Usefulness of dilated blood vessels in the tumor periphery for assessing the invasion depth of small-sized depressed colorectal cancer

Rintaro Hashimoto (MD)$^\dagger$, Tomoki (MD), Hidetaka Hamamoto (MD), Hajime Yamaoka (MD), Masato Nakahori (MD), Akimichi Chonan (MD)

Abstract
The relationship between dilated blood vessels in the tumor periphery and the tumor invasion depth is unclear. Therefore, the present study aimed to clarify the relationship between dilated blood vessels and the invasion depth of small-sized (<30 mm) colorectal cancer (CRC), and its implications on endoscopic treatment.

We performed a single-arm observational study of the diagnostic accuracy of the existence of dilated vessels in the tumor periphery of CRC lesions as an indicator of submucosal deep (SM-d, ≥1000 μm) carcinomas. Lesions were classified into two groups based on the existence of dilated vessels by two experienced endoscopists. The clinicopathological features, invasion depth, and lymphovascular invasion/poorly differentiated clusters were analyzed in all resected specimens.

Four hundred and two consecutive small-sized CRC lesions were included. The dilated vessels were observed in 96/402 (24%) lesions, and most of them (93/96) were found in depressed lesions. In depressed lesions, the histopathological diagnosis of the dilated vessels group showed SM-d or deeper invasion in 84/93 (90%) cases, whereas 3/20 (15%) had SM-d invasion in the nondilated vessels group ($P < 0.001$). When the dilated vessels were used as an indicator of SM-d or deeper invasion in depressed lesions, the sensitivity was 95.6%, specificity was 66.7%, and accuracy was 90.2%. No correlation was observed between the existence of dilated vessels and the lesion site, lesion diameter, and lymphovascular invasion/poorly differentiated cluster.

The existence of dilated blood vessels in the tumor periphery suggests SM-d or deeper invasion in depressed lesions.

Abbreviations: CI = confidence interval, CRC = colorectal cancer, LVI = lymphovascular invasion, MP = muscularis propria, NBI = narrow-band imaging, NICE classification = Narrow-band Imaging International Colorectal Endoscopic classification, PDC = poorly differentiated clusters, SM-d = submucosal deep, SM-s = submucosal slight.

Keywords: chromoendoscopy, colonoscopy, colorectal cancer, endoscopic resection, sensitivity, specificity

1. Introduction
It is very important to accurately estimate the depth of invasion of early stage colorectal cancer (CRC) to make proper therapeutic decisions because patients with intramucosal carcinoma and submucosal invasive carcinoma with an invasion depth of less than 1000 μm can be cured by endoscopic resection.$^{[1]}$

Chromoendoscopy, using Kudo and Tsuruta’s pit pattern classification, is a useful tool for making the differential diagnosis of colorectal tumors, including submucosal carcinoma.$^{[2,3]}$

Magnifying chromoendoscopy has been widely demonstrated to be effective in differentiating between colorectal neoplastic and non-neoplastic polyps, and assessing the invasion depth by using pit pattern analysis.$^{[4]}$

Some studies have shown the effectiveness of narrow-band imaging (NBI) magnifying endoscopy for determining the invasion depth.$^{[5]}$ Recently, the Narrow-band Imaging International Colorectal Endoscopic classification, which requires diagnoses of the vascular pattern and surface pattern, has also been used.$^{[6,7]}$ As vascular diagnosis is not included in the diagnostic criteria for pathological grading, its relationship with the depth of tumor invasion is indirect. For colorectal intratumoral vessels, as the pathological grading increases, a larger blood vessel diameter and varying density and irregularity on a scale are observed with the depth of tumor invasion.$^{[8]}$ However, the relationship between blood vessels in the tumor periphery and the tumor invasion depth is unclear.

2. Methods

2.1. Study design
This study was a single-arm observational study of diagnostic accuracy according to the Standards for the Reporting of Diagnostic Accuracy Studies initiative,$^{[9]}$ which was conducted at Sendai Kousei Hospital, a tertiary referral hospital. The aim of this study was to clarify the relationship between the existence of dilated blood vessels in the tumor periphery and the invasion depth of small-sized CRC. The protocol of this retrospective study was approved by our Institutional Review Board.

