

of the characteristics of lesions with an extremely low risk of lymph node metastasis have provided technical and theoretical bases for the en bloc resection of lesions larger than those resectable by conventional endoscopic mucosal resection (EMR) [2,4,12–15]. ESD is now indicated for the treatment of differentiated-type early gastric cancers with SM1 invasion that are 30 mm or less in diameter [2,4]. The indications for endoscopic resection are now gradually being extended to submucosal invasive gastric cancers that previously required surgical resection. To our knowledge, however, no previous study has evaluated the clinicopathologic features, prevalence of lymph node metastasis, lymphatic invasion, and ulceration of early gastric cancer containing both differentiated-type and undifferentiated-type components. Indications for the endoscopic treatment of mixed-histologic-type early gastric cancers have also not been evaluated.

Materials and methods

The study group comprised 376 patients with primary gastric cancer invading the submucosa (99 patients with SM1 invasion and 277 with SM2 invasion) who underwent surgical resection and a D2 lymphadenectomy according to the Japanese Classification of Gastric Carcinoma at Kitasato University East Hospital from 1995 through 2006. All lesions were thinly sliced at intervals of 3 to 5 mm. One section each of all dissected lymph nodes (at least 15 nodes per case) was stained with hematoxylin and eosin. The cut sections were examined histologically to assess the presence or absence of metastasis. To assess histologic type, all specimens were reviewed to determine the percentages of differentiated-type components (well and moderately differentiated tubular adenocarcinoma and papillary adenocarcinoma) and undifferentiated-type components (poorly differentiated adenocarcinoma and signet-ring cell carcinoma) [1]. The lesions were classified into the following four categories according to the proportions of intramucosal undifferentiated-type components: differentiated type (A), differentiated-type-predominant mixed type (B), undifferentiated-type-predominant mixed type (C), and undifferentiated type (D). The percentages of undifferentiated-type components were 0% in A, more than 0% but less than 50% in B, 50% or more but less than 100% in C, and 100% in D. A representative case of mixed-type gastric cancer is shown in Fig. 1.

The patterns of submucosal invasion were classified into four types (Fig. 2).

The horizontal length of submucosal invasion was measured histologically on the sections with greatest invasion (Fig. 3).

The number of lymphatic invasion sites and the number of lymphatic invasion sites per millimeter of submucosal invasion were calculated. Data on patient sex and age and tumor location, macroscopic type, and size were collected from the patients' medical records and pathology reports.

Immunohistochemistry

Lymphatic invasion by cancer cells was identified immunohistochemically using D2-40 antibody (DakoCytomation, Glostrup, Denmark) and an EnVision+ kit (DakoCytomation). One representative slide including the site of deepest invasion was selected for each case. Sections 4 μm thick were cut from the formalin-fixed, paraffin-embedded tissue. The sections were mounted on coated slides and deparaffinized in xylene. Nonspecific reactions were blocked with 0.3% hydrogen peroxide in methanol for 15

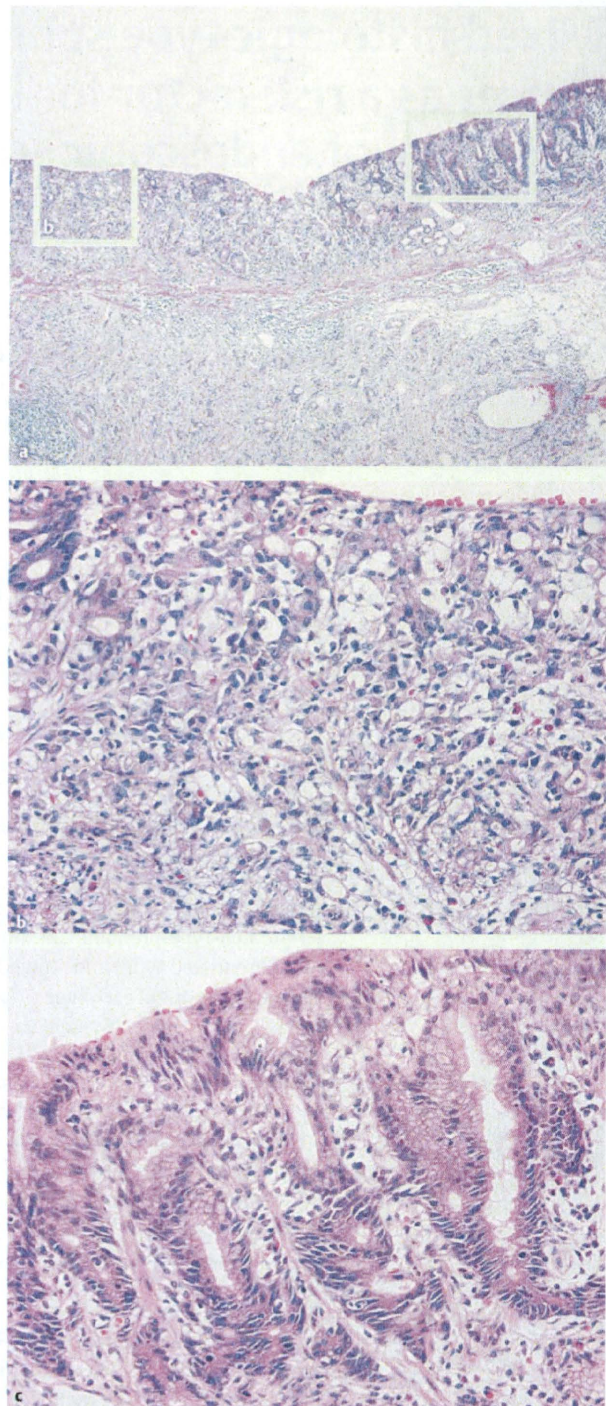


Fig. 1 Representative case of mixed-type (differentiated and undifferentiated) gastric carcinoma. **a** In the lamina propria, the right half of the tumor consists of the differentiated type and the left half of the undifferentiated type. The submucosal invasion consists of the undifferentiated type. **b** Box **b** in **a**, at higher magnification. **c** Box **c** in **a**, at higher magnification.

minutes and the blocking solution of the EnVision+ kit. Then, the sections were incubated with the D2-40 antibody at a 1:50 dilution in phosphate-buffered saline for 1 hour at room temperature. After incubation, staining was performed according to the manufacturer's instructions. 3-3'-Diaminobenzidine was used as the final chromogen, and nuclei were counterstained

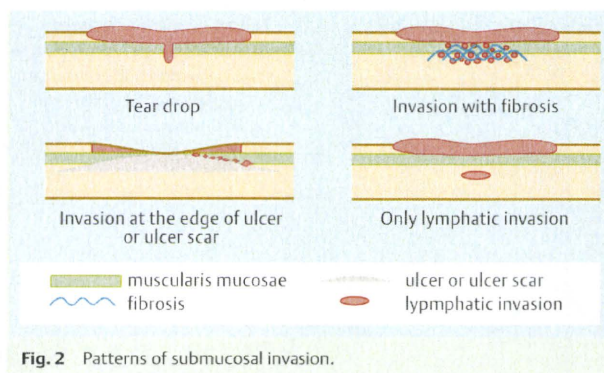


Fig. 2 Patterns of submucosal invasion.

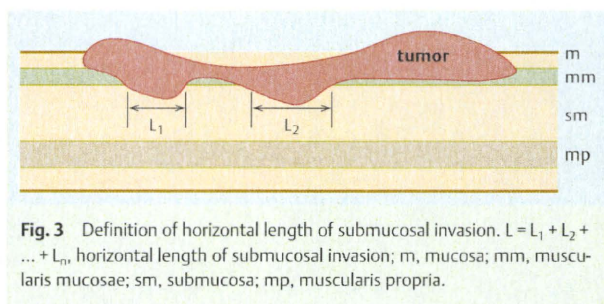


Fig. 3 Definition of horizontal length of submucosal invasion. $L = L_1 + L_2 + \dots + L_n$, horizontal length of submucosal invasion; m, mucosa; mm, muscularis mucosae; sm, submucosa; mp, muscularis propria.

with Mayer's hematoxylin to facilitate histopathologic assessment. Two pathologists (N.H. and T.M.) checked the slides to identify and count the number of lymphatic invasion sites, which appeared as cancer cell nests surrounded by D2-40-positive lymphatic endothelium (● Fig. 4).

