Identification of dihalogenated proteins in rat intestinal mucosa injured by indomethacin

Tomohisa Takagi,¹ Yuji Naito,^{1,*} Hitomi Okada,¹ Tetsuya Okayama,¹ Katsura Mizushima,¹ Shinya Yamada,¹ Kouhei Fukumoto,¹ Ken Inoue,¹ Megumi Takaoka,¹ Tomoko Oya-Ito,¹ Kazuhiko Uchiyama,¹ Takeshi Ishikawa,¹ Osamu Handa,¹ Satoshi Kokura,¹ Nobuaki Yagi,¹ Hiroshi Ichikawa,¹ Yoji Kato,² Toshihiko Osawa³ and Toshikazu Yoshikawa¹

¹Molecular Gastroenterology and Hepatology, Graduate School of Medical Science, Kyoto Prefectural University of Medicine, Kyoto 602-8566, Japan

²School of Human Science and Environment, University of Hyogo, Himeji 670-0092, Japan

³Department of Nutritional Science, Aichi Gakuin University, Nisshin 470-0195, Japan

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Previous studies have shown that activated neutrophils and their myeloperoxidase (MPO)-derived products play a crucial role in the pathogenesis of non-steroidal anti-inflammatory drug (NSAID)-related small intestinal injury. The aim of the present study is to identify dihalogenated proteins in the small intestine on indomethacin administration. Intestinal damage was induced by subcutaneous administration of indomethacin (10 mg/kg) in male Wistar rats, and the severity of the injury was evaluated by measuring the area of visible ulcerative lesions. Tissue-associated MPO activity was measured in the intestinal mucosa as an index of neutrophil infiltration. The dihalogenated proteins were separated by two-dimensional polyacrylamide gel electrophoresis (2D-PAGE) using novel monoclonal antibodies against dibromotyrosine (DiBrY), and they were identified by matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) peptide mass fingerprinting and a Mascot database search. Single administration of indomethacin elicited increased ulcerative area and MPO activity in the small intestine. 2D-PAGE showed an increased level of DiBrY-modified proteins in the indomethacin-induced injured intestinal mucosa and 6 modified proteins were found. Enolase-1 and albumin were found to be DiBrY modified. These proteins may be responsible for the development of neutrophil-associated intestinal injury induced by indomethacin.

Key Words: indomethacin, dibromotyrosine, albumin, enolase

Non-steroidal anti-inflammatory drugs (NSAIDs) are commonly used world over for the treatment of musculoskeletal pain and inflammation. A major limitation of the clinical utility of NSAIDs is their gastrointestinal toxicity and the increased risk of serious, possibly fatal, complications of the upper gastrointestinal tract, including bleeding, ulceration, and perforation of the stomach and duodenum.^(1–4) Furthermore, recent advances in gastrointestinal endoscopies such as capsule endoscopy and balloon endoscopy for investigation of the entire small intestine have revealed that NSAIDs can cause mucosal injuries in the small intestine as well as a variety of abnormalities such as ulcerations, perforation, bleeding, and diaphragm-like strictures.^(5,6) Therefore, it is important to investigate the precise pathogenesis of NSAID-induced intestinal injuries and determine preventive and therapeutic strategies for them.

Investigations using experimental animal models have shown that NSAIDs such as indomethacin damage the small intestine. Increasing evidence suggests that the intestinal injury caused by indomethacin is associated with abnormal intestinal permeability, bacterial translocation, activation of inflammatory cytokines, nitric oxide (NO) overproduction, and prostaglandin deficiency,

in addition to neutrophil accumulation.^(7–11) In particular, the infiltration of neutrophils into the intestinal mucosa may play a crucial role in the pathogenesis of NSAID-induced intestinal injury. Stadnyk *et al.* demonstrated that neutrophils were detectable in the small intestine of rats at 6 h after indomethacin administration and continued to accumulate until 48 h after administration.⁽¹²⁾ We also previously reported that neutrophil infiltration gradually increased in a time-dependent manner after indomethacin administration in rats.^(7,13,14) Interestingly, impaired leukocyte recruitment and neutrophil depletion resulted in the amelioration of NSAID-induced injury in mice.^(15,16) Thus, neutrophil-mediated inflammation can be considered to be involved in NSAID-induced intestinal injury.