2.2. Patients
This study included consecutive endoscopically or surgically resected 774 CRC lesions, of which the invasion depth was...
pathologically diagnosed as intramucosal (M), submucosal (SM), or muscularis propria (MP), in 652 patients at Sendai Kousei Hospital between January 2012 and November 2014. To assess the existence of dilated blood vessels in the tumor periphery correctly, we excluded lesions (1) without any endoscopic images at our hospital (12 lesions), (2) larger than 3 cm in diameter (184 lesions), and (3) without sufficient endoscopic images for judging the existence of dilated blood vessels by two experienced endoscopists (H.Y. and M.H.), who have each performed more than 5000 colonoscopies (165 lesions).

2.3. Endoscopic procedures

We used magnifying colonoscopies (PCF-Q 240ZI and PCF-Q260AZI, Olympus Co., Tokyo, Japan), attached to a processor (CV-260SL, Olympus Co.) and a light source (CLV-260SL, Olympus Co.) to assess the lesions in all cases. There were no adverse events in all cases.

2.4. Definitions

We defined the dilated blood vessels in the tumor periphery as the vessels thicker than those surrounding a type I pit that travels through two or more ducts over a type I pit (Fig. 1).

2.5. Evaluation

All lesions were classified into two groups according to the existence of dilated blood vessels in the tumor periphery. Furthermore, the gross type was classified as protruded (0-Ip, IsP, and Is), flat elevated (0-IIa), and depressed (0-IIc, IIa+IIc).[10] Clinicopathological features, such as the depth of invasion, lymphovascular invasion (LVI), and poorly differentiated clusters (PDCs) were analyzed in all resected specimens based on the World Health Organization’s criteria.[11] We measured the submucosal invasion depth according to the guidelines issued in 2014 by the Japanese Society for Cancer of the Colon and Rectum for treating CRC.[12] All lesions were independently assessed by two other experienced endoscopists (T.M. and H.H.), who have each performed more than 5000 colonoscopies. They were blinded to each pathological diagnosis, and they evaluated the existence of dilated blood vessels of the lesions from endoscopic images. When they did not agree, the existence of dilated blood vessels was regarded as negative.

2.6. Statistical methods

Data were analyzed by JMP Pro statistical software for Windows, version 11 (SAS, Tokyo, Japan). Continuous data were compared using unpaired Student’s t-tests, whereas categorical variables were tested using Mann-Whitney U test. Values of $P < 0.05$ (two-tailed) were considered significant. The diagnostic accuracy was expressed as point estimates of accuracy rate with Wald-type 95% confidence interval (CI).

3. Results

A total of 402 consecutive small-sized CRC lesions in 380 patients were included. The invasion depth of the lesions was as follows: M 227, SM-s 38, SM-d 98, and MP 39. The dilated vessels were observed in 96 of 402 lesions (Fig. 2), but most of them were recognized in depressed lesions (Table 1). Thus, we evaluated the relationship between the existence of dilated vessels and age, location, invasion depth, and tumor size of the depressed lesion, and we found that the existence of dilated vessels and the tumor depth are closely related (Table 2). The existence of dilated vessels suggests SM invasion, especially SM-d or deeper invasion, in depressed lesions. Thus, the existence of dilated vessels may be

![Figure 1. Dilated vessels (arrow) seen in the tumor periphery.](image-url)
accuracy was 90.2% (83.1, 95.0) (Table 4). The positive/negative diagnostic odds ratio was 43.5 (range 11.3–96 lesions), and the concordance rate between the two endoscopists was 98.2% (94/95, 95% CI: 89.1, 98.8), specificity was 95.6% (83.1, 95.0) (Table 3). The diagnostic accuracy and inter-observer concordance of both classifications are not so high. The dilated vessels evaluated in the current study were easier to observe than observation with blue laser imaging magnification has been also used to determine the invasion depth of colorectal neoplasms and the diagnostic effectiveness of this method was similar to that of NBI magnification.[13] Magnifying endoscopy requires close observation of the tumor surface’s blood vessels and structure, which can sometimes be difficult to observe in real-world clinical settings due to bleeding or intestinal tract washing liquid residues. In addition, the diagnostic accuracy and inter-observer concordance of both classifications are not so high.[14,15] The dilated vessels evaluated in the current study were easier to observe than the indicators that have been used thus far. The dilated vessels are a strong indicator of SM-d or deeper lesions (Table 3). The concordance rate between the two endoscopists was 98.2% (94/96 lesions), and the \( \kappa \) value was 0.93.

When dilated vessels were used as an indicator of SM-d or deeper invasion in depressed lesions, the sensitivity was 95.6% (95% CI: 89.1, 98.8), specificity was 90.2% (83.1, 95.0) (Table 4). The positive/negative likelihood ratio was 2.87/0.07 (range 1.56–5.26/0.02–0.18). The diagnostic odds ratio was 43.5 (range 11.3–168).

