An 81-year-old man was admitted to the hospital with hematochezia and ascites, but with no abdominal pain or skin rash. Colonoscopy revealed a circumferential ulcer with marked edema in the terminal ileum (Figure 1a). Duodenoscopy also showed multiple ulcers in the duodenum (Figure 1b). Histological examination of biopsy specimens indicated nonspecific inflammation. Paracentesis yielded hemorrhagic fluid, with no malignant cells and no growth of pathogens. A definite diagnosis could not be established initially, and the patient was treated conservatively.

On hospital day 8, palpable purpuric lesions abruptly developed at both extremities, hematochezia progressed, and renal function rapidly worsened. On the basis of these findings, Henoch-Schönlein purpura was diagnosed clinically, and was pathologically confirmed several days later (Figure 2). Despite intensive treatment, including methylprednisolone pulse therapy, the patient’s condition progressively deteriorated and he died on day 13.

Autopsy revealed massive hemorrhagic ascites and coagulated blood in the gastrointestinal tract. The mucosa in both the duodenum and the ileum was extensively peeling, but interestingly, the jejunal and colonic mucosa was intact (Figure 3). Immunohistochemistry demonstrated IgA depositions on the small vessels of the bowel wall, including not only the duodenum and the ileum, with extensive ulcers, but also the jejunum, with a normal appearance.

The presence of characteristic purpura can facilitate the diagnosis of Henoch-Schönlein purpura, but the diagnosis is more difficult when other manifestations of the disease precede the purpura [1–4]. In the present patient, it was difficult to obtain a final diagnosis of Henoch-Schönlein purpura, as the case had the following unusual in characteristics: old age at onset, delayed appearance of characteristic purpura.
Henoch–Schönlein purpura, an absence of abdominal pain, and the presence of hemorrhagic ascites. The patient also had an unusual distribution of gastrointestinal lesions, with the jejunum being spared despite systemic vasculitis. However, the reason for the absence of the jejunal lesions is not clear.

References


Corresponding author

S. Yazumi, M.D., Ph.D.
Dept. of Gastroenterology and Hepatology
Kyoto University
Graduate School of Medicine
Shogoin-Kawara-cho, Sakyo-ku
Kyoto 606-8507
Japan
Fax: +81-75-751-4303
Email: yazoo@kuhp.kyoto-u.ac.jp