Antibody to eosinophil cationic protein suppresses dextran sulfate sodium-induced colitis in rats

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AIM: To produce an antibody against rat eosinophil cationic protein (ECP) and to examine the effects of the antibody in rats with dextran sulfate sodium (DSS)-induced colitis.

METHODS: An antibody was raised against rat ECP. Rats were treated with 3% DSS in drinking water for 7 d and received the antibody or normal serum. The colons were examined histologically and correlated with clinical symptoms. Immunohistochemistry and Western blot analysis were estimated as a grade of inflammation.

RESULTS: The ECP antibody stained the activated eosinophils around the injured crypts in the colon mucosa. Antibody treatment reduced the severity of colonic ulceration and acute clinical symptoms (diarrhea and/or blood-stained stool). Body weight gain was significantly greater and the colon length was significantly longer in anti-ECP-treated rats than in normal serum-treated rats. Expression of ECP in activated eosinophils was associated with the presence of erosions and inflammation. The number of Ki-67-positive cells in the regenerated surface epithelium increased in anti-ECP-treated rats compared with normal serum-treated rats. Western blot analysis revealed reduced expression of macrophage migration inhibitory factor (MIF) in anti-ECP-treated rats.

CONCLUSION: Our results indicate that treatment with ECP antibody, improved DSS-induced colitis in rats, possibly by increasing the regenerative activity of the colonic epithelium and downregulation of the immune response, and suggest that anti-ECP may promote intestinal wound healing in patients with ulcerative colitis (UC).

Key words: Ulcerative colitis; Eosinophil cationic protein; Dextran sulfate sodium

INTRODUCTION

Eosinophil accumulation in the gastrointestinal tract is a common feature of numerous gastrointestinal disorders, including classic IgE-mediated food allergy, eosinophilic gastroenteritis, allergic colitis, eosinophilic esophagitis, inflammatory bowel disease (IBD) and gastroesophageal reflux disease. In IBD, eosinophils usually represent only a small percentage of the infiltrating leukocytes but their levels has been proposed as a negative prognostic indicator. Several studies have found an association between allergic colitis and later development of IBD, but this association is controversial.

Eosinophils and one of their granule proteins, eosinophil cationic protein (ECP) and eosinophil protein X (EPX), are generally recognized as being involved in the host defense against invading parasites. They are markedly cationic proteins with cytotoxic capacities that can potentially cause tissue destruction and could act as modulators of immune response. The eosinophil may also be involved in the pathogenesis of IBD because we have already reported activation of eosinophils in patients with active ulcerative colitis (UC), using the techniques of indirect immunoenzymatic method and electron microscopic examination of eosinophils, measurement of serum ECP, as well as increased percentages of hypodense eosinophils in the peripheral blood. Moreover, bowel biopsies from patients with IBD have demonstrated infiltration of eosinophils in the lamina

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propria and marked extracellular deposits of ECP\(^{[11,13]}\). There is also an excess release of the eosinophil proteins ECP and EPX in the luminal fluid and fecal material of patients with UC or Crohn’s disease\(^{[15]}\). However, the pathophysiological role played by activated eosinophils in the inflammatory process in UC has not yet been elucidated.

The aim of this study was to produce an antibody against rat ECP and to examine the antibody in dextran sulfate sodium (DSS)-induced colitis model. Rats treated with DSS develop severe colorectal damage mimicking IBD in humans\(^{[14,15]}\). We herein present evidence for the effectiveness of ECP antibody against DSS-induced colitis in rats, possibly by increasing the regenerative activity of the colonic epithelium and downregulation of the immune response.

MATERIALS AND METHODS

**Antibody raised against rat ECP**

TRAQWFA1QHISLNPPR\(^{[16]}\) was synthesized based on human ECP amino acid sequence\(^{[17]}\), which has homology to rat ECP\(^{[18]}\). Anti-ECP sera were generated by immunizing New Zealand white rabbits. In brief, rabbits were inoculated intradermally with 100 µg of ECP diluted in complete Freund’s adjuvant at weeks 2, 4, 6, 8, 10, 12, and 14. The serum was prepared according to the protocol provided by the manufacturer.