Histopathological analysis of the dilated vessels positive group showed submucosal cancer in 57 (39%) cases, LVI in 29 (51%), and PDC in 5 (8%). Similarly, the dilated vessels negative group had submucosal cancer in 79 (26%) cases, LVI in 33 (41%), and PDC in 12 (15%). Therefore, in relation to the existence of dilated vessels, LVI, and PDC, there were no significant differences between the groups (Table 5).

### 4. Discussion

Magnifying chromoendoscopy using pit pattern classification is considered the most accurate method for determining of the depth of invasion of early-stage CRC.[4] The NBI classification can be easily used with or without magnifying endoscopy, and it has been advocated in recent years.[15–17] Lately, narrow-band light observation with blue laser imaging magnification has been also used to determine the invasion depth of colorectal neoplasms and the diagnostic effectiveness of this method was similar to that of NBI magnification.[13] Magnifying endoscopy requires close observation of the tumor surface’s blood vessels and structure, which can sometimes be difficult to observe in real-world clinical settings due to bleeding or intestinal tract washing liquid residues. In addition, the diagnostic accuracy and inter-observer concordance of both classifications are not so high.[14,15] The dilated vessels evaluated in the current study were easier to observe than the indicators that have been used thus far. The dilated vessels are also useful if obtaining a frontal internal view of the lesions is impossible or bowel preparation is insufficient.

Depressed lesions have a tendency to rapidly invade the submucosal layer, even when they are small. Oka et al reported that the prevalence of SM invasion in depressed lesions is about

![Figure 2: STARD diagram showing the number of enrolled lesions in this study.](image-url)

### Table 1

| Lesions (n=402) | Dilated vessels + (n=96) | Dilated vessels – (n=306) | P value |
|----------------|-------------------------|---------------------------|---------|
| Sex (M/F)      | 57/39                   | 179/127                   | 0.89    |
| Age            | 64.5 (38–85)            | 66.8 (38–89)              | 0.52    |
| Location       |                         |                           |         |
| Right colon    | 32                      | 118                       | 0.30    |
| Left colon     | 35                      | 111                       |         |
| Rectum         | 29                      | 77                        |         |
| Macroscopic typea |                   |                           |         |
| Protruded      | 2                      | 257                       | <0.001  |
| Flat elevated  | 0                      | 31                        |         |
| Depressed      | 94                      | 18                        |         |
| Size (mm)      |                         |                           |         |
| <5             | 5                      | 2                         | 0.13    |
| 6–10           | 26                     | 35                        |         |
| 11–20          | 48                     | 246                       |         |
| 21–30          | 17                     | 23                        |         |

Depressed = IIc/IIa+IIc, F = female, flat elevated = IIa, M = male, protruded = IIb/Iic/Iic.  

*a* Borrman Type I and II are included in “protruded” and “depressed,” respectively.

### Table 2

| Lesions (n=112) | Dilated vessels + (n=94) | Dilated vessels – (n=18) | P value |
|----------------|-------------------------|--------------------------|---------|
| Gender (M/F)   | 56/38                   | 10/8                     | 0.84    |
| Sex            | 64.6 (38–85)            | 62.1 (40–88)             | 0.45    |
| Location       |                         |                           |         |
| Right colon    | 31                      | 5                         | 0.73    |
| Left colon     | 35                      | 10                        |         |
| Rectum         | 28                      | 3                         |         |
| Depth          |                         |                           |         |
| M              | 2                       | 8                         | <0.001  |
| SM-s           | 5                       | 6                         |         |
| SM-d           | 52                      | 4                         |         |
| MP             | 35                      | 0                         |         |
| Size (mm)      |                         |                           |         |
| <5             | 4                       | 2                         | 0.61    |
| 6–10           | 25                      | 4                         |         |
| 11–20          | 50                      | 11                        |         |
| 21–30          | 15                      | 1                         |         |

F = female, M = male, M = muscularis propria, SM-d = submucosal deep, SM-s = submucosal slight.

### Table 3

| Lesions (n=112) | Dilated vessels + (n=94) | Dilated vessels – (n=18) | P value |
|----------------|-------------------------|--------------------------|---------|
| M/SM-s         | 7                       | 14                       | <0.001  |
| SM-d/MP        | 87                      | 4                        |         |

M/SM-s = mucosal/submucosal slight, SM-d/MP = submucosal deep/muscularis propria.
the blood vessel diameter in colorectal tumors is generally thicker unclear. Still, it is surmised that dilated vessels may originate from invasion in depressed lesions, even if the tumor size is small. Thus, this information may help physicians strongly suspect deep dilated vessels in depressed lesions, although the existence was no correlation between the tumor size and existence of dilated vessels in depressed lesions. Thus, we still consider dilated vessels the possible usefulness of recognizing dilated vessels in the tumor to the center for pit-pattern diagnosis. However, our data suggest the possible usefulness of recognizing dilated vessels in the tumor periphery, which was not well understood until now.