Ethical approval

This work using pathological samples in Kitasato University East Hospital (with the informed consent of patients) was approved by our Medical School and University Ethics Committee.

Statistical analysis

To assess differences in tumor size and horizontal length of submucosal invasion among the four histologic groups (A, B, C, D), the Kruskal-Wallis test was used, followed by evaluation with the Mann-Whitney *U* test for multiple comparisons. Resulting *P* values were corrected according to the Bonferroni method. The χ^2 test was used to compare other clinicopathologic features. Multivariate logistic-regression analysis was performed with SPSS (version 11.0 in 2001; SPSS, Chicago, Illinois, USA). *P* values of less than 0.05 were considered to indicate statistical significance.

Results

The prevalence of lymph node metastasis was 16.5% (62/376) overall, 4.0% (4/99) in patients with SM1 invasion, and 20.9% (58/277) in those with SM2 invasion. Univariate analysis demonstrated that tumor size (> 30 mm), lymphatic invasion, histologic type, depth of invasion (SM2), and histologic type of the invasion front (undifferentiated type) differed significantly between patients with and those without lymph node metastasis (● Table 1).

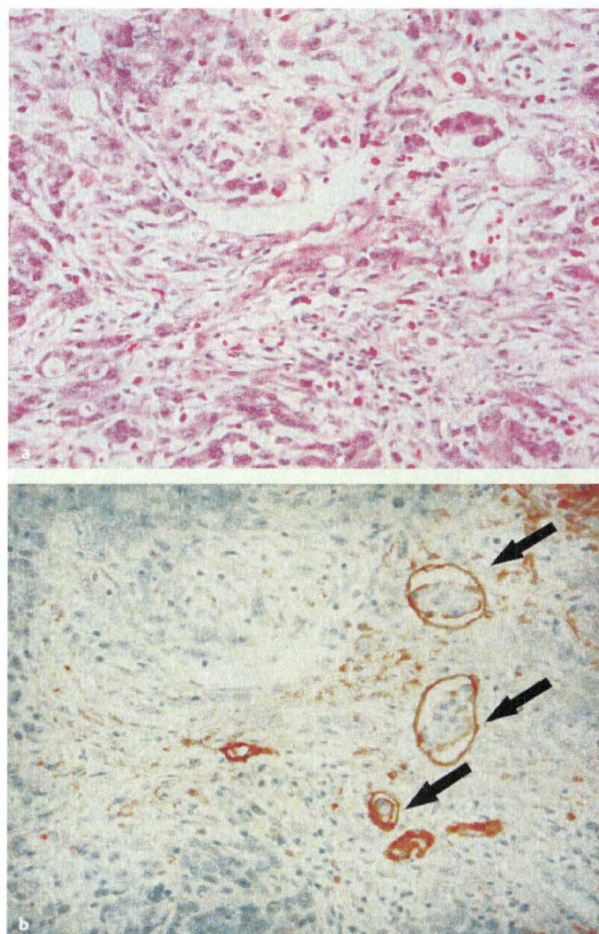


Fig. 4 a Submucosal invasion front of an undifferentiated-type gastric carcinoma. b Lymphatic invasion of cancer cells after D2-40 immunostaining. Cancer cells can be easily identified in lymphatic vessels (arrows).

Multivariate analysis of these risk factors indicated that lymphatic invasion, depth of invasion (SM2), tumor size (> 30 mm), and histologic type (C, undifferentiated-type-predominant mixed type) were independent risk factors for lymph node metastasis (● Table 2).

Relation between histologic type and lymph node metastasis

The prevalence of lymph node metastasis was 5.4% (7/129) in differentiated type (A), 19.2% (20/104) in differentiated-type-predominant mixed type (B), 36.5% (23/63) in undifferentiated-type-predominant mixed type (C), and 15.0% (12/80) in undifferentiated type (D). The prevalence of lymph node metastasis was highest in undifferentiated-type-predominant mixed type (C) ($P < 0.001$ vs. A, $P = 0.019$ vs. B, and $P = 0.0029$ vs. D) (● Table 3).

Submucosal cancers with a depth of invasion of SM1, no lymphatic invasion, a tumor size equal to or less than 30 mm, and consisting of less than 50% of undifferentiated components were free of lymph node metastasis (0/41; 95% confidence interval 0%–8.6%).

	n	Lymph node metastasis, n (%)		p-value
		Present	Absent	
Sex				
Male	271	39 (14.4)	232	0.078
Female	105	23 (21.9)	82	0.284
Age				
< 60	229	34 (14.8)	195	
≥ 60	147	28 (19.0)	119	
Location				0.171
Upper third	217	32 (14.7)	185	
Middle third	56	9 (16.1)	47	
Lower third	103	21 (20.4)	82	
Macroscopic type				0.758
Elevated	72	11 (15.3)	61	
Depressed	304	51 (16.8)	253	
Tumor size				< 0.001
≤ 30 mm	193	18 (9.3)	175	
> 30 mm	183	44 (24.0)	139	
Pattern of submucosal invasion*				0.473
1	3	0 (0.0)	3	
2	347	58 (16.7)	289	
3	24	3 (12.5)	21	
4	2	1 (50.0)	1	
Histologic type †				< 0.001
A	129	7 (5.4)	122	
B	104	20 (19.2)	84	
C	63	23 (36.5)	40	
D	80	12 (15.0)	68	
Lymphatic invasion				< 0.001
Yes	131	45 (34.4)	86	
No	245	17 (6.9)	228	
Histologic type of invasion front				< 0.001
Differentiated	183	17 (9.3)	166	
Undifferentiated	193	45 (23.3)	148	
Depth of invasion ‡				< 0.001
SM1	99	4 (4.0)	95	
SM2	277	58 (20.9)	219	

* 1, teardrop type; 2, invasion with fibrosis; 3, invasion at the edge of ulcer or ulcer scar; 4, only lymphatic invasion.

† A, differentiated type; B, differentiated-type-predominant mixed type; C, undifferentiated-type-predominant mixed type; D, undifferentiated type.

‡ SM1, depth of invasion from the muscularis mucosae < 500 μm; SM2, depth of invasion from the muscularis mucosae ≥ 500 μm.

Table 1 Relations between clinicopathologic factors and regional lymph node metastases in 376 cases of gastric cancer with submucosal invasion: results of univariate analysis.

Clinicopathologic feature	Relative risk (95% CI)	SE	P value
Lymphatic invasion (presence vs. absence)	5.28 (2.73–10.22)	0.337	< 0.001
Depth of invasion (SM2 vs. SM1)	3.41 (1.14–10.23)	0.560	0.029
Tumor size (> 30 vs. ≤ 30 mm)	2.58 (1.35–4.91)	0.329	0.004
Histologic type (C vs. non-C)	2.35 (1.12–4.96)	0.380	0.024

CI, confidence interval; SE, standard error.

Table 2 Risk factors for regional lymph node metastasis in 376 cases of gastric cancer with submucosal invasion: results of multivariate analysis.