On the other hand, neutrophils have granules containing peroxidases such as myeloperoxidase (MPO). MPO is known to catalyze the formation of hypochlorous acid (HOCl) and hypobromous acid (HOBr) using hydrogen peroxide (H₂O₂) and Cl[–] or Br[–], respectively. These reactive intermediates may react with proteins,^(17,18) lipids,^(19,20) and nucleotides,^(21–23) and they reportedly cause tyrosine halogenation; such halogenations give rise to products such as dibromotyrosine (DiBrY),^(24,25) which is a tyrosine molecule modified by bromine at the 3- and 5-positions and is one of the major oxidative products derived from neutrophil MPO.

The role of tyrosine halogenation in the development of neutrophil-mediated inflammatory damage such as NSAID-induced intestinal injuries remains unclear. In this study, we identified the DiBrY-modified proteins involved in indomethacin-induced intestinal injuries by using a proteomics-based approach.

Materials and Methods

Experimental animals. Male Wistar rats weighing 190–210 g were obtained from Shimizu Laboratory Supplies Co., Ltd. (Kyoto, Japan). The animals were housed at 22°C in a controlled environment with 12 h of artificial light per day, and they were allowed *ad libitum* access to rat chow and water. The experiments were performed on 5–6 non-fasting rats per group without anesthesia. Animal maintenance and all experimental procedures were carried out in accordance with the NIH guidelines for the use of experimental animals. All experimental protocols were approved by the Animal Care Committee of the Kyoto Prefectural University of Medicine (Kyoto, Japan).

Induction of small intestinal lesions. The animals were

*To whom correspondence should be addressed.
E-mail: ynaito@koto.kpu-m.ac.jp

subcutaneously administered 10 mg/kg indomethacin (Sigma Chemical; St. Louis, MO) and killed 24 h later under deep ether anesthesia. To determine the extent of injury, 1% Evans blue was injected intravenously 30 min before euthanasia; the jejunum and ileum were then removed, opened along the antimesenteric attachment, and examined for lesions under a dissecting microscope with square grids. The area (in mm²) of visible lesions was macroscopically measured, totaled per 20 cm of the small intestine, and expressed as an ulcer index. The degree of intestinal injury was evaluated by an independent observer who was blinded to the experimental conditions. For histological examination, formalin-fixed tissue was stained with hematoxylin and eosin (H&E). Staining was evaluated by light microscopy by a pathologist who was also blinded to the experimental conditions.

Measurement of MPO activity. Tissue-associated MPO activity was determined by a modification of the method of Grisham *et al.*⁽²⁶⁾ as an index of neutrophil accumulation. The intestinal mucosa was scraped off using two glass slides and then homogenized with 1.5 ml of 10 mmol/l potassium phosphate buffer (pH 7.8) containing 30 mmol/l KCl in a Teflon Potter-Elvehjem homogenizer. The mucosal homogenates were centrifuged at 20,000 × *g* for 15 min at 4°C to pelletize the insoluble cellular debris. The pellet was then dissolved in an equivalent volume of 0.05 M potassium phosphate buffer (pH 5.4) containing 0.5% hexadecyltrimethylammonium bromide. The samples were centrifuged at 20,000 × *g* for 15 min at 4°C and the supernatants collected. MPO activity was assessed by measuring the H₂O₂-dependent oxidation of 3,3',5,5'-tetramethylbenzidine. One unit of enzyme activity was defined as the amount of MPO required to cause a 1.0/min change in absorbance of at 645 nm and 25°C. The level of MPO activity in the mucosal homogenates was expressed as unit per milligram of protein. The total protein in the tissue homogenates was measured using a Bio-Rad Protein Assay kit (Bio-Rad Laboratories, KK; Tokyo, Japan) according to the manufacturer's protocol.

