Cholesterol embolization syndrome (CES) is a rare syndrome with a high incidence of morbidity and mortality [1]. Because of the variability in clinical presentations, the diagnosis is often difficult. CES presents as multiorgan dysfunction and should be considered in the differential diagnosis of the symptom complex after vascular thrombolytic therapy [2]. Skin and renal involvement are the most common presentations, such as livedo reticularis and rash [3]. CES of the digestive systems is not uncommon, with rates varying from 18% to 48% [4, 5]. We encountered a rare case of rectal perforation due to CES, which was diagnosed only after the patient had died.

A 91-year-old woman with rectal bleeding was admitted for shunt formation and hemodialysis for progressive renal failure. She suddenly had massive melena, and computed tomography showed rectal perforation, which was also suspected based on the communicating irregular air density in adipose tissue on the left side of the rectum (Fig. 1)

Colonoscopy revealed bleeding, perforation, and fistula formation in the rectum (Fig. 2).

The rectal perforation was limited to the surrounding intrapelvic adipose tissue, and surgical treatment was planned, but the patient died of renal complications and disseminated intravascular coagulation. On pathologic examination, a 9 mm × 5 mm ulcer was found in the lower rectum, and at the base of the ulcer, arteriolar cholesterol emboli with cholesterol clefts were observed along with granulation tissue (Fig. 3).

Cholesterol emboli and clefts were also found in the spleen and renal arterioles. From these pathologic findings, a rectal perforation resulting from CES was diagnosed.

The diagnosis of CES can be confirmed by means of biopsy of the target organs [3]; the biopsy sections of the lesion areas demonstrate cholesterol emboli and clefts. A thorough clinical history and physical findings are important steps in establishing a diagnosis. A correct diagnosis is important because it may prevent inappropriate treatment of the patient [3–5].

Competing interests: None

H. Mori1, H. Kobara1, M. Kobayashi1, A. Muramatsu1, T. Nomura1, T. Yachida1, K. Izuiti2, Y. Suzuki2, J. Gong1, T. Masaki1

1 Department of Gastroenterology and Neurology, Kagawa Medical University School of Medicine, Kagawa, Japan

2 Department of Gastroenterological Surgery, Kagawa Medical University School of Medicine, Kagawa, Japan
References


Bibliography

Endoscopy 2010; 42: E352–E353
© Georg Thieme Verlag KG Stuttgart · New York · ISSN 0013-726X

Corresponding author

H. Mori
Department of Gastroenterology and Neurology
Kagawa University School of Medicine
1750–1 Ikenobe
Miki Kita Kagawa 761–0793
Japan
Fax: +81-87-891-2158
hiro4884@med.kagawa-u.ac.jp