Relations of histologic type and tumor size to lymphatic invasion

The prevalence of lymphatic invasion was highest in undifferentiated-type-predominant mixed type (C) ($P < 0.001$ vs. A, $P = 0.02$ vs. B, and $P = 0.03$ vs. D). Tumor size (47.2 ± 34.5 mm) and the width of submucosal invasion (8.6 ± 7.4 mm) were greatest in undifferentiated-type-predominant mixed type (C) and differed significantly from the respective values in differentiated type (A, 31.5 ± 20.5 , $P = 0.018$; 4.7 ± 5.1 ; $P < 0.001$) (Table 3). The number of lymphatic invasion sites per millimeter of submucosal invasion did not differ significantly among the histologic types. However, the presence of one or more lymphatic invasion sites per millimeter of submucosal invasion was associated with a higher prevalence of lymph node metastasis ($P = 0.030$) (Table 4).

Discussion

In the present study, we histologically classified early gastric cancers according to the percentage of undifferentiated components and found that the prevalence of lymph node metastasis was higher with the undifferentiated-type-predominant mixed type (C) than with the other histologic types. We also found that lymphatic invasion, SM2 invasion, tumor size, and undifferentiated-type-predominant mixed type were independent risk factors for lymph node metastasis in patients who had gastric cancer with submucosal invasion.

Previous clinicopathologic studies of gastric cancer, head and neck cancer, and breast cancer have demonstrated that lymphatic invasion is an extremely important risk factor for lymph node metastasis [16–18]. The advent of D2–40 immunostaining has

Table 3 Relations of histologic type and size of tumor to lymphatic invasion in 376 cases of gastric cancer.

Tumor type	Presence of lymph node metastasis, n (%)	Presence of lymphatic invasion, n (%)	Tumor size, mm, mean \pm SD	Horizontal length of submucosal invasion, mm, mean \pm SD	Number of lymphatic invasion sites, /mm, mean \pm SD
A (n = 129)	7 (5.4)	24 (18.6)	31.5 \pm 20.5	4.7 \pm 5.1	0.50 \pm 0.48
B (n = 104)	20 (19.2)	40 (38.5)	39.5 \pm 25.9	6.0 \pm 5.5	1.1 \pm 2.3
C (n = 63)	23 (36.5 [*])	36 (57.1 ^{**})	47.2 [†] \pm 34.5	8.6 [‡] \pm 7.4	1.2 \pm 2.7
D (n = 80)	12 (15.0)	31 (38.8)	37.0 \pm 22.2	6.7 \pm 6.1	0.46 \pm 0.69

P values were determined using the Mann-Whitney U test with Bonferroni correction.

^{*}P < 0.001 vs. A, P = 0.013 vs. B, P = 0.003 vs. D (χ^2 test).

^{**}P < 0.001 vs. A, P = 0.019 vs. B, P = 0.029 vs. D (χ^2 test).

[†]P = 0.018 vs. A.

[‡]P < 0.001 vs. A.

	n	Lymph node metastasis		P value
		Present	Absent	
Number of lymphatic invasion sites \geq 1	24	13 (54.1 [*])	11	< 0.001
Number of lymphatic invasion sites < 1	107	33 (30.8)	74	
Cases without lymphatic invasion	245	16 (6.9)	229	

^{*}P = 0.030 vs. number of lymphatic invasion sites < 1.

Table 4 Relation between lymph node metastasis and number of lymphatic invasion sites per millimeter of horizontal length of submucosal invasion in 376 cases of gastric cancer.

facilitated the identification of lymph vessels, and many studies have reported that immunostaining is useful for the evaluation of gastric cancer [19–23]. We therefore clinicopathologically studied the relationship of the histologic type of gastric cancer to lymph node metastasis from the viewpoints of lymphatic invasion and histogenesis.

Undifferentiated-type-predominant mixed type (C) had the highest prevalence of lymphatic invasion. The presence of lymphatic invasion was characterized by a high prevalence of lymph node metastasis. In this study, the existence of one or more lymphatic invasion sites per millimeter of submucosal invasion was associated with an increased prevalence of lymph node metastasis. The higher prevalence of lymph node metastasis in undifferentiated-type-predominant mixed type (C) than in the other groups was attributed to the significant differences in the presence of lymphatic invasion among these groups.

The fact that the highest prevalence of lymphatic invasion is seen in undifferentiated-type-predominant mixed type (C) gastric cancer might be related to the size of the tumor and the horizontal length of the submucosal invasion, which were larger in this group than in the others. Lymph vessels are densely present from the muscularis mucosae to the upper layer of the submucosa [19]. With tumor-cell proliferation and extension to the muscularis mucosae and submucosa, the incidence of lymphatic invasion increases.

Gastric cancer generally shows greater histologic diversity than other types of cancer. Even tumors confined to the mucosa show histologic diversity, which tends to increase with deeper invasion and increased tumor diameter [24,25]. This notion is supported by the findings of Inoshita et al., who studied histologic diversity in gastric cancer. In elderly patients, they found that the differentiated type predominates in early gastric cancer, but that histologic diversity increases with progression to advanced cancer, resulting in higher proportions of undifferentiated type [26]. Peng and Honda et al. reported that undifferentiated-type gastric cancer with tubular components (C: undifferentiated-type-predominant mixed type) and undifferentiated-type gastric cancer (D) arise from different genetic pathways. They proposed that mixed-type gastric cancer can arise from either differentiated

cells or undifferentiated cells [27,28]. Studies of mucin phenotype have reported that some cases of differentiated-type gastric cancer with gastric phenotype are transformed into undifferentiated-type gastric cancer during tumor growth and development, increasing the risk of lymph node metastasis [29–34]. However, further studies are needed to delineate the relation between histogenesis and the risk of metastasis.

At present, the indications for endoscopic resection include the treatment of submucosal cancers that meet the following four conditions: a tumor size of 30 mm or less, SM1 invasion, differentiated type, and no lymphatic invasion [2,4]. In our study, lymph node metastasis was not associated with submucosal cancers that met all of the following criteria: SM1 invasion, no lymphatic invasion, a tumor size of 30 mm or less, and less than 50% undifferentiated-type components (0/41; 95% confidence interval, 0% to 8.6%). Such cancers can be curatively treated by local endoscopic resection.

Gastric cancer is associated with underlying conditions such as acid-induced changes and chronic gastritis caused by persistent *Helicobacter pylori* infection. Disease progression is often accompanied by the formation of an ulcer or ulcer scar. It is challenging to predict preoperatively the percentage of undifferentiated components and to accurately diagnose the depth of tumor invasion on the basis of the fine surface characteristics of gastric cancers. In particular, the accuracy of endoscopic ultrasonography for predicting whether a tumor is confined to the mucosa or invades the submucosa is only about 80% [35]. Submucosal tumor invasion is frequently discovered on histopathologic examination after endoscopic resection. If an early gastric cancer is found to have SM1 invasion on histopathologic examination after endoscopic resection, the presence of a tumor that is 30 mm or greater in size, lymphatic invasion, or a 50% or higher percentage of undifferentiated components suggests an increased risk of lymph node metastasis. Additional surgical resection with lymph node dissection should therefore be considered.

Undifferentiated early gastric cancer has a high risk of lymph node metastasis [2–5]. To date, surgical resection has been the treatment of choice, but the feasibility of endoscopic resection is now being considered. Park et al. reported that endoscopic resec-

tion can be indicated for undifferentiated-type cancers that are 15 mm or less in diameter and have a submucosal invasion depth of up to 500 µm because these characteristics are associated with a low risk of lymph node metastasis [36]. In their study, however, histologic types and lymphatic invasion were not adequately evaluated. In our study, the incidence of lymph node metastasis was higher in undifferentiated-type-predominant mixed type (C) than in pure undifferentiated type (D) gastric cancer. Endoscopic resection should therefore not be conducted in patients with undifferentiated-type gastric cancer without detailed histopathologic studies, including assessments of histologic diversity and lymphatic invasion.

