0-18.0 g/dL), and 369000/mm3 (normal range 150000-450000/mm3). Other laboratory findings including those of blood chemistry and urine analysis were in the normal range. Serum carcinoembryonic antigen level was increased up to 23.25 ng/dL.

Endoscopy revealed diffuse infiltration with nodular mucosal thickening of the stomach wall, involving the lower two-thirds of the stomach body (Figure 1). Based on endoscopy, endoscopic ultrasound (Figure 2A) and abdominal CT scan (Figure 2B), advanced gastric cancer (cT3N1M0) was suspected. Hematoxylin and eosin (H&E) staining of the endoscopic biopsy revealed poorly cohesive tumor cells spreading into the gastric mucosa, suggesting signet ring cell carcinoma. However, no intracytoplasmic mucin was found in the tumor cells, with scant to moderate pinkish cytoplasm. Normal stomach glandular tissue was found in the biopsy specimen, with no cancer cells connected to the glandular structure (Figure 3A and B). These findings were not consistent with typical gastric signet ring cell carcinoma. Because the patient was diagnosed with invasive lobular carcinoma, archival breast tumor tissue was re-evaluated for comparison.

Breast tissue pathology showed a similar appearance to the endoscopic biopsy specimen, such as a de-cohesive pattern with cells arranged in an Indian file pattern, and a centrally located enlarged nucleus (Figure 3C). In the immunohistochemical (IHC) test, the tumor cells showed diffuse strong nuclear staining for GATA3 binding protein (GATA3) (Figure 3D). IHC results of gross cystic disease fluid protein-15 (GCDFP-15) (Figure 3E), E-cadherin (Figure 3F), estrogen receptor (ER, Figure 3G) and progesterone receptor (PR, Figure 3H) were negative. HER-2 IHC staining showed weak membranous staining consistent with equivocal (+2) positivity (Figure 3I). Silver in situ hybridization (SISH) for HER-2 gene was performed, and the dual-probe HER2/Chr17 ratio was 3.2 (161/51), consistent with HER-2 amplification (Figure 3J).

Based on the pathologic findings, breast cancer metastasizing to the stomach was diagnosed. The stomach metastasis developed 4 years after surgery and 2 years after the initiation of an aromatase inhibitor use. As systemic treatment, docetaxel combined with trastuzumab was considered but trastuzumab was not available due to insurance guidelines. Docetaxel (150 mg/m2 intravenously [I.V.], day 1) was administered every 3 wk. After 2 cycles of systemic chemotherapy, follow up abdominal CT scans showed decreased stomach wall thickness, and perigastric lymph nodes showed a partial response (PR) based on the Response Evaluation Criteria in Solid Tumors (RECIST) (Figure 4). During 2 cycles of systemic chemotherapy, the patient’s symptoms of indigestion and epigastric discomfort regressed. Currently, the patient is in persistent PR state and 6 cycles of docetaxel have been administered.

**DISCUSSION**

Cancer metastasizing to the gastrointestinal (GI) tract is reported to be rare, but breast cancer is the second most common cancer metastasizing to the GI tract after lung cancer[2,4]. However, the incidence of breast cancer with GI tract metastasis is reported to be 1% or lower[5,6]. Invasive lobular carcinoma tends to metastasize to the GI tract more frequently compared to invasive ductal carcinoma[7]. The most common metastatic sites in the GI tract are the colon and rectum, stomach, small intestine and esophagus, in that order[4]. In Korea, 7 cases of breast cancers metastasizing to the GI tract have been reported, with 5 cases of breast cancer with gastric metastases and 2 cases of synchronous stomach and colorectal metastases[3]. The clinical characteristics of the previous cited cases are summarized in Table 1[2,3,8-19].

Most breast cancer patients with gastric metastasis present with GI symptoms[3,16], similar to primary gastric cancer. In our case, the patient complained of indigestion, early satiety, and weight loss. Endoscopy with sufficient mucosal biopsy is mandatory for the diagnosis. Diffuse infiltration of the gastric wall with linitis plastica formation may be found[2], but approximately 50% of patients may have shallow mucosal lesion indistinguishable from benign gastric mucosal lesions[20]. Our patient showed extensive nodular mucosal thickening with a thickened gastric fold, with a primary suspicion of advanced gastric cancer.