Sample preparation and two-dimensional polyacrylamide gel electrophoresis (2D-PAGE). The intestinal tissue samples (200 mg) were homogenized in 2 ml homogenization buffer (8 M urea, 4% 3[(3-Cholamidopropyl)dimethylammonio]propanesulfonic acid (CHAPS), 40 mM Tris) containing nuclease and protein inhibitors (GE Healthcare UK Ltd.; Buckinghamshire, England) using a homogenizer at 25,000 rpm. The homogenized samples were transferred to an ultracentrifuge tube, and the nucleic acids were removed by centrifugation (20 min at 20,000 × *g* and 25°C). The samples were then precipitated using the PlusOne™ 2D Clean-Up kit as recommended by the manufacturer (GE Healthcare UK Ltd.). The protein concentration in the supernatant fraction was determined using the Bradford assay with bovine serum albumin as the standard.

The samples were solubilized in 6 M urea, 20 mM dithiothreitol, 30% glycerol, 45 mM Tris base, 1.6% lithium dodecyl sulfate (LDS) (Invitrogen Japan KK, Tokyo, Japan), and 0.002% bromophenol blue and were subsequently heated at 70°C for 10 min. Protein lysates (50 µg) were separated by 2D-PAGE. Immobilized pH gradient (IPG) strips of pH 4–7 or pH 5.3–6.3 (Invitrogen Japan KK) were rehydrated overnight with the protein samples. The proteins were separated on the basis of their respective isoelectric points by isoelectric focusing (IEF) using ZOOM® IPGRUNNER™ (Invitrogen Japan KK) at a maximal voltage of 2000 V and a current limit of 50 µA per gel. Following IEF, the IPG strips were incubated in equilibration buffer I (6 M urea, 130 mM dithiothreitol, 30% glycerol, 45 mM Tris base, 1.6% LDS, 0.002% bromophenol blue; Genomic Solutions) and once in equilibration buffer II (6 M urea, 135 mM iodoacetamide, 30% glycerol, 45 mM Tris base, 1.6% LDS, 0.002% bromophenol blue; Genomic Solutions) for 15 min each. The equilibrated IPG strips were applied to 4–12% Bis-Tris gradient gels (Invitrogen Japan KK), and the NuPAGE® MOPS buffer (Invitrogen Japan KK) was

used at 200 V for 55 min to separate the proteins in the second dimension on the basis of their molecular size. Following electrophoresis, the gels were transferred onto nitrocellulose and immunoblotted with the anti-DiBrY monoclonal antibody (3A5), which was kindly gifted to us by Prof. Toshihiko Osawa.⁽²⁴⁾

In-gel protein digestion and peptide mass fingerprinting. Protein spots of interest were excised using Xcise Proteomics Systems (Shimadzu Corp., Kyoto, Japan) from the preparative gel stained with Deep Purple™ Total Protein Stain (GE Healthcare UK Ltd). The excised spots were washed thrice with 50 mmol/l ammonium bicarbonate and 50% acetonitrile (ACN), dehydrated with 100% ACN. The proteins were subjected to in-gel digestion with 10 µg/ml trypsin (Promega KK; Tokyo, Japan) in 50 mmol/l ammonium bicarbonate at 30°C overnight. Tryptic peptides were extracted from the gel slices with 1% trifluoroacetic acid and 50% ACN. After concentration and desalting using a Millipore ZipTip_µ-C18 (Nihon Millipore KK; Tokyo, Japan), the resulting peptides were mixed with an equal volume of 10 mg/ml 2,5-dihydroxybenzoic acid (DHBA), and the peptide mass spectra were obtained using the AXIMA-QIT™ MALDI-TOF-MASS (Shimadzu) platform for peptide mass fingerprinting. Protein identification was carried out using the Mascot search engine (http://www.matrixscience.com/search_form_select.html).

