Roles of Endogenous Prostaglandins and Cyclooxygenase Isozymes in Healing of Indomethacin-Induced Small Intestinal Lesions in Rats

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ABSTRACT

The role of prostaglandins (PGs)/cyclooxygenase (COX) in the healing of indomethacin-induced small intestinal ulcers was examined in rats. Animals were given indomethacin (10 mg/kg s.c.) and killed 1, 2, 3, 5, and 7 days later. Indomethacin (2 mg/kg), 5-(4-chlorophenyl)-1-(4-methoxyphenyl)-3-trifluoro-methylpyrazole (SC560; COX-1 inhibitor; 3 mg/kg), and rofecoxib (COX-2 inhibitor; 3 mg/kg) were given p.o. once daily for 6 days, during the first 3 days or last 3 days of the experimental period. All COX inhibitors given for 6 days significantly impaired the healing of these ulcers. Healing was also impaired by rofecoxib given for the first 3 days or by SC560 given for the last 3 days. The expression of COX-2 mRNA in the intestine was up-regulated after ulceration, persisting for 3 days and dissipating thereafter. Mucosal PGE₂ contents decreased within 3 h after ulceration, recovered 24 h later, and increased above normal 1–3 days later. The PGE₂ content at 4 days after ulceration was decreased by rofecoxib but not SC560, whereas that at 7 days was suppressed by SC560 but not rofecoxib. Vascular content in the ulcerated mucosa decreased when the healing was impaired by COX inhibitors. The deleterious effect of indomethacin on healing was mimicked by a prostacyclin E receptor (EP) 4 antagonist and reversed by coadministration of PGE₂ as well as an EP4 agonist. In conclusion, endogenous PGs play a role in the healing of intestinal ulcers through EP4 receptors, yet the COX isozyme involved differs depending on the stage of healing; COX-2 in the early stage and COX-1 in the late stage.

Nonsteroidal anti-inflammatory drugs (NSAIDs), the most frequently prescribed drugs worldwide, have been used in patients as analgesic and antinociception as well as anti-inflammation drugs, but the major limitation to their use is gastrointestinal side effects. These drugs not only damage the gastrointestinal mucosa but also impair the healing of pre-existing ulcers as well (Wang et al., 1989; Levi et al., 1990; Schmassmann et al., 1995; Wallace et al., 2000; Tanaka et al., 2002a). These effects of NSAIDs are considered to be brought about by a deficiency of prostaglandins (PGs) due to inhibition of cyclooxygenase (COX). COX exists in two isoforms, COX-1 expressed constitutively in various tissues, including the stomach and intestine, and COX-2 expressed in few tissues but rapidly induced in response to growth factors and cytokines (Feng et al., 1993; Kargman et al., 1993; O’Neill and Ford-Hutchinson, 1993; Singer et al., 1998).

It is known that the deleterious effect of NSAIDs on healing is shared by selective COX-2 inhibitors (Mizuno et al., 1997; Ukawa et al., 1998; Shigeta et al., 1998; Halter et al., 2001; Araki et al., 2002), suggesting an important role for COX-2/PGs in the healing of pre-existing ulcers. Mizuno et al. (1997) first demonstrated that both COX-2 mRNA and protein were strongly expressed in stomachs in which ulcers had been induced. More recently, Jones et al. (1999) reported that both COX-1 and COX-2 are important for the regulation of angiogenesis and that selective COX-2 inhibitors and non-selective NSAIDs inhibit angiogenesis through direct effects on endothelial cells. However, since the repair of epithelial damage after irradiation was retarded in COX-1 knockout mice (Blikslager et al., 2002), it is possible that the mecha-
nism of ulcer healing is regulated by endogenous PGs produced not only by COX-2 but also by COX-1. Unfortunately, most of the above-mentioned findings were obtained in the stomach but not the small intestine. Thus, the relative role of COX-1 and COX-2 in the healing of the intestinal lesions remains unknown.