Pathologic findings of breast cancer metastasizing to the stomach are morphologically similar to poorly cohesive gastric carcinoma, especially in invasive lobular carcinoma[21,22]. However, some morphological differences are present. In metastatic mammary carcinoma, sialomucin is present in the intracytoplasmic lumina with a central location of the nucleus. In contrast, primary gastric signet ring cell carcinoma contains clear intracytoplasmic acid mucin that pushes the nucleus to the periphery[21]. In the present case, the nuclei of the tumor cells were located in the center, and there were no clear intracytoplasmic inclusions.

Also, IHC study is helpful for differential diagnosis. Gross cystic disease fluid protein 15 (GCDFP-15) staining was traditionally used for differential diagnosis of mammary origin carcinoma. However, it shows relatively low sensitivity (55%-76%) for detecting a breast origin cancer[23]. Recently, GATA3 is widely known as a mammary cancer and urothelial cancer marker. GATA3 expression shows 100% positivity in involving breast lobular carcinoma and 96% positivity in breast ductal carcinoma. However, only 5% of tumors are positive for GATA3 in gastric adenocarcinoma[24]. In our case, although GCDFP-15 staining was negative, GATA3 showed diffuse strong nuclear positivity, consistent with a mammary origin of the carcinoma.

Metastatic breast cancer involving the stomach is treated with systemic agents such as cytotoxic chemotherapeutic agents or hormonal agents. Surgical resection of the stomach has a limited role in treatment, and does not affect the survival outcomes of patients presenting with gastric metastasis[4]. However, surgical treatment may have a role in palliative treatment such as relieving obstructive symptoms.

Breast cancer patients have a superior survival outcome compared to other cancers, raising the possibility of a double primary cancer during the clinical course. However, metastasis of primary breast cancer must also be considered. In a breast cancer patient who complains of gastrointestinal symptoms, prompt endoscopy and biopsy are necessary for an accurate diagnosis. Sufficient pathologic review of gastric biopsy and previous breast specimens, with immunohistochemical examination is warranted. When metastasis of breast cancer to the stomach is suspected, appropriate systemic treatment is necessary for further treatment.

**COMMENTS**

***Case characteristics***

A 65-year-old female patient who was diagnosed as metastatic breast cancer visited the hospital for evaluation of epigastric discomfort.

***Clinical diagnosis***

Epigastric discomfort and indigestion.

***Differential diagnosis***

Gastric ulcer, primary gastric cancer showed be differentiated by endoscopic biopsy.

***Laboratory diagnosis***

Serum carcinoembryonic antigen was increased up to 23.25 ng/dL.

***Imaging diagnosis***

Endoscopy showed diffuse infiltration with nodular mucosal thickening of stomach wall.

***Pathological diagnosis***

Metastatic invasive lobular carcinoma to stomach was diagnosed by immunohistochemical stain.

***Treatment***

Docetaxel 150 mg/m2 intravenous, every 3 wk.

***Related reports***

Breast cancer rarely metastasize to gastrointestinal tract and should be diagnosed by careful review of the pathologic specimen. If patient have underlying breast cancer, metastatic breast cancer should be considered other than primary gastric cancer during the diagnosis.

***Term explanation***

GATA3 refers to GATA3 binding protein used for differential marker for diagnosis of breast cancer. Partial response (PR) means more than 30% decrease in the sum of the longest diameters of target lesions during response evaluation.

***Experiences and lessons***

Early differential diagnosis of metastatic breast cancer to stomach is important for appropriate systemic chemotherapy and avoidance of unnecessary surgery.

***Peer-review***

This is generally an interesting and useful paper.