We believe that confirmation of lymphatic invasion by means of D2–40 immunostaining may lead to more accurate identification of cases at high risk of lymph node metastasis.

We conclude that histologically mixed-type gastric cancer with submucosal invasion can be considered for endoscopic resection provided that the following four conditions indicating a low risk of metastasis are met: a lower than 50% proportion of undifferentiated components, a tumor size of 30 mm or less, SM1 invasion, and no lymphatic invasion. Given the limited number of cases in this study, these findings should be confirmed by more data in the future.

Acknowledgment

We are indebted to the members of the Department of Pathology, Kitasato University East Hospital, for their excellent technical assistance.

Competing interests: None

References

- 1 Japanese Gastric Cancer Association. Japanese classification of gastric carcinoma: 2nd English edition. *Gastric Cancer* 1998; 1: 10–24
- 2 Gotoda T, Yanagisawa A, Sasako M et al. Incidence of lymph node metastasis from early gastric cancer: estimation with a large number of cases at two large centers. *Gastric Cancer* 2000; 3: 219–225
- 3 Yamao T, Shirao K, Ono H et al. Risk factors for lymph node metastasis from intramucosal gastric carcinoma. *Cancer* 1996; 77: 602–606
- 4 Gotoda T, Sasako M, Ono H et al. Evaluation of the necessity for gastrectomy with lymph node dissection for patients with submucosal invasive gastric cancer. *Br J Surg* 2001; 88: 444–449
- 5 Kurihara N, Kubota T, Otani Y et al. Lymph node metastasis of early gastric cancer with submucosal invasion. *Br J Surg* 1998; 85: 835–839
- 6 Korenaga D, Haraguchi M, Tsujitani S et al. Clinicopathological features of mucosal carcinoma of the stomach with lymph node metastasis in eleven patients. *Br J Surg* 1986; 73: 431–433
- 7 Ono H, Kondo H, Gotoda T et al. Endoscopic mucosal resection for treatment of early gastric cancer. *Gut* 2001; 48: 225–229
- 8 Tanabe S, Koizumi W, Mitomi H et al. Clinical outcome of endoscopic aspiration mucosectomy for early gastric cancer. *Gastrointest Endosc* 2002; 56: 708–713
- 9 Maehara Y, Orita H, Okuyama T et al. Predictors of lymph node metastasis in early gastric cancer. *Br J Surg* 1992; 79: 245–247
- 10 Sano T, Kobori O, Muto T. Lymph node metastasis from early gastric cancer: endoscopic resection of tumour. *Br J Surg* 1992; 79: 241–244
- 11 Ohkuwa M, Hosokawa K, Boku N et al. New endoscopic treatment for intramucosal gastric tumors using an insulated-tip diathermic knife. *Endoscopy* 2001; 33: 221–226
- 12 Karita M, Tada M, Okita K. The successive strip biopsy partial resection technique for large early gastric and colon cancers. *Gastrointest Endosc* 1992; 38: 174–178
- 13 Inoue H, Takeshita K, Hori H et al. Endoscopic mucosal resection with cap-fitted panendoscope for esophagus, stomach, and colon mucosal lesions. *Gastrointest Endosc* 1993; 39: 58–62
- 14 Torii A, Sasaki M, Kajiyama T et al. Endoscopic aspiration mucosectomy as curative endoscopic surgery: analysis of 24 cases of early gastric cancer. *Gastrointest Endosc* 1995; 42: 475–479
- 15 Hanazaki K, Wakabayashi M, Sodeyama H et al. Clinicopathologic features of submucosal carcinoma of the stomach. *J Clin Gastroenterol* 1997; 24: 150–155
- 16 Bando E, Yonemura Y, Taniguchi K et al. Outcome of ratio of lymph node metastasis in gastric carcinoma. *Ann Surg Oncol* 2002; 9: 775–784
- 17 Bimer P, Obermair A, Achindl M et al. Selective immunohistochemical staging of blood and lymphatic vessels reveals independent prognostic influence of blood and lymphatic invasion in early-stage cervical cancer. *Clin Cancer Res* 2001; 7: 93–97
- 18 Clemente CG, Boracchi P, Andreola S et al. Peritumoral lymphatic invasion in patients with node negative mammary duct carcinoma. *Cancer* 1992; 69: 1396–1403
- 19 Sako A, Kitayama J, Ishikawa M et al. Impact of immunohistochemically identified lymphatic invasion on nodal metastasis in early gastric cancer. *Gastric Cancer* 2006; 9: 295–302
- 20 Schoppmann SF, Birner P, Studer P et al. Lymphatic microvessel density and lymphovascular invasion assessed by antipodoplanin immunostaining in human breast cancer. *Anticancer Res* 2001; 21: 2351–2356
- 21 Kahn HJ, Bailey D, Marks A. A new monoclonal antibody, D2-40, for detection of lymphatic invasion in primary tumors. *Lab Invest* 2002; 82: 1255–1257
- 22 Arigami T, Natsugome S, Uenosono Y et al. Lymphatic invasion using D2-40 monoclonal antibody and its relationship to lymph node micrometastasis in pN0 gastric cancer. *Br J Cancer* 2005; 93: 688–693
- 23 Yonemura Y, Endou Y, Tabuchi K et al. Evaluation of lymphatic invasion in primary gastric cancer by a new monoclonal antibody, D2-40. *Hum Pathol* 2006; 37: 1193–1199
- 24 Luinetti O, Fiocca R, Villani L et al. Genetic pattern, histological structure, and cellular phenotype in early and advanced gastric cancers: evidence for structure-related genetic subsets and for loss of glandular structure during progression of some tumors. *Hum Pathol* 1998; 29: 702–709
- 25 Ishiguro S, Kasugai T, Terada N. Change of histological type of gastric carcinoma: from differentiated carcinoma to undifferentiated carcinoma [in Japanese with English abstract]. *Stomach and Intestine* 1996; 31: 1437–1443
- 26 Inoshita N, Yanagisawa A, Arai T et al. Pathological characteristics of gastric carcinomas in the very old. *Jpn J Cancer Res* 1998; 89: 1087–1092
- 27 Peng DF, Sugihara H, Mukaisho K et al. Genetic lineage of poorly differentiated gastric carcinoma with a tubular component analysed by comparative genomic hybridization. *J Pathol* 2004; 203: 884–895
- 28 Honda T, Tamura G, Endoh Y et al. Expression of tumor suppressor and tumor-related proteins in differentiated carcinoma, undifferentiated carcinoma with tubular component and pure undifferentiated carcinoma of the stomach. *Jpn J Clin Oncol* 2005; 35: 580–586
- 29 Egashira Y, Shimoda T, Ikegami M. Mucin histochemical analysis of minute gastric differentiated adenocarcinoma. *Pathol Int* 1999; 49: 55–61
- 30 Saito A, Shimoda T, Nakanishi Y et al. Histologic heterogeneity and mucin phenotypic expression in early gastric cancer. *Pathol Int* 2001; 51: 165–171
- 31 Tajima Y, Shimoda T, Nakanishi Y et al. Gastric and intestinal phenotypic marker expression in gastric carcinomas and its prognostic significance: immunohistochemical analysis of 136 lesions. *Oncology* 2001; 61: 212–220
- 32 Kabashima A, Yao T, Maehara Y et al. Relationship between biological behavior and phenotypic expression in undifferentiated-type gastric carcinomas. *Gastric Cancer* 2005; 8: 220–227
- 33 Kushima R, Hattori T. Histogenesis and characteristics of gastric-type adenocarcinomas in the stomach. *J Cancer Res Clin Oncol* 1993; 120: 103–111
- 34 Yoshikawa A, Inada K, Yamachika T et al. Phenotypic shift in human differentiated gastric cancers from gastric to intestinal epithelial cell type during disease progression. *Gastric Cancer* 1998; 1: 134–141
- 35 Hizawa K, Iwai K, Esaki M et al. Is endoscopic ultrasonography indispensable in assessing the appropriateness of endoscopic resection for gastric cancer? *Endoscopy* 2002; 34: 973–978
- 36 Park YD, Chung YJ, Chung HY et al. Factors related to lymph node metastasis and the feasibility of endoscopic mucosal resection for treating poorly differentiated adenocarcinoma of the stomach. *Endoscopy* 2008; 40: 7–10

