Increased proliferation of middle to distal colonic cells during colorectal carcinogenesis in experimental murine ulcerative colitis

TAKUYA INOUE1, MITSUYUKI MURANO1, TAKANORI KURAMOTO1, KUMI ISHIDA1, KEN KAWAKAMI1, YOSUKE ABE1, EUJIRO MORITA1, NAOKO MURANO1, KEN TOSHINA1, TAKASHI NISHIKAWA1, KENTARO MAEMURA2, CHIKAO SHIMAMOTO1, ICHIRO HIRATA3, KEN-ICHI KATSU1 and KAZUHIDE HIGUCHI1

1Second Department of Internal Medicine, 2Department of Anatomy, Osaka Medical College, Osaka 569-8686; 3Department of Internal Medicine, Fujita Health University, 1-98 Dengakugakubo, Kutsukake-cho, Toyoake, Aichi 470-1192, Japan

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Abstract. Patients with ulcerative colitis (UC) exhibit an increased risk for the development of cancer of the colon and rectum. This association is widely attributed to colonic inflammation. However, the severity of colonic inflammation necessary for the development of dysplasia and/or cancer remains unknown. In this study, we investigated the pattern of cell proliferation in colorectal carcinogenesis in an experimental murine model of UC. Chronic colitis was induced by administration of four cycles of dextran sulfate sodium (DSS) (each cycle: 5% or 2% DSS for 7 days and then distilled water for 14 days). Mice were sacrificed after every cycle and at 120 days following the completion of the fourth cycle. Colonic cell proliferation was immunohistochemically evaluated using the thymidine analogue bromodeoxyuridine and the labeling index (LI) was determined. The incidence of dysplasia and/or cancer was 28%, 6.7%, and 0% in the 5% DSS, 2% DSS, and normal control groups respectively. All gross lesions were present in the middle to distal colon. Disease activity index and total LI after four cycles of DSS were significantly higher in the 5% DSS group compared to the 2% DSS group. In the 5% DSS group, the LI was significantly higher in the middle colon than in the proximal colon. Simple repeated administration of the non-genotoxic colon carcinogen DSS induced dysplasia and/or cancer. In addition, we have demonstrated the presence of regional differences in proliferation pattern between the middle and the proximal colon during carcinogenesis in experimental murine UC. These findings may provide insight into the development of colorectal cancer in humans with long-standing UC.

Introduction

Patients with ulcerative colitis (UC) exhibit an increased risk for the development of cancer of the colon and rectum with the risk of colorectal cancer increasing as the extent of colon affected and duration of disease increase. Indeed, the incidence of colorectal cancer in patients with long-standing UC is higher than that of sporadic colorectal cancer (1). Although this association is widely attributed to colonic inflammation, it is unknown whether the degree of inflammation relates to the development of dysplasia and/or cancer. Thus, experimental models of colitis that develop dysplasia and cancer similar to human UC are needed to investigate the dysplasia-cancer sequence and, although there are many animal models of inflammatory bowel disease (IBD), few are applicable to such study. Moreover, in most of these animal models, dysplasia and cancer are induced by pretreatment with genotoxic colon carcinogens such as azoxymethane (AOM) (2-4) or dimethylhydrazine (DMH) (5,6) followed by dextran sulfate sodium (DSS) administration. These studies suggest that chronic or repeated mucosal inflammation induced by administration of DSS may result in the rapid development of colorectal cancer. On the other hand, in the absence of genotoxic colon carcinogens, the long-term treatment of mice with the non-genotoxic compound DSS results in dysplasia and cancer which has clinical and histopathological similarities to human UC (7-9).

Although experimental chronic colitis and colorectal cancer induced in rodents by administration of DSS has been extensively investigated, the mechanisms by which DSS induces both colonic inflammation and cancer are still unknown. For example, it is unclear whether dysplastic or proliferative changes occur first. However, abnormal proliferation is present in the normal-seeming mucosa of human UC (10). Therefore, abnormal epithelial cell proliferation is implicated in chronic colitis associated with subsequent carcinogenesis (11). Hyperproliferation of colonic epithelial cells was observed in tritiated thymidine-labeled specimens from carcinogen-treated mice (12). However, there have been no reports on the pattern of cell proliferation during carcinogenesis in experimental murine UC induced by simple repeated DSS administration.
Bromodeoxyuridine (BrdU) is a thymidine analogue which becomes incorporated into DNA during the S-phase of the cell cycle and can be clearly detected using a monoclonal antibody (13). Several studies have demonstrated that the labeling index (LI) estimated by BrdU immunohistochemistry is equivalent to that obtained by thymidine autoradiography (14) and BrdU labeling has been considered to accurately represent the level of proliferative activity (15).