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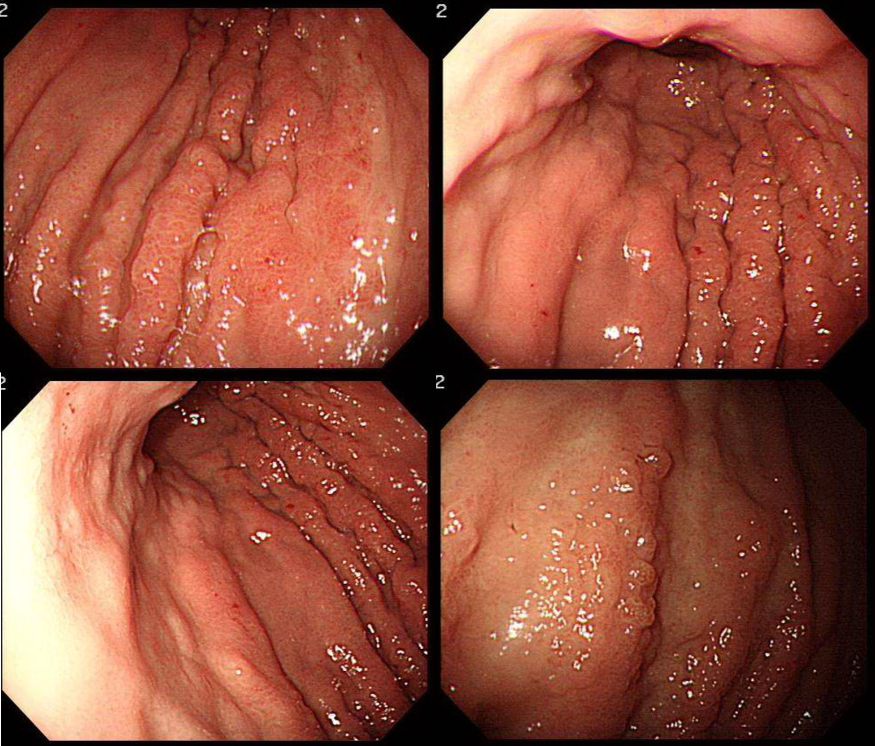
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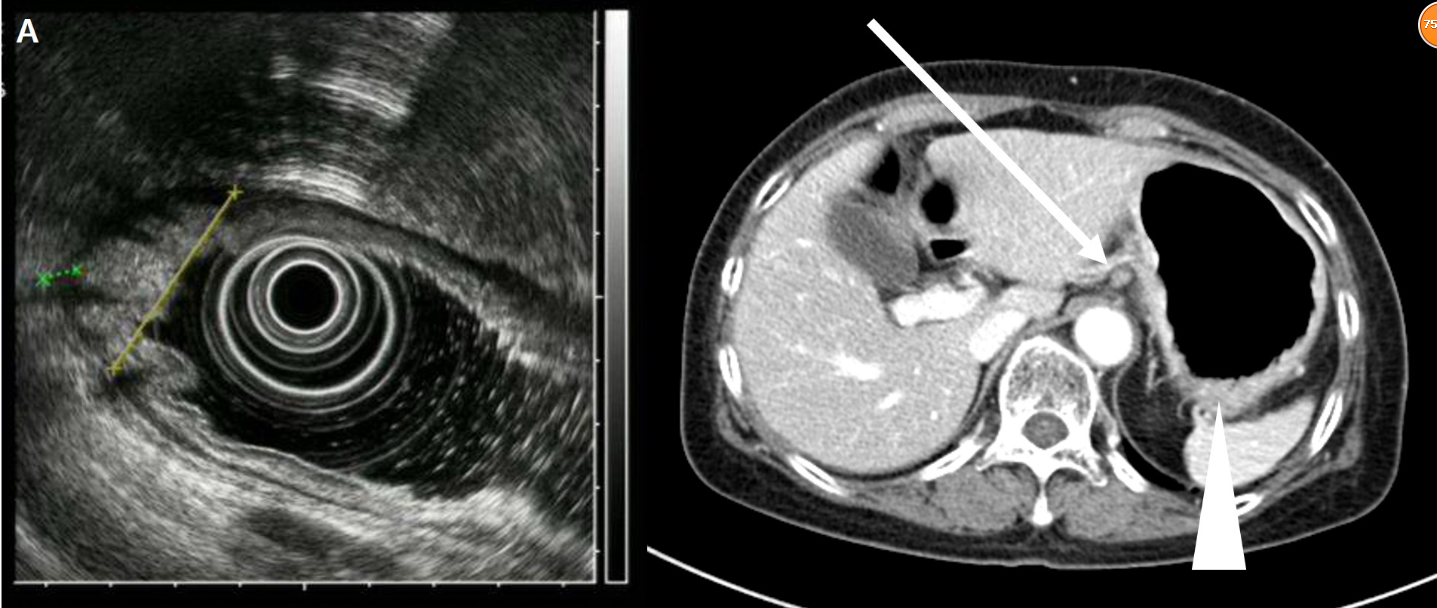