**Induction of experimental colitis**

Male Wistar rats/IZM (8 wk old) were obtained from Charles River Japan, Inc. Rats were kept in a specific pathogen-free environment at the Animal Center in accordance with the rules and regulations of the Institutional Animal Care and Use Committee of Nagasaki University. Food as well as drinking water with or without DSS (MW 5 000; Wako Pure Chemical Industries, Osaka, Japan) were provided ad libitum. The rats were treated with 3% DSS in drinking water for 7 d\(^{[19]}\), and then received either antibodies or normal serum intraperitoneally. The rats were weighed daily and visually inspected for gross rectal bleeding and diarrhea. The colons were examined histologically and correlated with clinical symptoms. Hematoxylin and cosin (HE)-stained sections were prepared from the distal colon. The severity of ulceration was quantified by the Ulcer Index (ulcer length/circumference length, %) in the distal colon. Furthermore, the severity of colitis was evaluated by assessment of colon length and histological examination. Similarly, we evaluated the effect of anti-ECP antibodies with regard to clinical signs and pathologic features.

**Administration of antibodies specific for ECP**

The polyclonal ECP antibody (0.25 mL/rat at d 0 and 1, 0.5 mL/rat at d 2, 3, 4, 5, 6, and 7 after DSS treatment) or non-immune rabbit serum was injected intraperitoneally.

**Immunohistochemical studies**

Anti-ECP antibody as eosinophils, anti-ED1 antibody (Serotec, Oxford, UK) as activated macrophages, anti-Ki-67 antibody (MIB-5, Dako Cytomation Denmark A/S, Denmark) as proliferative cells were used. The sections were deparaffinized and rehydrated. Then, they were microwaved for 10 min in 0.01 mol/L citrate buffer, pH 6.0, to unmask the antigenicity, and trypsin digested for 15 min, for Ki-67 and for ED1 as pre-treatment, respectively. The slides were placed in methanol with 0.3% H\(_2\)O\(_2\) for 10 min to block endogenous peroxidase activity. They were then incubated with ECP, Ki-67, and ED1 for 60 min and further processed for immunohistochemistry using the Histofine Simple Stain kit (Nichirei, Tokyo, Japan). Peroxidase activity was developed in diaminobenzidine as a chromogen. In the procedures described above, tissue sections that were not incubated with ECP, ED1, or Ki-67 served as negative controls.

**Western blot analysis**

Western blots were prepared for expression of ED1 and macrophage migration inhibitory factor (MIF). For this purpose, the colonic tissue was collected from the lesion area, and then suspended in 5 volumes of ice-cold 50 mmol/L Tris-HCl (pH 7.2) containing 150 mmol/L NaCl, 1% NP-40, 1% sodium deoxycholate, and 0.05% SDS, broken into pieces on ice, and subjected to three freeze-thaw cycles.

**After centrifugation at 15 000 g for 10 min at 4 °C,** 2-mercaptopethanol and bromophenol blue were added to the supernatant at final concentrations of 2% and 0.001%, respectively. The tissue extracts (30 µg protein) were subjected to 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis, and the separated proteins transferred onto Hybond ECL nitrocellulose membrane. After blocking nonspecific binding sites with 5% skim milk, the membrane was incubated with 1 000× diluted antibody, against ED1 and MIF (N-20, Santa Cruz Biotechnology, Inc.) at room temperature for 1 h. The bound antibodies were detected using an enhanced chemiluminescence detection kit (ECL Plus, Amersham Life Science, Buckinghamshire, UK) and the amount of each protein was quantified by densitometric analysis\(^{[20]}\).

**Statistical analysis**

All data were expressed as mean±SE. Differences between groups were examined for statistical significance using the Student’s t-test or Cochran and Cox test. A P value less than 0.05 denoted the presence of a statistically significant difference.

**RESULTS**

**Clinicopathological findings**

The body weight gain was significantly greater in anti-ECP-treated rats at d 4 and 7 after DSS treatment compared with normal serum-treated rats (Figure 1). ECP antibody clearly reduced acute clinical symptoms (diarrhea and/or blood-stained stool, Table 1). Moreover, ECP antibody improved various mucosal parameters; it reduced colonic ulceration severity (Ulcer Index) from 61.1±25.5% (DSS) to 43.1±29.4% (DSS+anti-ECP). The colon length was 13.4±0.4 cm in anti-ECP-treated rats and 12.3±0.6 cm in normal serum-treated rats at 7 d after DSS treatment (Table 2). DSS-induced shortening of the colon length was significantly abrogated in anti-ECP-treated rats (P<0.05).