Statistical analysis. Results are presented as the mean ± (standard error of the mean [SEM]). Overall differences between groups were determined by one-way analysis of variance (ANOVA). Whenever the one-way ANOVA was significant, differences between individual groups were analyzed by Bonferroni's multiple comparisons test. Differences of *p* < 0.05 were considered significant. All analyses were performed using the GraphPad Prism 4 program (GraphPad Software Inc.; San Diego, CA) for Macintosh.

Results

Intestinal ulceration after indomethacin administration.

A single administration of 10 mg/kg indomethacin elicited multiple erosions in the small intestine (Fig. 1B); a normal untreated small intestine is shown in Fig. 1A for comparison. The ulcer index gradually increased with time, and significant increases in this index were noted 6, 12, and 24 h after indomethacin administration (Fig. 1C).

Histological findings and time course of changes in MPO activities. The histological features observed were defects of the villi, epithelial stratification, basal lamina degeneration, and infiltration of the mucosa by inflammatory cells (neutrophils) (Fig. 2B); none of these were observed in the normal intestinal

Fig. 1. Representative macroscopic findings and ulcer index after indomethacin administration. To determine the extent of intestinal injury, 1% Evans blue was injected intravenously 30 min before euthanasia. The representative macroscopic findings indicate that the small intestinal injuries were induced 24 h after indomethacin administration (B); a normal untreated intestine is shown in (A). The ulcer index after indomethacin administration was evaluated as described in Materials and Methods (C). Data represent the mean ± (SEM) of 7 rats. **p* < 0.01 compared to the sham group (0 h).

Fig. 2. Representative histological findings and neutrophil accumulation in the intestinal mucosa after indomethacin administration. Histological appearance of the intestinal tissue of the sham group rats (A) and rats treated with indomethacin (B). Histological examination revealed that indomethacin administration induced ulceration in the small intestine, which was associated with the infiltration of numerous inflammatory cells. Hematoxylin and eosin (H&E) staining ($\times 40$). The time course of tissue-associated myeloperoxidase (MPO) activity after indomethacin administration was determined as an index of neutrophil accumulation in the intestinal mucosa (C). Data represent the mean $\pm$ (SEM) of 8 rats. * $p < 0.05$ compared to the sham group (0 h).

mucosa (Fig. 2A). Neutrophil accumulation was evaluated by measuring the MPO activity in the intestinal mucosa homogenates. The MPO activity in the intestinal mucosa was markedly increased by indomethacin treatment and the increase was significant 24 h after administration (Fig. 2C).

Identification of DiBrY-modified proteins in intestinal mucosa with indomethacin-induced injury. To evaluate the expression of DiBrY-modified proteins in the intestinal mucosa, we performed 2D-PAGE by using a pH 4–7 IPG strip and 4–12% Bis-Tris gradient gels. Fig. 3 (upper panel; A, B) shows individual proteins separated by 2D-PAGE and stained with Deep Purple™ Total Protein Stain. Subsequently, the gels were transferred onto nitrocellulose and were immunoblotted with anti-DiBrY monoclonal antibody (Fig. 3, lower panel; C, D). We focused on analyzing DiBrY-modified proteins between pI 5.3 and 6.3 of size between 20 and 60 kDa. (Fig. 3, lower panel, boxed image).

Next, the individual proteins were separated by 2D-PAGE between pI 5.3 and 6.3 (Fig. 4). As shown in Fig. 4B, the immunoblot analysis revealed 6 DiBrY-modified proteins (indicated by arrows). These 6 proteins were analyzed by MALDI-TOF using peptide-mass fingerprinting, and a database search was performed with Mascot (Table 1). Enolase-1 and albumin were found to be DiBrY-modified in the intestinal mucosa with indomethacin-induced injury.

Fig. 3. Image of 2D-PAGE analytical gels and immunoblotting with anti-dibromotyrosine (DiBrY) monoclonal antibody. (A, B) 2D-PAGE gels were stained using Deep Purple™ Total Protein Stain. Subsequently, they were transferred onto nitrocellulose and immunoblotted with the anti-DiBrY monoclonal antibody (C, D). Boxes mark image areas that were investigated in subsequent experiments at pH 5.3–6.3.