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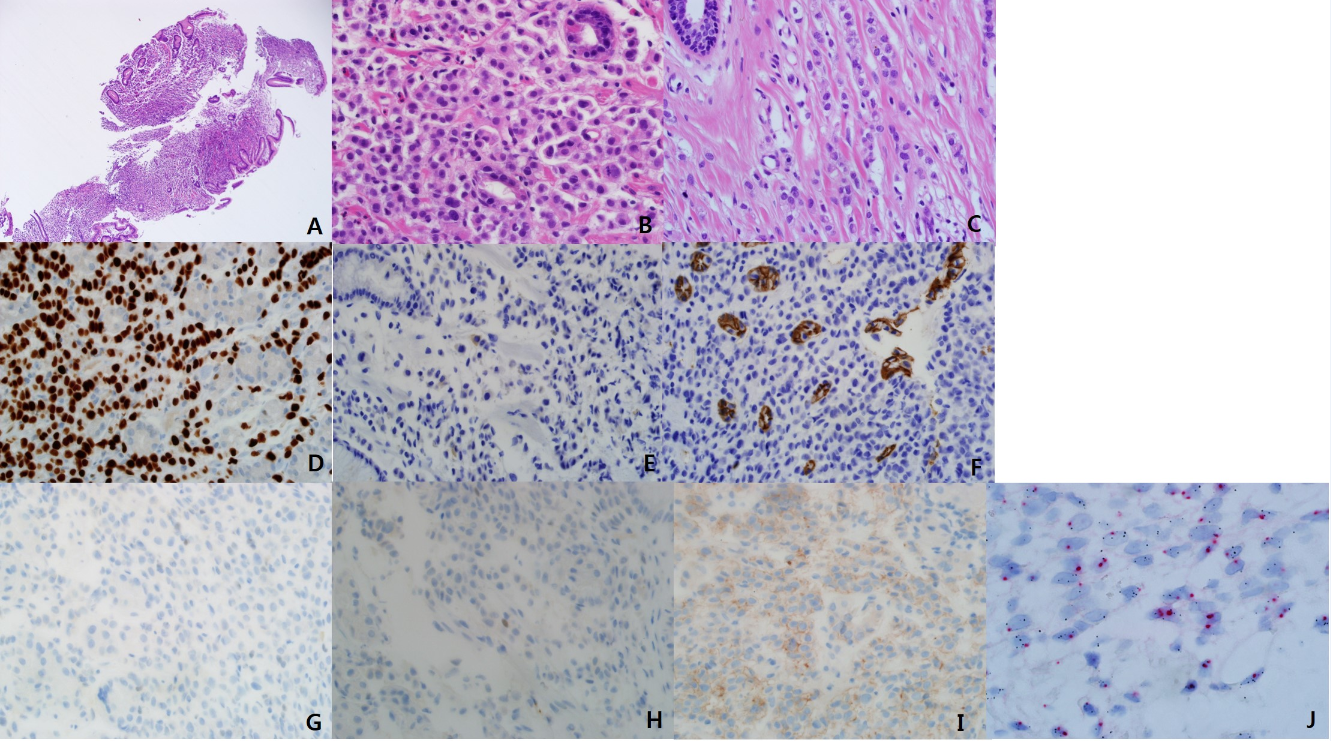
Table 1 Clinical characteristics of representative gastric metastasis of breast cancer