In the present study, we established a murine model of long-standing UC induced by simple repeated administration of the non-genotoxic colon carcinogen DSS. This resulted in the induction of dysplasia and/or cancer similar to UC-associated neoplasia although the number of lesions was relatively small. We then investigated the influence of the severity of colonic inflammation on colorectal carcinogenesis and regional differences in cell proliferation that might be associated with a higher incidence of tumors.

Materials and methods

Animals. Seven-week-old female BALB/c mice (CELA Japan, Tokyo, Japan) weighing 20-25 g were used in this study. The animals were maintained in an animal colony with controlled temperature (23˚C) and light (12/12-h light and dark cycle) at the Osaka Medical College (OMC), Osaka, Japan, and were permitted free access to standard mice chow pellets (MM-3, Funabashi, Chiba, Japan) and tap water.

Induction of experimental colitis. DSS (molecular weight 5000) was obtained from Meitou Sangyou (Osaka, Japan). Mice were given water containing 5% or 2% DSS instead of tap water on the indicated days.

Protocol for induction of colorectal tumors and experimental procedures. The experimental design is shown in Fig. 1. Chronic colitis was induced in mice by four cycles of oral administration of DSS with each cycle comprising DSS treatment for 7 days and distilled water for the following 14 days. Mice received intraperitoneal BrdU (B5002 Sigma Chemicals, St. Louis, MO, USA) at a dose of 40 mg/kg body weight 1 h before sacrifice. Cohorts of mice were sacrificed after every cycle to determine the labeling index, and at 120 days following the completion of the fourth cycle to evaluate the incidence of dysplasia and/or cancer. Mice were divided into three groups based on the amount of DSS they received: 5% DSS, 2% DSS and a normal control group.

Evaluation of the severity of clinical colitis. The disease activity index (DAI) was determined in all animals during the first administration cycle of DSS by scoring the extent of body weight loss, stool haemoccult reactivity or gross bleeding, and stool consistency in accordance with the method described by Murthy et al (16). This method of scoring is a comprehensive functional measure and correlates well with the degree of inflammation. The individuals who examined the mice and determined the DAI scores were blinded as to the experimental group to which the animal belonged.

Pathologic examination. After sacrifice, the entire colorectum from the colocecal junction to the anal verge was excised and rinsed in phosphate-buffered saline (PBS). The specimen was opened longitudinally, examined for gross lesions and all gross lesions were recorded. The colon was divided into three equal portions (proximal, middle and distal) and each segment was fixed in 10% formalin. Subsequently, three transverse sections of each segment were submitted for histological processing. All slides were stained with hematoxylin and eosin.

According to the histological criteria outlined by the Inflammatory Bowel Disease - Dysplasia Morphology Study Group (17), dysplasia was categorized on the basis of microscopic findings. This included architectural alteration beyond the degree of reparative changes in chronic colitis, which often resembled the glandular arrangement of adenomas, and cytologic abnormalities that consisted principally of cellular and nuclear pleomorphism, nuclear hyperchromatism, loss of nuclear polarity and marked stratification of nuclei. The cancer category included ‘early invasive’, which was defined as cancer cells invading the muscularis mucosa and/or the submucosa. The indefinite category consisted of ‘cytologically atypical epithelium’, which was recognized for the most part to be reparative and/or regenerative in nature.

BrdU immunohistochemistry. Expression of BrdU in the intestinal mucosa was assessed by the labeled streptavidin biotin method using an LSAB kit (Dako, Carpinteria, CA,
Table I. Colorectal length and gross lesion position.

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<th>DSS (5%)</th>
<th>DSS (2%)</th>
<th>Normal control</th>
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<td>Colon length (cm)</td>
<td>8.2±0.93</td>
<td>9.1±0.6</td>
<td>12.1±0.6</td>
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<tr>
<td>Distance from anus of gross lesions (cm)</td>
<td>2.7±1.2</td>
<td>2.3</td>
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USA). The entire colon was divided into three equal portions (proximal, middle and distal) and each longitudinal segment was fixed in 10% formalin, embedded in paraffin wax and cut into tissue sections of 4 μm thickness. Tissue sections were mounted on microscope slides, dewaxed in xylene (3x3 min) and dehydrated with 100% ethanol. After washing with PBS, sections were treated with 2 mol/l HCL for 30 min and then treated twice with 0.1 mol/l borate buffer (pH 8.5) for 5 min. After washing with PBS, endogenous peroxidase activity was blocked using 0.3% hydrogen peroxidase in 10% methanol for 30 min and blocking reagent was added for 15 min. Sections were incubated at 4°C overnight with the primary mouse monoclonal antibody (1 in 20 dilution in PBS) against BrdU (Dako). After washing with PBS, sections were incubated with a biotinylated anti-mouse immunoglobulin antibody (Dako) at room temperature for 30 min. Sections were then washed in PBS and visualized using streptavidin-biotin-horseradish peroxidase (Dako) and 3, 3' diaminobenzidine (Dako). Finally, sections were counterstained with Meyer's hematoxylin solution, dehydrated and cover-slipped with permanent mounting medium for microscopic examination.