Prospective clinical trial of magnetic-anchor-guided endoscopic submucosal dissection for large early gastric cancer (with videos)

Takuji Gotoda, MD, Ichiro Oda, MD, Katsunori Tamakawa, PhD, Hirohisa Ueda, PhD, Toshiaki Kobayashi, MD, PhD, Tadao Kakizoe, MD, PhD

Tokyo, Japan

Background: The treatment of early gastric cancer (EGC) by endoscopic submucosal dissection (ESD) has been rapidly gaining popularity in Japan. However, the procedure needs a high quality of skill. To facilitate complicated ESD by using a single working-channel gastroscope (“one-hand surgery method”), the magnetic-anchor-guided ESD (MAG-ESD) controlled by an extracorporeal electromagnet was reported to be successful in a porcine model.

Objectives: The purpose of this prospective clinical trial was to evaluate the feasibility of MAG-ESD for large EGC located on the gastric body in human beings.

Design: Prospective clinical trial at a single center.

Setting: National Cancer Center Hospital, Tokyo, Japan.

Subjects: From January 2005 to May 2006, 25 patients with EGC > 20 mm in diameter, located in the gastric body, and intestinal-type histology were enrolled. Patients with a cardiac pacemaker, advanced malignancy in other organs, severe cardiac and/or pulmonary diseases, and uncontrolled hypertension and/or diabetes mellitus were excluded from this study.

Interventions: Similar to a standard ESD, the MAG-ESD procedure was performed with the patient under conscious sedation by intravenous injection of midazolam (3–5 mg) and pentazocine (15 mg).

Main Outcome Measurements: Unfavorable events and other intraoperative complications caused by the magnetic anchor or the magnetic force were recorded and evaluated. Two GI endoscopists (T.G., I.O.) assessed whether the magnetic anchor facilitated gastric ESD according to 2 criteria: “supportive” and “not supportive.” The en bloc resection rate, complications, total operation time, bleeding, perforation, and recurrence rate were also evaluated. The total operation time was measured from insertion to withdrawal of the endoscope, including the retrieving of the magnetic anchor or anchors.

Results: All tumors were resected en bloc, without any perforations or severe uncontrollable bleeding. All magnetic anchors were safely retrieved. Two endoscopists assessed that the MAG system was supportive in 23 patients. None of the patients experienced physiologic and mental abnormalities as a result of long-term magnetic-field exposure. During a median follow-up of 20 months (15–32 months), neither delayed adverse effects nor allergies caused by the stainless steel of the magnetic anchor were observed.

Conclusions: MAG-ESD is a feasible and safe method that allowed an excellent visualization by suitable tissue tension and facilitated gastric ESD in patients with EGC. The system should be miniaturized to make it applicable in daily clinical practice. (Gastrointest Endosc 2009;69:10-5.)

Abbreviations: EGC, early gastric cancer; ESD, endoscopic submucosal dissection; IT-knife, insulation-tipped diathermic knife; MAG-ESD, magnetic-anchor-guided endoscopic submucosal dissection.

DISCLOSURE: The authors report that there are no disclosures relevant to this publication. This study was supported by a grant-in-aid for the Research on Advanced Medical Technology of the Ministry of Health, Labor and Welfare, and a grant-in-aid for the Third Term Comprehensive 10-year Strategy for Cancer Control, of the Ministry of Health, Labor and Welfare, Japan.

Copyright © 2009 by the American Society for Gastrointestinal Endoscopy
0016-5107/\$36.00
doi:10.1016/j.gie.2008.03.1127

It has been reported that endoscopic submucosal dissection (ESD) of early gastric cancer (EGC) improves the rate of successful en bloc resection.^{1,2} An ESD by using an insulation-tipped diathermic knife (IT-knife), developed at the National Cancer Center Hospital, was the first of such techniques.^{3,4} Other endoscopic devices for ESD have been developed.⁵⁻⁷ ESD has been rapidly gaining popularity in Japan, primarily because of its ability to remove larger EGC en bloc, thus reducing a local recurrence caused by a piecemeal resection.⁸ However, it is still an

investigational technique and requires a high level of skill from the endoscopists.⁹⁻¹¹

Endoscopic resection should be safe, effective, and applicable to a wide variety of clinical situations. In particular, when EGC is located in the gastric body, an ESD is more complicated, and the rate of a complete resection is lower than in the gastric antrum.¹² The more difficult extension of the wall and the collection of fluid, including blood and/or gastric juice, hinder the performance of the ESD procedure. Optimal extension of the wall and visualization of the lesion is mandatory for a safe and feasible ESD.

To facilitate a complicated standard ESD procedure performed by using a single working-channel gastroscope (one-hand surgery), the magnetic-anchor-guided ESD (MAG-ESD) controlled by an extracorporeal electromagnet, was developed.¹³ We reported that MAG-ESD facilitated the ESD procedure in the porcine model. The purpose of this prospective clinical trial was to evaluate the feasibility of MAG-ESD for large EGC in human beings.

PATIENTS AND METHODS

Patients

The purpose of this prospective clinical trial was to evaluate the feasibility of MAG-ESD. Twenty-five patients with EGC >20 mm diameter, located in the gastric body, were enrolled. The patients were first seen on an outpatient basis, and the tumor was assessed by a gastroscopy. From January 2005 to May 2006, all patients with EGC >20 mm in diameter, located in the gastric body, and with intestinal-type histology underwent an ESD on an inpatient basis at the National Cancer Center Hospital, Tokyo, Japan. The ethics committee approved the study, and a detailed written informed consent was obtained from each patient. The presented study was conducted according to the Declaration of Helsinki.

The patients with a cardiac pacemaker, advanced malignancy in other organs, severe cardiac and/or pulmonary diseases, uncontrolled hypertension, and/or diabetes mellitus were excluded from this study. Pregnant or lactating women, and those who wished to become pregnant during the study were also excluded. Patients with tumors with recurrent disease, fibrosis, deeper invasion, or diffuse-type histology were excluded.

Standard ESD

The standard ESD procedure was initially started by using a standard gastroscope with a single working channel (GIF Q260 or Q240; Olympus Optical Co, Ltd, Tokyo, Japan).¹⁴ Marking dots were placed approximately 5 mm outside the margin of the lesions by using a needle-knife (KD-1L-1; Olympus) and forced coagulation current 20 W (IC C200; ERBE, Tübingen, Germany). First, injection

Capsule Summary

What is already known on this topic

- Endoscopic submucosal dissection (ESD) is useful in the en bloc removal of large gastric lesions, thus reducing the risk of a local recurrence caused by piecemeal resection.
- Magnetic-anchor-guided ESD (MAG-ESD), controlled by an extracorporeal electromagnet, facilitates the standard ESD procedure performed by using a single working-channel gastroscope.

What this study adds to our knowledge

- In 25 patients with gastric cancer lesions >20 mm in diameter who underwent magnetic-anchor-guided ESD, all tumors were resected en bloc, without any perforations or severe uncontrollable bleeding, and all magnetic anchors were safely retrieved.
- No patient experienced physiologic or mental abnormalities as a result of long-term magnetic field exposure.

of diluted epinephrine (1:100,000) was performed to raise the submucosal layer and to insert the tip of the IT-knife into the submucosal layer. Then, a small initial incision was made by a standard needle-knife by using 80 W; effect 3 Endocut (ICC200; ERBE). Mucosal cutting at the periphery of the marking dots was circumferentially performed with an IT-knife (KD-610L; Olympus) with 80 W Endocut. After additional submucosal injection of diluted epinephrine, the submucosal layer below the lesion was directly dissected by using the same IT-knife. The final aim was to achieve en bloc resection.

