Successful treatment of Cronkhite–Canada syndrome using anti-tumor necrosis factor antibody therapy

Cronkhite–Canada syndrome (CCS) is a rare nonhereditary syndrome characterized by gastrointestinal polyposis and ectodermal changes [1]. Although several treatments, such as steroids, are available, the prognosis is poor, with a 5-year mortality rate of 55% [2].

A 57-year-old man, who suffered from alopecia, anorexia, severe diarrhea, dystrophic nail changes, and pigmentation on the hands, presented to our hospital. Colonoscopy revealed numerous polyps throughout the colon (● Fig. 1a). Gastroscopy and small-bowel series also showed severe gastric and small-bowel polyposis, and histology showed juvenile-like polyps with mild inflammation. On the basis of these findings, the patient was diagnosed with CCS.

He was treated with medications including corticosteroids, antiplasmin agents, and azathioprine; however, the symptoms persisted. He developed intussusception 7 years after diagnosis (● Fig. 1b) and underwent ileocecal resection. As he continued to suffer from frequent relapses, with diarrhea and malnutrition with severe hypoalbuminemia, a different therapeutic strategy was needed to control the disease progression. Supported by a single report finding of high levels of tumor necrosis factor alpha (TNF-α) in tissue affected by CCS [3], implying that anti-TNF-α therapy could be useful for CCS patients, the patient was administered infliximab at 200 mg every 2 weeks. His general condition improved, and colonoscopy performed 20 months after induction of anti-TNF therapy showed complete remission with the disappearance of polyposis (● Fig. 1c). He now receives 200 mg of infliximab every 2 weeks, and has been symptom free for 3 years since the initial administration.

There is one report that described strong TNF-α expression in intestinal mucosa affected by CCS, implicating the potential usefulness of anti-TNF-α antibody for patients with CCS. However, to the best of our knowledge, this treatment has not been tested until now [3]. Although we could not detect TNF-α expression in the polyps of this patient by immunohistochemistry (● Fig. 2), given the patient’s dramatic response to therapy, it can be assumed that TNF plays an important role in disease development, perhaps upstream of polyp formation. This case indicates that further basic and clinical research is warranted, and that anti-TNF therapy could be an important new strategy for treatment of CCS.

Competing interests: None

Daisuke Watanabe 1, Makoto Ooi 1, Namiko Hoshi 1, Michitaka Kohashi 1, Tomoo Yoshie 1, Nobunao Ikehara 1, Masaru Yoshida 2, Emmy Yanagita 1, Takashi Yamasaki 1, Tomoo Itoh 1, Takeshi Azuma 1

1 Division of Gastroenterology, Department of Internal Medicine, Kobe University Graduate School of Medicine, Kobe, Japan
2 Division of Metabolomics Research, Department of Internal Medicine, Kobe University Graduate School of Medicine, Kobe, Japan
3 Division of Diagnostic Pathology, Department of Pathology, Kobe University Graduate School of Medicine, Kobe, Japan

Fig. 1 Colonoscopy in a patient with Cronkhite–Canada syndrome. a Before initial treatment. b 3 days before ileocecal resection. c 20 months after anti-tumor necrosis factor alpha therapy.

Fig. 2 Immunohistochemistry for tumor necrosis factor alpha (TNF-α). Colon tissue of patient with Cronkhite–Canada syndrome was stained with: a anti-TNF-α antibody; b isotype control.
References


Bibliography

DOI http://dx.doi.org/10.1055/s-0034-1377539
Endoscopy 2014; 46: E476–E477
© Georg Thieme Verlag KG
Stuttgart · New York
ISSN 0013-726X

Corresponding author
Takeshi Azuma, MD, PhD
Division of Gastroenterology
Department of Internal Medicine
Kobe University
7-5-1, Kusunoki-cho
Chuo-ku, Kobe 650-0017
Japan
Fax: +81-78-3826309
azumat@med.kobe-u.ac.jp