**ECP expression in normal colonic mucosa and in colitis**

HE- and immunohistochemically-stained sections showed...
eosinophils and ECP-positive eosinophils in close proximity to damaged crypts in the lamina propria and partially in the extracellular interstitium of the colon of rats with DSS-induced colitis (Figures 2A and B). The ECP antibody stained activated eosinophils. In the normal colon, eosinophil infiltration was hardly observed and ECP-positive activated eosinophils were never seen in the colonic tissue (data not shown).

**Histological findings**

ECP antibody treatment clearly suppressed DSS-induced colitis as confirmed in HE-stained sections (Figures 2C and D). Moreover, immunohistochemical studies for Ki-67 and ED1 revealed that the ECP antibody treatment increased the regenerative activity of the colonic epithelium and downregulation of immune response (Figure 3). Expression of ECP in activated eosinophils was associated with the presence of erosions and inflammation. The number of Ki-67-positive cells in the regenerative surface epithelium/circumference significantly increased in anti-ECP-treated rats (21.3±10.4) compared with normal serum-treated rats (14.0±6.5) at d 7 after DSS treatment (P<0.05, Table 3 and Figure 3).

The number of ED1-positive macrophages in the lamina propria of anti-ECP-treated rats at d 7 after DSS treatment was less than that in normal serum-treated rats. Moreover, the size of ED1-positive macrophages in the lamina propria of anti-ECP-treated rats at d 7 after DSS treatment was smaller than that of normal serum-treated rats.

**Western blot analysis**

ECP antibody treatment significantly downregulated
macrophage activation during 7-d treatment of DSS (Figure 4).

Figure 3 Colonic mucosa in normal serum-treated rats (A, C, and E) and anti-ECP1-treated rats (B, D, and F) at 7-d treatment of DSS. HE staining (A and B), immunostaining for Ki-67 (C and D) and ED1 (E and F) as described under Materials and methods. Treatment with ECP antibody increased the number of Ki-67-positive cells in the regenerated surface epithelium and reduced the size of activated macrophages present in the lamina propria. Original magnification, ×400.

Figure 4 Detection of ED1 (A) and MIF (B) by Western blot analysis in colonic tissues after 7 d of DDS treatment in normal serum-treated rats (lanes 1-4) and anti-ECP-treated rats (lanes 5-7). The colonic tissue was collected from the lesion area and examined by Western blot analysis as described under Materials and methods. The relative expression levels of ED1 and MIF in the damaged colonic tissue were significantly lower in anti-ECP-treated rats than in normal serum-treated rats. The relative protein expression was quantified by densitometric analysis (n = 7, *P<0.05 vs DSS+normal serum).

DISCUSSION
A variety of clinical and experimental models has revealed that eosinophils promote pro-inflammatory changes mediated by their ability to release various cytotoxic substances and a variety of lipid mediators and cytokines. In type-2 helper T (Th2) cells-associated gastrointestinal inflammatory conditions, increased levels of eosinophils occur in the lamina propria in an eotaxin 1-dependent manner[21]. After mucosal allergen challenge, eosinophils under the regulation of IL-5 accumulate in the esophagus, an organ normally devoid of eosinophils. However, the main role of IL-5 in DSS-induced colonic inflammation is to attract a population of eosinophils that do not appear to
contribute significantly to the initiation or development of tissue damage in this model of colitis\(^{[29]}\).

In 1974, the existence of an eosinophil-granule protein, ECP, with a highly cytotoxic action, was reported by Olsson and Venge\(^{[3]}\). ECP is antibacterial\(^{[28]}\), helminthotoxic\(^{[24]}\), elicits the Gordon phenomenon when injected intratracheally into rabbits, and is cytotoxic to tracheal epithelium\(^{[25,26]}\). Although the mechanism of its cytotoxicity is not completely understood, it is suggested to be due to the pore-forming activity of ECP, which destabilizes lipid membranes\(^{[27]}\), but is unrelated to RNase activity\(^{[28]}\). We demonstrated here that a polyvalent antibody to eosinophil granule protein ECP exhibits beneficial effects on various mucosal parameters of DSS-induced colitis in rats. ECP expressed in activated eosinophils of the colonic mucosa in DSS-induced colitis were associated with the presence of injured crypts and inflammation. The body weight gain was significantly greater in anti-ECP-treated rats, compared with control rats. The ECP antibody reduced acute clinical symptoms (diarrhea and/or blood-stained stool), severity of colonic ulceration and shortening of the colon length. Immunohistochemical studies for Ki-67 and ED1 revealed that ECP antibody treatment increased the regenerative activity of the colonic epithelium and downregulation of the immune response.