All patients were sedated by intravenous injection of midazolam (3–5 mg) and pentazocine (15 mg), and, if necessary, conscious sedation was maintained with an additional injection of midazolam.

Magnetic anchor and extracorporeal electromagnetic control system

The magnetic anchor (Pentax Co, Tokyo, Japan) consists of 3 parts: a hand-made magnetic weight, made of magnetic stainless steel (SYS420F), microforceps, and a connecting thread. A 1.0 × 1.0 × 1.5-cm weight was designed to facilitate gastric ESD by use of an extracorporeal hands-free electromagnet, whereby magnetic forces allow a suitable counter-traction for submucosal dissection (Fig. 1). The anchor weight used for this procedure was approximately 6 g.

The magnetic control system (Fig. 2) consists of an electromagnet with up-and-down motion; a movable examination table was made by Tamakawa Co (Sendai, Japan) for use in a standard endoscopic room. The magnetic control system consisted of a 0.68 kOe/100A extracorporeal

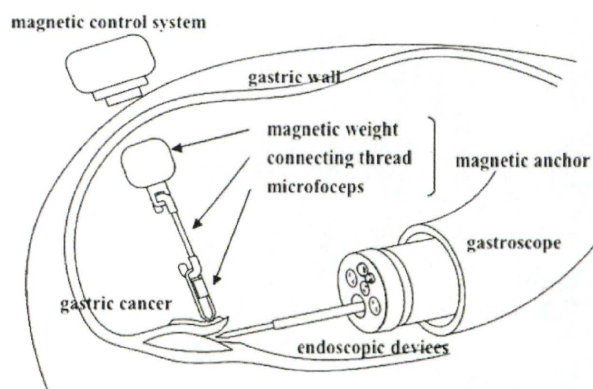


Figure 1. Concept of the MAG-ESD.

electromagnet, 350 mm in diameter, positioned at 10 cm from the center of the magnetic yoke. In this manner, the position of the electromagnet was adjusted according to the patient's physique. The examination table was able to move freely to be able to control the magnetic weight so as to achieve ideal mucosal lifting to allow the gastric submucosal dissection.

MAG-ESD

According to the standard ESD, after circumferential mucosal cutting by using an IT-knife, the procedure was switched to an MAG-ESD, controlled by a high-power electromagnet placed outside the body of the patient (Fig. 3). First, an overtube (Sumitomo Bakelite, Tokyo, Japan) was inserted into the esophagus. Second, a tube catheter was passed through the working channel of the gastroscop. A magnetic anchor, with a magnetic weight, a microforceps, and a connecting thread, was attached to the tip of the catheter. The gastroscop that carries the magnetic anchor was reinserted. Inside the stomach, the magnetic weight was pushed out from the catheter. According to the direction of gravity, the microforceps connected to the magnetic weight was placed at the mucosal edge (Video 1, available online at www.giejournal.org). The submucosal dissection by using an IT-knife was performed by suitable tissue tension with hands-free stabilization and visualization (Video 2, available online at www.giejournal.org).

If experienced endoscopists, who have performed more than 100 gastric ESDs, requested additional magnetic anchors to maneuver the traction direction of the exfoliated gastric tissue, then any numbers of magnetic anchors were attached. To maintain suitable tissue tension, either the patients were rotated or the direction of the magnetic anchor was repositioned by using the movable examination table. After endoscopic resection, both the resected tissue and the magnetic anchor or anchors were retrieved into the overtube by using a grasping forceps and were removed from the stomach.

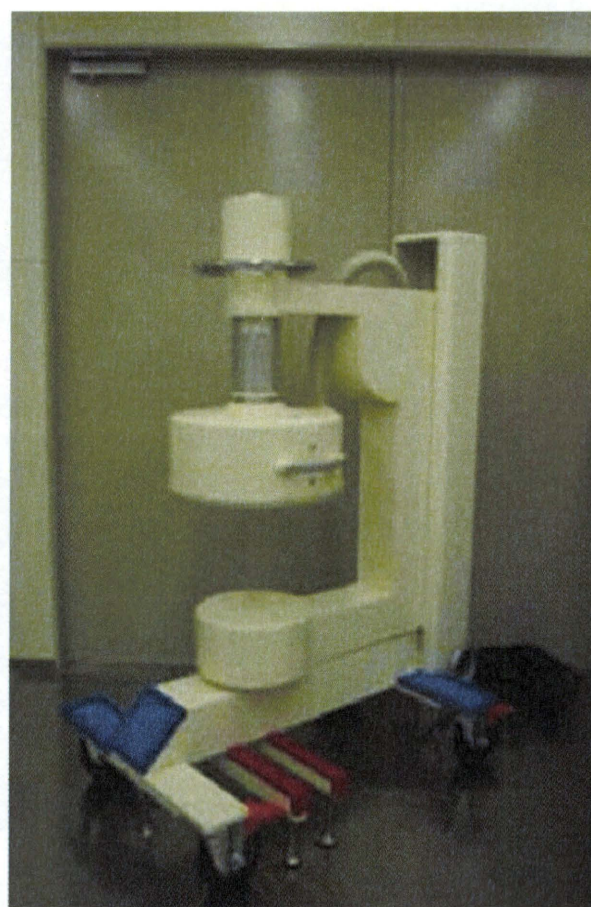


Figure 2. Extracorporeal electromagnetic control system.

Assessments

The demographic and clinical features of each patient were recorded in a case report form. Unfavorable events and other intraoperative complications caused by the magnetic anchor or the magnetic force were recorded and evaluated. We defined serious adverse events as those that lead to death, threat to life, notable disability, prolonged hospital stay, or hospitalization. Patients were followed-up until adverse events either dissipated or returned to pretreatment levels. Two GI endoscopists (T.G., I.O.) assessed, according to the 2 criteria, whether the magnetic anchor facilitated a gastric ESD. Once the dedicated endoscopists evaluated that the MAG-traction-facilitated gastric ESD compared with the standard gastric ESD technique, it was defined as "supportive." When the ESD procedure was not effectively influenced by using the MAG system, it was defined as "not supportive." The en bloc resection rate, complications, total operation time, bleeding, perforation, and recurrence rate were also evaluated. The total operation time was measured from gastroscop insertion to withdrawal, including retrieving the magnetic anchor or anchors.

