

Depressed-type early adenocarcinoma of the terminal ileum

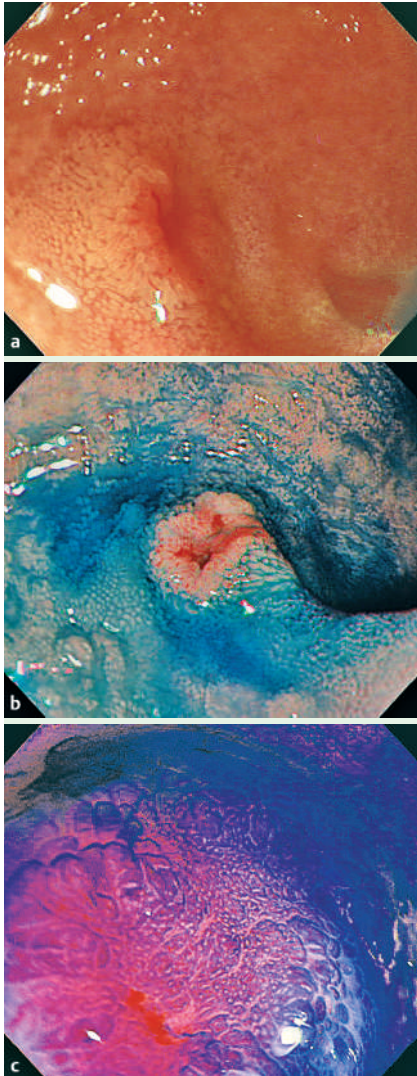


Fig. 1 **a** Conventional endoscopic view showed a shallow depressed lesion in the terminal ileum. **b** Chromoendoscopic view with indigo carmine dye showed a star-shaped demarcation line of the depressed lesion. **c** Magnifying endoscopic view with crystal violet staining showed regular arrangement of small round and tubular pit patterns.

Video 1

Video clip shows conventional and magnifying endoscopic features of depressed type adenocarcinoma of the terminal ileum, and procedure of endoscopic mucosal resection.

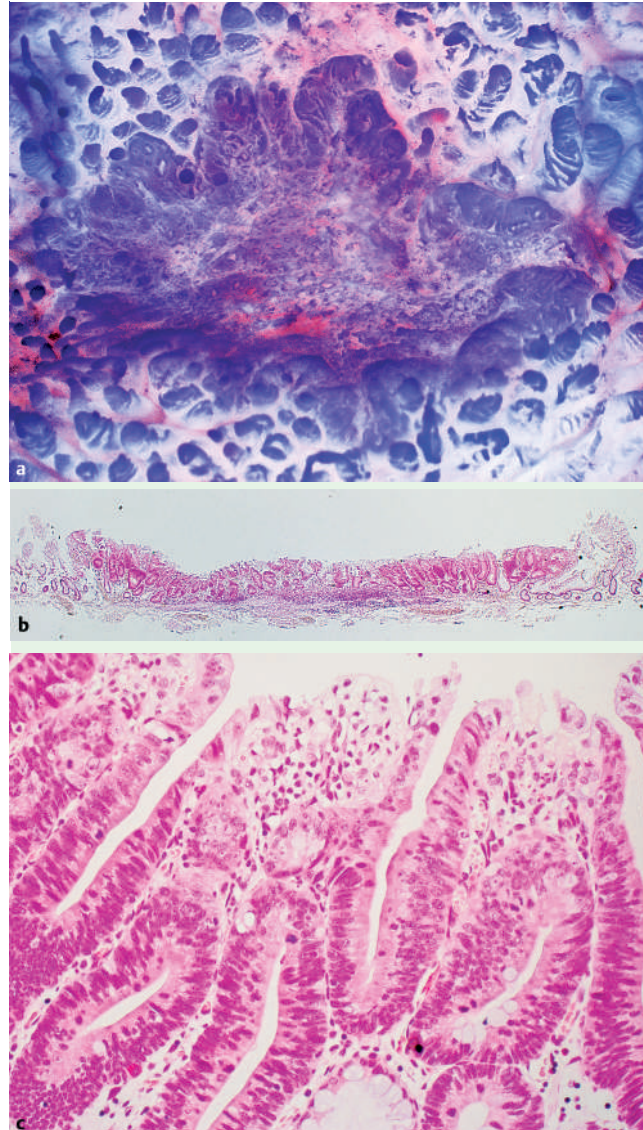


Fig. 2 **a** Stereomicroscopic view showed a star-shaped depressed lesion with marginal elevation. **b** Histopathologic cross section revealed an intramucosal depressed neoplasm with lamina propria invasion (H & E, original magnification $\times 10$). **c** High-power view revealed well-differentiated adenocarcinoma (H&E, original magnification $\times 200$).

Depressed-type neoplasm of the jejunum and the ileum has not yet been recognized. To the best of our knowledge, this is the first case of depressed-type primary adenocarcinoma of the ileum. A 62-year-old man underwent total colonoscopy. A small reddish depressed lesion with marginal elevation, 5 mm in diameter, was detected in the terminal ileum (▶ Fig. 1 a). A chromoendoscopic view with indigo carmine dye showed a star-shaped demarcation line of the depressed lesion (▶ Fig. 1 b). A magnifying endoscopic view with crystal violet stain-

ing showed small round and tubular pit patterns (▶ Fig. 1 c). With these findings, we diagnosed intramucosal neoplasm. An endoscopic mucosal resection was performed, and the resected lesion measured 5 × 4 mm. Stereomicroscopic view showed a star-shaped, depressed lesion with marginal elevation (▶ Fig. 2 a). A histopathologic cross section revealed an intramucosal depressed neoplasm with lamina propria invasion (▶ Fig. 2 b). A high-power view revealed well-differentiated adenocarcinoma (▶ Fig. 2 c). Cancer-cell nuclei were positive in p53 im-

munohistochemical staining. A polymerase chain reaction-single strand conformation polymorphism (PCR-SSCP) study of p53 genes revealed some mutations of exon 6, 7, and 8. K-ras codon 12 mutations (PCR-restriction fragment length polymorphism [RFLP]) were not observed. Morphological appearance was classified as type 0-IIc in the Paris endoscopic classification [1], and mimicked a depressed-type colorectal cancer advocated by Kudo [2]. It had been reported that K-ras mutations were absent in depressed-type colorectal cancers [3,4]. It was reported that rates of p53 positivity in depressed-type colorectal neoplastic lesions were higher in carcinomas and high-grade neoplasms than in low-grade neoplasms [5]. Therefore the characterizations of the genetic change, such as p53 and K-ras, were mimicking depressed-type colorectal cancer.

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References

- 1 The Paris endoscopic classification of superficial neoplastic lesions. Esophagus, stomach and colon: November 30 to December 1, 2002. *Gastrointest Endosc* 2003; 58: S3–S43
- 2 Kudo S. Endoscopic mucosal resection of flat and depressed type of early colorectal cancer. *Endoscopy* 1993; 25: 455–461
- 3 Saito K, Arai K, Mori M. p53 overexpression and K-ras codon 12 mutations in submucosal invasive depressed-type colorectal cancer. *Oncol Rep* 2000; 7: 741–744
- 4 Kaneko K, Kurahashi T, Makino R *et al*. Pathological features and genetic alterations in colorectal carcinomas with characteristics of nonpolypoid growth. *Br J Cancer* 2004; 91: 312–318
- 5 Hayakawa M, Shimokawa K, Kusugami K *et al*. Clinicopathological features of superficial depressed-type colorectal neoplastic lesions. *Am J Gastroenterol* 1999; 94: 944–949

Bibliography

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