Mucosal repair involves both the rapid migration of cryptal enterocytes into the injured area of the mucosa, and replacement of the mucosa by cell replication\(^{[29]}\). Various peptide growth factors regulate epithelial cell function within the mucosal epithelium of the gastrointestinal tract. Recently, Sinha et al\(^{[30]}\), reported preliminary data suggesting that epidermal growth factor enemas are effective in the treatment of active left-sided UC. Furthermore, Dignass et al\(^{[31]}\), demonstrated that hepatocyte growth factor (HGF) modulates intestinal epithelial cell proliferation and migration, thus enhancing epithelial cell restitution, the initial step of gastrointestinal wound healing, in an in vitro model. Moreover, it was demonstrated that administration of recombinant human HGF lessened colitis-associated weight loss in rats as well as improved the clinical signs of colitis in vivo\(^{[30]}\). In our rat model, mucosal erosion was observed on d 5 and accelerated on d 7, whereas the colon length was reduced on d 7. Importantly, the ECP antibody enhanced epithelial regeneration, leading to a reduction in size of colonic mucosal erosions, although it was administered concomitantly with DSS. In this regard, enhanced mucosal repair allows for a more rapid recovery of epithelial barrier function leading to reduced exposure to various luminal agents that contribute to persistent colitis. Accordingly, treatment with ECP antibody should reduce the inflammatory response to these luminal stimuli.

MIF was originally identified as a lymphokine derived from activated T cells that inhibits the random migration of macrophages in vitro and is involved in delayed-type hypersensitivity\(^{[33]}\). Moreover, it is postulated that tumor necrosis factor (TNF)-\(\alpha\) and interferon (IFN)-\(\gamma\) upregulate MIF production in macrophages and, conversely, MIF induces TNF-\(\alpha\) production, forming a proinflammatory loop within the cytokine network\(^{[34]}\). Gut monocyte macrophages are responsible for production of MIF, which in turn stimulates Th-1-type cytokines and features characteristic of colitis. Swollen macrophages induced by DSS have been seen in the mucosa with gland dropout and inflammatory cell infiltration even under intact colonic epithelium\(^{[35]}\). It seems that macrophage dysfunction, alterations of luminal bacteria, and DSS toxic effects on the colonic epithelium acted together to result in inflammatory ulcerative changes of the colonic mucosa. Therefore, in our study, the ECP antibody reduced macrophage activation such as the size of ED1-positive macrophages and the expressions of ED1 and MIF proteins in damaged colonic mucosa induced by DSS, resulting in downregulation of the immune response.

ECP is one of the major components of eosinophilic granules with a molecular mass ranging from 16 to 21.4 kDa. It exhibits various biological effects both in vitro and in vivo\(^{[36,37]}\). It is classified as a member of the ribonuclease (RNase) A supergene family based on the homology of both nucleotide and amino acid sequences. The homology of amino acid sequence between rat ECP\(^{[29]}\) and human ECP\(^{[37]}\) is 54%. When the amino acid sequence of rat ECP was compared with that of human ECP, eight structural cysteines and three amino acids, H15, K38, and H128, that are required for RNase activity\(^{[37]}\) were found to be highly conserved. The length of the signal peptide of rat ECP is equivalent to that of human ECP. Maeda et al\(^{[34]}\), demonstrated that ECP is growth-inhibitory and this ability of ECP to bind to heparin or other carbohydrate on the cell surface is dependent on tryptophan residues, W10 and W35. Interestingly, W10 is located at the P2 subsite of the catalytic domain of RNase and controls the weak RNase activity of the protein\(^{[38]}\). ECP alters the coagulation cascade\(^{[39]}\), augments fibrinolysis\(^{[40]}\), and regulates the classical pathway of complement\(^{[41]}\). Our results showed that the ECP antibody clearly reduced acute clinical symptoms of blood-stained stool. Our evaluation of the ECP molecule including W10 and H15 in human ECP enabled us to test the beneficial effects of anti-ECP treatment on inflammatory disease, in which uncontrolled cell growth could contribute to a delay in wound healing. Further studies are needed to investigate this issue.

In summary, we demonstrated that treatment with an antibody to ECP improved DSS-induced colitis in rats possibly by increasing the regenerative activity of the colonic epithelium and downregulation of the immune response. This supports the concept that humanized anti-ECP treatment could also be an effective therapy for IBDs.

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