In conclusion, the existence of dilated vessels in the tumor periphery suggests SM invasion, especially SM-d or deeper invasion, in depressed lesions. This information may help physicians decide treatment and encourage careful management of depressed lesions.

References

[1] Watanabe T, Itabashi M, Shimada Y, et al. Japanese Society for Cancer of the Colon and Rectum (JSCCR) guidelines 2010 for the treatment of colorectal cancer. Int J Clin Oncol 2012;17:1-29.

[2] Kudo S, Hirota S, Nakajima T, et al. Colorectal tumours and pit pattern. J Clin Pathol 1999;47:880-5.

[3] Fu KI, Sano Y, Kato S, et al. Chromoendoscopy using indigo carmine dye spraying with magnifying observation is the most reliable method for differential diagnosis between non-neoplastic and neoplastic colorectal lesions: a prospective study. Endoscopy 2004;36:1089-3.

[4] Matsuda T, Fujii T, Saito Y, et al. Efficacy of the invasive/non-invasive pattern by magnifying chromoendoscopy to estimate the depth of invasion of early colorectal neoplasms. Am J Gastroenterol 2008;103: 2700–6.

[5] Kanao H, Tanaka S, Oka S, et al. Narrow-band imaging magnification predicts the histology and invasion depth of colorectal tumors. Gastrointest Endosc 2009;69:631.

[6] Hewett DG, Kaltenbach T, Sano Y, et al. Validation of a simple classification system for endoscopic diagnosis of small colorectal polyps using narrow-band imaging. Gastroenterology 2012;143: 599–607.

[7] Hayashi N, Tanaka S, Hewett DG, et al. Endoscopic prediction of deep submucosal invasive carcinoma: validation of the narrow-band imaging international colorectal endoscopic (NICE) classification. Gastrointest Endosc 2013;78:625–32.

[8] Fukuzawa M, Saito Y, Matsuda T, et al. Effectiveness of narrow-band imaging magnification for invasion depth in early colorectal cancer. World J Gastroenterol 2010;16:1727–34.

[9] Bossuyt PM, Reitsma JB, Bruns DE, et al. The STARD statement for reporting studies of diagnostic accuracy: explanation and elaboration. BMJ 2015;351:h5527.

[10] Endoscopic Classification Review Group. Update on the Paris classification of superficial neoplastic lesions in the digestive tract. Endoscopy 2005;37:570–8.

[11] Hamilton SR, Aaltoinen LA, editors. World Health Organization Classification Of Tumors. Pathology and Genetics of Tumours of the Digestive System. Lyon, France: IARC Press; 2010. pp. 104-109

[12] Japanese Society for Cancer of the Colon and RectumJapanes guidelines for the treatment of colorectal carcinoma. 2014; Tokyo: Kanehara Shuppan Co, 48–52.

[13] Yoshida N, Hisabe T, Inada Y, et al. The ability of a novel blue laser imaging system for the diagnosis of invasion depth of colorectal neoplasms. J Gastroenterol 2014;49:73–80.

[14] Sakamoto T, Saito Y, Nakajima T, et al. Comparison of magnifying chromoendoscopy and narrow-band imaging in estimation of early colorectal cancer invasion depth: a pilot study. Dig Endosc 2011;23:118–23.

[15] Zhang JJ, Gu LY, Chen XY, et al. Endoscopic diagnosis of invasion depth for early colorectal carcinomas: a prospective comparative study of narrow-band imaging, acetic acid, and crystal violet. Medicine (Baltimore) 2015;94:e328.

[16] Oka S, Tanaka S, Nakados A, et al. Endoscopic features and management of diminutive colorectal submucosal invasive carcinoma. Dig Endosc 2014;26:78–3.

[17] Konoedranga MA, Fiet E, Gaumann A. 3D microvascular architecture of pre-cancerous lesions and invasive carcinomas of the colon. Br J Cancer 2001;84:1354–62.

[18] Kashida H, Kudo SE. Early colorectal cancer: concept, diagnosis, and management. Int J Clin Oncol 2006;11:1–8.