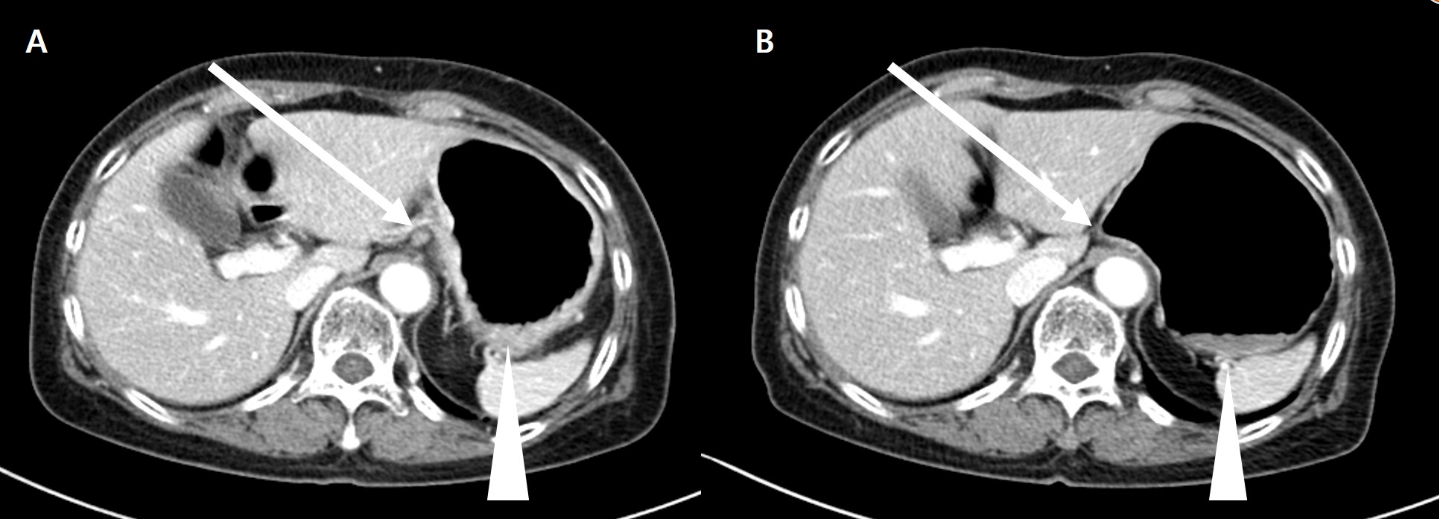
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| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Ref.** | **Age** | **Duration after initial diagnosis** | **Clinical presentation** | **Endoscopy** | **Pathology** | **IHC** | | | **Surgery** | **Treatment** | **Other metastases site** | **Overall survival** |
| **ER** | **PR** | **C-erbB2** |
| Our case | 65 | 4 | Epigastric Discomfort Indigestion | diffuse infiltrative mucosal lesion extensive nodular thickening | ILC | neg | neg | pos | No |  | bone | - |
| Pera *et al*[18] | 45 | 7 | Epigastric pain heart burn | erosion of gastric wall | ILC | pos | pos | - | Subtotal gastrectomy | H | - | - |
| Jones *et al*[2] | 51 | 3 | No symptom | polyp at antrum wall | ILC | neg | neg | neg | Total gastrectomy | Palliative | bone | - |
| 61 | 6.9 | Dysphagia weight loss | fungating mass | ILC | pos | pos | neg | No | C,R | brain, bone, pleura | - |
| Eo *et al*[11] | 48 | 9 | Nausea anorexia | elevated mucosal lesion | IDC | pos | pos | neg | No | C | liver, bone, pleura | - |
| Arragoiz *et al*[8] | 70 | 1 | Diarrhea constipation | mucosal thickening | ILC | pos | neg | neg | No | H | lung, rectum | - |
| Koike *et al*[16] | 42 | 5 | Epigastric pain | mucosal erosion | ILC | pos | pos | neg | No | C | - | - |
| 54 | 6 | Epigastric pain | mucosal erosion | ILC | pos | pos | neg | No | C, H | liver, bone, peritoneum | 5 |
| 54 | 3 | Epigastric pain vomiting | submucosal tumor | IDC | pos | pos | pos | No | C | bone | 2.3 |
| Geredeli *et al*[12] | 47 | 3 | Increased serum CEA, CA15-3 |  | ILC | neg | neg | neg | Subtotal gastrectomy | C | bone | - |
| Buka *et al*[9] | 58 | 1.2 | Abdominal pain weight loss | polypoid infiltration | ILC | pos | pos | neg | Total gastrectomy | C, R | colon, pleura | 7.2 |
| Lee *et al*[17] | 48 | 5.7 | Melena | mucosal erosion | - | - | - | - | - | C | bone, liver | - |
| Yim *et al*[19] | 48 | Initial diagnosis | Epigastric discormfort | mucosal erosion | ILC | neg | neg | - | No | C | bone | - |
| Jeon *et al*[14] | 49 | 5 | Melena | volcano shaped ulcers | IDC | pos | neg | - | No | C | bone |  |
| Kim *et al*[15] | 53 | 10 | Dyspepsia lower abdominal pain small caliper of stool | mucosal erosion | IDC | neg | neg | - | No | C, H | kidney, ovary, colon, bone, peritoneal LN | 2.4 |
| Hwang *et al*[13] | 66 | 17 | Back pain | flat mucosal lesion | ILC | neg | pos |  | Endoscropic mucosal resection | C | bone | - |
| Choei *et al*[10] | 56 | 4 | Upper abdominal discomfort | mucosal erosion | IDC | neg | pos | pos | - | C, H | - | 1.3 |
| Yu *et al*[3] | 63 | 10 | Melena small caliper of stool | linitis plastica flat ulcer | ILC | pos | pos | pos | No | C, H | Colon, bone marrow | - |