**Evaluation of the labeling index (LI).** Only complete well-orientated longitudinally sectioned crypts displaying the lumen at the top and the muscularis mucosa at the base were used for analysis. Labeled cells were defined as epithelial cells whose nuclei were stained brown at a magnification of x400. The LI for the whole crypt and for each compartment (proximal, middle and distal) was determined after counting the number of BrdU-labeled cells per 10 crypts and dividing the number of labeled cells by the total cell number and multiplying by 100.

**Statistical analysis.** All results are expressed as mean ± SD. Comparisons were performed using one-way ANOVA followed by Tukey-Kramer’s test. Categorical data were analyzed by the chi-squared test. Statistical significance was defined as P<0.05.

**Results**

**Changes of DAI score.** Most mice in the 5% DSS group exhibited loose and hemoccult-positive stools 2 days after DSS administration (day 2). Clinical symptoms of colitis including bloody stool, diarrhea and loss of body weight progressed further until day 8 but then gradually disappeared during the subsequent 14-day period of drinking distilled water without DSS. Accordingly, the DAI score gradually increased from day 2 to day 8 and usually reverted to normal by day 21. Treatment of mice with 5% DSS significantly increased the DAI score from day 4 to 14 compared to the 2% DSS group (Fig. 2).

**Colorectal length, incidence and distribution of dysplasia and/or cancer and tumor size.** The colorectal length was significantly shortened compared to the normal control group after 120 days from the completion of the fourth cycle of DSS administration. The colorectal length in the 5% DSS, 2% DSS and normal control groups was 8.2±0.9 cm, 9.1±0.6 cm and 12.1±0.6 cm, respectively. All gross lesions had the shape of the sessile type and were observed in the middle or distal colon (Table I) whilst all mice with dysplasia and/or cancer had lesions limited to only one colon segment (Fig. 3). The size of gross lesions in the 5% and 2% DSS groups was 4480±1184 μm and 2000 μm, respectively. The incidence of dysplasia and/or cancer was presented in Table II. The incidence of dysplasia and/or cancer in the 5% and 2% DSS group was statistically significant higher in the 5% DSS group than in the 2% DSS group. Concerning regional differences, the LI was significantly greater in the middle colon (8.5±1.65%) than in the proximal colon (6.3±1.95%) in the 5% DSS group after four cycles of DSS administration (Table III).

**BrdU labeling in the colorectum.** The time course study of *in vivo* BrdU uptake is shown in Fig. 4. Although BrdU immunoreactive cells amounted to only 0.71±0.34% of total cells in the whole colon of normal control mice, the percentage of BrdU-positive cells after four cycles of DSS administration increased to 7.69±1.86% and 4.56±2.21% in the 5% DSS and 2% DSS groups respectively. The total LI was significantly higher in the 5% DSS group than in the 2% DSS group. Concerning regional differences, the LI was significantly greater in the middle colon (8.5±1.65%) than in the proximal colon (6.3±1.95%) in the 5% DSS group after four cycles of DSS administration (Table III).

**Discussion**

DSS is a synthetic sulfate polysaccharide composed of dextran with sulfate glucose and is known to induce colitis, and DSS-induced murine colitis has been used as a model of human UC (5,8). The mechanisms of DSS-induced colitis are believed to involve macrophage dysfunction, altered luminal bacterial flora and toxic effects upon the colonic epithelium.
Colorectal cancer is one of the most serious complications of UC and the risk of UC-associated neoplasia increases as the extent and duration of the disease increase. Indeed, the incidence of colorectal cancer in patients with long-standing UC is higher than that of sporadic colorectal cancer (1). A few investigators have described the appearance of dysplasia and/or cancer in mice following simple repeated administration of the non-genotoxic colon carcinogen DSS and suggested that repeated mucosal erosion with necrosis and regeneration of the colonic epithelium seems to increase the susceptibility of the mucosal epithelia for dysplasia and/or cancer development (7). In the present study, simple repeated administration of the non-genotoxic colon carcinogen DSS induced dysplasia and/or cancer in mice, though the neoplastic lesions were relatively few in number compared to those induced by genotoxic carcinogens such as AOM and DMH. Interestingly, in contrast to human UC, there were no mice that exhibited more than one neoplasm in this study. Using
In conclusion, the present data demonstrate that simple repeated administration of the non-genotoxic colon carcinogen DSS induced dysplasia and/or cancer and that the high level of inflammation could be an important factor in the earlier stage of initiation of dysplasia. Although further studies are needed to clarify the relationship between continued inflammation and apoptosis, a persistent abnormal level of background cell proliferation of the mucosa together with regional differences in cell proliferation may play an important role in the development of dysplasia and/or cancer in this murine model of colitis. These findings may also provide insight into the development of colorectal cancer in humans with long-standing ulcerative colitis.

Acknowledgements

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