Fig. 4. Image of 2D-PAGE between pH 5.3 and 6.3. (A, B) 2D-PAGE between pH 5.3 and 6.3 gels were stained using Deep Purple™ Total Protein Stain. (C, D) Immunoblotting analysis revealed 6 DiBrY-modified proteins (arrows) in the intestinal mucosa injured by indomethacin treatment.

Table 1. List of dibromotyrosine-modified proteins identified in indomethacin-induced injured intestinal mucosa

Spot No.	Protein identity	pI	MW (Da)	MASCOT score	Sequence coverage (%)
1	enolase-1	6.16	47440	78	29
2	enolase-1	6.16	47440	58	32
3	Albumin isoform CRA_a	6.72	53060	70	21
4	Albumin isoform CRA_a	6.72	53060	93	34
5	Albumin isoform CRA_a	6.72	53060	119	46
6	Albumin isoform CRA_a	6.72	53060	82	23

pI, Isoelectric point; MW, Molecular Weight.

MASCOT score described the significance of the search results from the search engine MASCOT and a significance threshold of about 60 is typical. The sequence coverage was defined as the ratio of the number of the identified amino acids to the total number acid of the protein (%).

Discussion

In this study, we performed 2D-PAGE to identify DiBrY-modified proteins in intestinal mucosa injured because of indomethacin. We confirmed that 2 proteins (enolase-1 and albumin) exhibited altered expression, as shown in Table 1. It is well known that albumin possesses potent antioxidant properties and non-specifically binds to free radicals.^(27,28) Enolase-1 modification has also been reported in many reports.^(29–33) The present proteome analysis might provide important, novel clues for understanding NSAID-induced intestinal injuries and reveal candidates for therapeutic targets.

Enolase, also known as phosphopyruvate dehydratase, is a glycolytic enzyme. Mammalian enolase has 3 isoforms: enolase-1 (α -enolase), enolase-2 (γ -enolase), and enolase-3 (β -enolase). The expression of these isoforms is developmentally regulated in a tissue-specific manner. Enolase-1 is widely distributed in a variety of tissues, whereas enolase-2 and enolase-3 are found exclusively in neurons or neuroendocrine tissue and muscle tissues, respectively.⁽³⁴⁾ Interestingly, recent investigations have revealed that enolase enzymes possess various different regulatory properties in addition to their role in glycolysis and gluconeogenesis. Enolase-1 plays a key role in anaerobic metabolism and acts as a plasminogen receptor (Plg-Rs),⁽³⁵⁾ indicating that it may play a role in tissue invasion. In fact, the enhanced binding and activation of plasminogen by neoplastic cells has been attributed to enhanced enolase-1 expression, and abnormal expression of enolase-1 is associated with tumor progression in lung cancer.^(36–38) These findings collectively indicate that, enolase-1 may be a potential therapeutic target for cancer. Further, although the role of enolase-1 remains unclear in intestinal inflammation, Wygrecka *et al.*

demonstrated that lipopolysaccharide (LPS) induced the expression of enolase-1 in monocytes, and the increased cell-surface expression of enolase-1 mediated the invasion of inflammatory cells.⁽³⁹⁾

In proteomics studies, enolase-1 has been frequently identified to be differentially expressed in 2D-PAGE-based experiments on human and animal tissues. In particular, enolase-1 was found to be excessively carbonylated,^(30,31) nitrated,⁽³³⁾ and 4-hydroxynonenal (HNE)-modified^(29,32) in various diseases. In the present study, we identified DiBrY-modified enolase-1 in indomethacin-induced intestinal injury. Although further studies on the precise function of DiBrY-modified enolase-1 are required, the protein may be considered to play an important role in the pathogenesis of intestinal inflammation.

In conclusion, we identified DiBrY-modified proteins, enolase-1 and albumin, in intestinal tissue with indomethacin-induced injury. Although further examination is required to gain a better understanding of the role of these modified proteins, DiBrY is expected to be a novel biomarker for inflammation-related tissue damage and may be involved in neutrophil accumulation.

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