ILC: Invasive lobular carcinoma; IDC: Invasive ductal carcinoma; IHC: Immunohistochemical stain; pos: Positive; neg: Negative; C: Chemotherapy; H: Hormonal treatment.

**Figure 1 Upper endoscopy shows diffuse infiltrative mucosal lesion with extensive nodular thickening of the stomach wall, involving lower two-thirds of body.**

**Figure 2 Endoscopic ultrasound shows subserosal invasion of the gastric lesion with lymph node involvement (A, B).** Abdomen CT scan shows infiltrative gastric lesion involving cardia and angle of stomach (arrowhead) with enlarged perigastric lymph node (arrow).

**Figure 3 Pathologic features of endoscopic biopsy specimen.** Discohesive tumor cells are infiltrated in the stroma of the stomach mucosal tissue (H&E × 40, A). Tumor cells show enlarged centrally located nucleus without intracytoplasmic clear mucin. The tumor cells had no connection to the remained normal gastric mucosal tissue (H&E × 400, B). Previous breast cancer pathology was reviewed (C). Discohesive tumor cells were arranged in indian file. The tumor cells had enlarged centrally located nucleus without intracytoplasmic mucin (H&E × 400, C). Immunohistochemical stains and molecular test of tumor was done (D-J). Diffuse strong nucleus expression of GATA3 was observed (GATA3 × 400, D). Focal, less than one percentage cytoplasmic expression of GCDFP was detected (GCDFP × 400, E). Negative stain for E-cadherin (E-cadherin × 400, F). Negative stains for ER and PR (ER 400, PR × 400, G, H). Immunohistochemical stain for HER-2 was equivocal (HER-2 × 400, I). Silver in situ hybridization (SISH) for determination of HER2 gene status. Occasional HER2 gene amplified cells were noted in the mixture with normal HE2 gene expressing cells (SISH × 1000, J).

**Figure 4 Response evaluation after 2 cycles of docetaxel chemotherapy (A, B).** Abdominal CT scan shows decreased perigastric lymph modes (arrows) and gastric mucosal thickening (arrowheads).