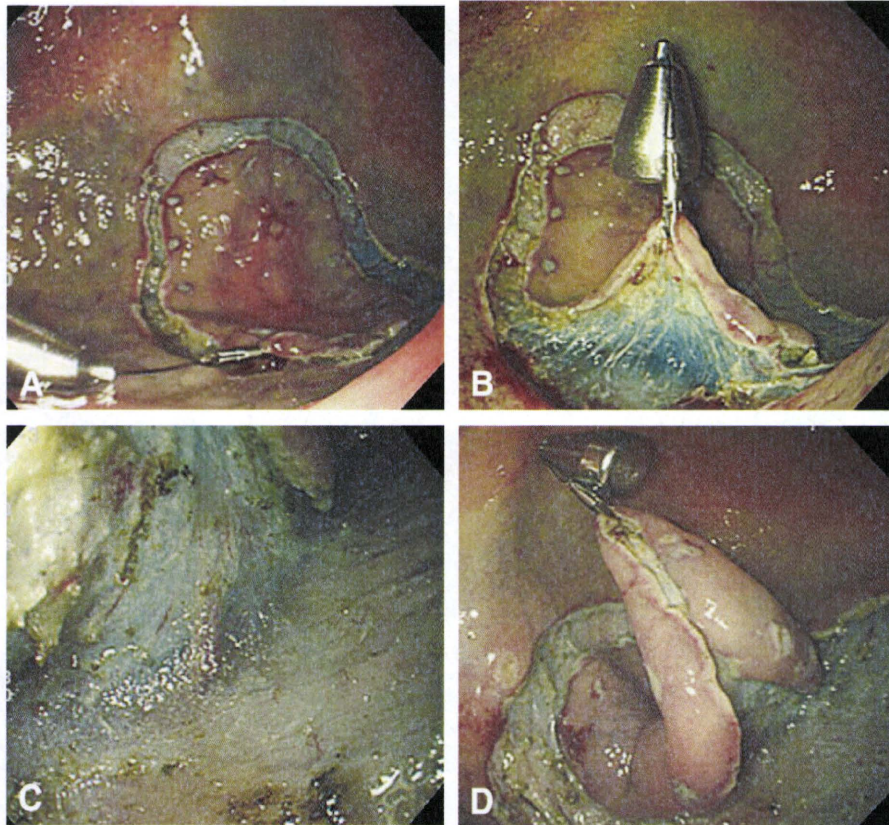


Figure 3. Magnetic-anchor-assisted ESD for large EGC. **A,** Fitting the magnetic anchor onto the tip of the gastric mucosa before applying the magnetic force. **B,** Lifting of the gastric tissue and stretched submucosal layer under strong counter-traction by the magnetic anchor. **C,** Good visualization of vessel in submucosal layer under counter-traction by magnetic force. **D,** Controllable traction by the magnetic anchor with a magnetic field.

RESULTS

The MAG-ESD technique was performed in 25 patients (M/F, 17/8; median age 70 years, range 48-85 years; median tumor size, 30 mm, range 20-70 mm).

The results of the MAG-ESDs are shown in Table 1. All tumors were resected en bloc, without any perforations or severe uncontrollable bleeding. The median size of resected specimen was 55 mm (33-125 mm). The median procedure time was 80 minutes (50-240 minutes). One resection was histologically confirmed as being noncurative because of deep submucosal invasion with positive vertical margins and lymphatic-vessel involvement. This patient underwent additional radical surgery.

One magnetic anchor was required in 21 cases, and 2 magnetic anchors were used in 4 cases. All magnetic anchors were safely retrieved. Two endoscopists assessed that the MAG system was supportive in 23 patients. In particular, the MAG system effectively facilitated an ESD for all 9 tumors located on the greater curvature of the gastric body. However, the magnetic anchor was not helpful in 2 patients. In one case, it was difficult to inflate the gastric lumen because of air leakage through the hiatus hernia.

TABLE 1. Results of 25 patients treated by MAG-ESD

En bloc resection rate	25/25 (100%)
Median resection size (mm)	55 (range 33-125)
Complications	0/25 (0%)
Median time consumption (min)	80 (range 50-240)
Exposure time for magnetic field (min)	30 (range 10-110)
Endoscopist's assessment	
Supportive	23
Not supportive	2

In another case, it was impossible to pull the gastric tissue toward the proper direction, even after changing the patient's position.

None of the patients experienced physiologic and mental abnormalities as a result of long-term magnetic-field exposure, neither before nor after the procedure. After a mean of 30 minutes (range 10–110 minutes) of exposure

to the magnetic field, no adverse effects of standard ESD procedure were observed regarding pulmonary and cardiac function. During a median follow-up of 20 months (range 15–32 months), neither delayed adverse effects nor allergies were observed because of the stainless steel of the magnetic anchor.

Eight weeks after an MAG-ESD, all artificial defects caused by ESD were completely cured. Neither recurrent cancer nor distant metastases were observed in any of the patients during follow-up.

DISCUSSION

The present study is, to our knowledge, the first clinical trial by using MAG-ESD for EGC in human beings. The feasibility of the technique for gastric cancer treatment was already evaluated in an animal study. The MAG-ESD technique permits excellent visualization of the submucosal layer, because it is possible to achieve suitable tissue tension. This simplifies a gastric ESD, even for large lesions located in the gastric body. The long-term exposure to the magnetic field did not cause any unwanted physiologic or mental effects. Furthermore, no delayed complications or allergies related to the stainless steel of the magnetic anchor were observed. All the tumors were resected en bloc, without any perforation or severe uncontrollable bleeding.

Endoscopic resection is comparable in many respects to conventional surgery, with the advantages of being less invasive and more cost efficient.^{15,16} Endoscopic removal of cancer was initially attempted by using colorectal polypectomy with a high-frequency electric surgical cautery.¹⁷ The use of endoscopic polypectomy to treat pedunculated or semipedunculated EGC was first described in 1974 in Japan. In 1984, the technique of EMR, the so-called strip biopsy, was devised for endoscopic snare polypectomy.¹⁸ Today, EMR is established and widely accepted as a minimally invasive treatment for EGC.¹⁹ Although several techniques have been reported to make EMR procedures easier and safer,^{20,21} these cannot be used to remove, en bloc, lesions larger than 2 cm in diameter.^{22,23} Piecemeal resection may cause the pathologist to inadequately stage the specimen. Furthermore, there is a high risk of a recurrence after a piecemeal resection.^{24,25}

An ESD is superior to a standard EMR and provides en bloc specimens with a standard single-channel gastroscop. After an endoscopic resection, pathologic assessment of depth of cancer invasion, degree of cancer differentiation, and lymphatic or blood-vessel involvement allows an accurate prediction of the risk of lymph-node metastasis.²⁶ The risk of developing lymph-node or distant metastasis is then weighed against the risk of surgery.^{27–29}

Endoscopic resection should be safe, effective, and applicable to a wide variety of clinical situations. However, an ESD still requires an experienced endoscopist with a high level of skill, especially when using a single working-chan-

nel gastroscop. Recently, the technique of percutaneous traction-assisted EMR by using a laparoscopic port to create a strong counter-traction was reported.^{30–32} However, all previous trials showed that the technique was complicated, invasive, and did not make ESD easier.

Magnets and magnetic fields were used to direct the catheter tip during catheter procedures.³³ A magnetic anchoring system was used to achieve laparoscopic surgery by using a single trocar.³⁴ Very recently, the feasibility of using magnetically anchored instruments was reported as a promising technique to facilitate natural orifice transluminal endoscopic surgery in a porcine model.³⁵ These magnets may also provide a way to alter tissue contours without any direct contact. A direct-current-generated magnetic field, as used in magnetic resonance imaging, is regarded as the least invasive or even the most appropriate noninvasive procedure that can be medically applied.

In 21 of our patients, only one magnetic anchor was needed to achieve the desired result, either by rotating the patient or by moving the examination table. In 4 cases, 2 magnetic anchors were required. In 2 cases, a second magnetic anchor was helpful. With the other 2 cases, however, the second anchor did not help, because the MAG system did not provide adequate visualization for submucosal dissection or allow suitable maneuvering of the endoscopic devices. This was caused by underinflation of the gastric cavity. Therefore, to obtain better visualization during an MAG-ESD, the prevention of air leakage because of a hiatus hernia should be achieved.

Another limitation of this procedure was that the extracorporeal electromagnetic control system is too large and cumbersome. Although it was possible to achieve hands-free fixation of the mucosa by using the magnetic anchor tractioned with the extracorporeal electromagnet, the system should be miniaturized to allow wider clinical application.

In conclusion, this prospective clinical trial proved that MAG-ESD can feasibly be used in human beings. The MAG-ESD technique was able to obtain excellent visualization by suitable tissue tension and to facilitate the procedures. Further innovations are warranted to apply the MAG procedure in daily clinical practice.

ACKNOWLEDGMENTS

We thank Professor Stefan Seewald (Department of Interdisciplinary Endoscopy, University Medical Center Hamburg-Eppendorf, Hamburg, Germany) for his helpful suggestions for fluent medical English and encouragement with the preparation of the article.

REFERENCES

1. Ono H, Kondo H, Gotoda T, et al. Endoscopic mucosal resection for treatment of early gastric cancer. *Gut* 2001;48:225–9.

2. Ohkuwa M, Hosokawa K, Boku A, et al. New endoscopic treatment for intramucosal gastric tumors using an insulated-tip diathermic knife. *Endoscopy* 2001;33:221-6.
3. Hosokawa K, Yoshida S. Recent advances in endoscopic mucosal resection for early gastric cancer [Japanese with English abstract]. *Jpn J Cancer Chemother* 1998;25:476-83.
4. Gotoda T, Kondo H, Ono H, et al. A new endoscopic mucosal resection (EMR) procedure using an insulation-tipped diathermic (IT) knife for rectal flat lesions. *Gastrointest Endosc* 1999;50:560-3.
5. Oyama T, Kikuchi Y. Aggressive endoscopic mucosal resection in the upper GI tract: Hook knife EMR method. *Minim Invasive Ther Allied Technol* 2002;11:291-5.
6. Yahagi N, Fujishiro M, Kakushima N, et al. Endoscopic submucosal dissection for early gastric cancer using the tip of an electro-surgical snare (thin type). *Dig Endosc* 2004;16:34-8.
7. Yamamoto H, Kawata H, Sunada K, et al. Successful en bloc resection of large superficial tumors in the stomach and colon using sodium hyaluronate and small-caliber-tip transparent hood. *Endoscopy* 2003;35:690-4.
8. Oka S, Tanaka S, Kaneko I, et al. Advantage of endoscopic submucosal dissection compared with EMR for early gastric cancer. *Gastrointest Endosc* 2006;64:877-83.
9. Rösch T, Sarbia M, Schmacher B, et al. Attempted endoscopic en bloc resection of mucosal and submucosal tumors using insulated-tip knives: a pilot series. *Endoscopy* 2004;36:788-801.
10. Choi IJ, Kim CG, Chang HJ, et al. The learning curve for EMR with circumferential mucosal incision in treating intramucosal gastric cancer. *Gastrointest Endosc* 2005;62:860-5.
11. Gotoda T, Friedland S, Hamanaka H, et al. A learning curve for advanced endoscopic resection. *Gastrointest Endosc* 2005;62:866-7.
12. Matsushita M, Hajiro K, Okazaki K, et al. Endoscopic mucosal resection of gastric tumors located in the lesser curvature of the upper third of the stomach. *Gastrointest Endosc* 1997;45:512-5.
13. Kobayashi T, Gotoda T, Tamakawa K, et al. Magnetic anchor for more effective endoscopic mucosal resection. *Jpn J Clin Oncol* 2004;34:118-23.
14. Gotoda T. A large endoscopic resection by endoscopic submucosal dissection (ESD) procedure. *Clin Gastroenterol Hepatol* 2005;3:571-3.
15. Soetikno R, Gotoda T, Nakanishi Y, et al. Endoscopic mucosal resection. *Gastrointest Endosc* 2003;57:567-9.
16. Ludwig K, Klautke G, Bernhard J, et al. Minimally invasive and local treatment for mucosal early gastric cancer. *Surg Endosc* 2005;19:1362-6.
17. Deyhle P, Largiader F, Jenny P. A method for endoscopic electroresection of sessile colonic polyps. *Endoscopy* 1973;5:38-40.
18. Tada M, Murakami A, Karita M, et al. Endoscopic resection of early gastric cancer. *Endoscopy* 1993;25:445-51.
19. Gotoda T. Endoscopic resection of early gastric cancer: the Japanese perspective. *Curr Opin Gastroenterol* 2006;22:561-9.
20. Inoue H, Takeshita K, Hori H, et al. Endoscopic mucosal resection with a cap-fitted panendoscope for esophagus, stomach, and colon mucosal lesions. *Gastrointest Endosc* 1993;39:58-62.
21. Akiyama M, Ota M, Nakajima H, et al. Endoscopic mucosal resection of gastric neoplasms using a ligating device. *Gastrointest Endosc* 1997;45:182-6.
22. Korenaga D, Haraguchi M, Tsujitani S, et al. Clinicopathological features of mucosal carcinoma of the stomach with lymph node metastasis in eleven patients. *Br J Surg* 1986;73:431-3.
23. Ell C, May A, Gossner L, et al. Endoscopic mucosectomy of early cancer and high-grade dysplasia in Barrett's esophagus. *Gastroenterology* 2000;118:670-7.
24. Tanabe S, Koizumi W, Mitomi H, et al. Clinical outcome of endoscopic aspiration mucosectomy for early stage gastric cancer. *Gastrointest Endosc* 2002;56:708-13.
25. Eguchi T, Gotoda T, Oda I, et al. Is endoscopic one-piece mucosal resection essential for early gastric cancer? *Dig Endosc* 2003;15:113-6.
26. Gotoda T, Sasako M, Ono H, et al. An evaluation of the necessity of gastrectomy with lymph node dissection for patients with submucosal invasive gastric cancer. *Br J Surg* 2001;88:444-9.
27. Gotoda T, Yanagisawa A, Sasako M, et al. Incidence of lymph node metastasis from early gastric cancer: estimation with a large number of cases at two large centers. *Gastric Cancer* 2000;3:219-25.
28. Etoh T, Katai H, Fukagawa T, et al. Treatment of early gastric cancer in the elderly patient: results of EMR and gastrectomy at a national referral center in Japan. *Gastrointest Endosc* 2005;62:868-71.
29. Soetikno R, Kaltenbach T, Yeh R, et al. Endoscopic mucosal resection for early cancers of the upper gastrointestinal tract. *J Clin Oncol* 2005;23:4490-8.
30. Ohashi S. Laparoscopic intraluminal surgery for early gastric cancer: is it a new concept in laparoscopic intraluminal surgery. *Surg Endosc* 1995;9:169-71.
31. Ohgami M, Otani Y, Kumai K, et al. Curative laparoscopic surgery for early gastric cancer: five years experience. *World J Surg* 1999;23:187-93.
32. Kondo H, Gotoda T, Ono H, et al. Percutaneous traction-assisted EMR by using an insulation-tipped electro-surgical knife for early stage gastric cancer. *Gastrointest Endosc* 2004;59:284-8.
33. Faddis MN, Blume W, Finney J, et al. Novel, magnetically guided catheter for endocardial mapping and radiofrequency catheter ablation. *Circulation* 2002;106:2980-5.
34. Zeltser IS, Bergs R, Fernandez R, et al. Single trocar laparoscopic nephrectomy using magnetic anchoring and guidance system in the porcine model. *J Urol* 2007;178:288-91.
35. Scott DJ, Tang SJ, Fernandez R, et al. Completely transvaginal NOTES cholecystectomy using magnetically anchored instruments. *Surg Endosc* 2007;21:2308-16.

Received December 7, 2007. Accepted March 31, 2008.

Current affiliations: Endoscopy Division (T.G., I.O.), National Cancer Center Hospital, Tokyo, Tamakawa Corporation (K.T.), Sendai, Pentax Corporation (H.U.), Tokyo, Cancer Screening Technology Division (T. Kobayashi), Research Center for Cancer Prevention and Screening, National Cancer Center (T. Kakizoe), Tokyo, Japan.

Reprint requests: Takuji Gotoda, MD, Endoscopy Division, National Cancer Center Hospital, 5-1-1 Tsukiji, Chuo-ku, Tokyo 104-0045, Japan.

If you want to chat with an author of this article, you may contact him at tgotoda@ncc.go.jp.