In the present study, we examined the roles of COX/PGs in the healing of indomethacin-induced intestinal lesions in rats, using selective COX-1 and COX-2 inhibitors. Since it has been recently shown that PGE$_2$ enhanced the healing of gastric ulcers through the activation of EP4 receptors (Tanaka et al., 2005), we also examined the effect of various EP agonists and antagonist on the healing of intestinal lesions and investigated which EP receptor subtype is involved in the healing process.

Materials and Methods

Animals. Male Sprague-Dawley rats (220–260 g; Nippon Charles River, Shizuoka, Japan) were used. Studies were carried out using four to nine animals without fasting in a conscious state, unless otherwise specified. All experimental procedures described here were approved by the Experimental Animal Research Committee of the Kyoto Pharmaceutical University.

Induction of Small Intestinal Ulceration. The animals were administered indomethacin (10 mg/kg s.c.) and killed 1, 2, 3, 5, and 7 days later. Various COX inhibitors, such as indomethacin (a non-selective COX inhibitor; 2 mg/kg), SC560 (a selective COX-1 inhibitor; 3 mg/kg), and rofecoxib (a selective COX-2 inhibitor; 3 mg/kg), were given p.o. once daily for the first 3 days, for 6 days, starting 1 day after the treatment with indomethacin (10 mg/kg). In some cases, both SC560 and rofecoxib were given together. Control animals were given the vehicle (hydroxy propyl cellulose solution) p.o. once daily for 6 days, starting 1 day after the treatment with indomethacin (10 mg/kg). In addition, the animals treated with various COX inhibitors for the first 3 days or the last 3 days were given the vehicle p.o. once daily for the rest of the experimental period. In another experiment, the effect of the EP4 antagonist AE3-208 (Amano et al., 2003) on the healing of indomethacin-induced intestinal lesions was examined. AE3-208 (3 mg/kg) was given i.p. twice daily for 6 days after ulceration caused by the administration of indomethacin (10 mg/kg). The effects of various EP agonists on the healing of the intestinal lesions were also examined in the presence of indomethacin (2 mg/kg/day). PGE$_2$ (1 mg/kg), 17-phenyl PGE$_2$ (EP1 agonist; 1 mg/kg), butaprost (EP2 agonist; 3 mg/kg), NT-012 (EP3 agonist; 3 mg/kg), or AE1-329 (EP4 agonist; 0.1 mg/kg) was given i.p. twice daily for 6 days, starting 1 day after induction of the intestinal lesions, whereas indomethacin (2 mg/kg) was given p.o. once daily for 6 days. The doses of the EP agonists were selected according to our previous studies (Takeuchi et al., 1997; Araki et al., 2000; Aoi et al., 2004). Control animals were given saline i.p. twice daily for 6 days of the experimental period. To reveal the damage in each case, 1 ml of Evans blue dye (w/w) was injected i.v. 30 min before sacrifice. Animals were killed under deep ether anesthesia, and the small intestine was excised. The tissue samples were then immersed in 10% neutralized formalin, embedded in paraffin, sectioned at 5 μm, and stained with H&E or Azan. Both macro- and microscopical observations were compared between normal rats giving saline in place of indomethacin (10 mg/kg), control rats given indomethacin (10 mg/kg) to produce intestinal ulcers, and the animals treated with various COX inhibitors after ulceration.

Determination of Mucosal PGE$_2$ Content. Levels of PGE$_2$ in the small intestinal mucosa were measured on various days after the administration of indomethacin (10 mg/kg). The animals were killed under deep ether anesthesia at various time points (3 h or 1, 2, 3, 5, or 7 days) after the administration of indomethacin, and small intestinal tissue was isolated, weighed, and placed in a tube containing 100% ethanol plus 0.1 M indomethacin (Tanaka et al., 2002a). Then, the tissue was homogenized with a Polytron homogenizer (IKA, Tokyo, Japan) and centrifuged at 10,000 rpm for 10 min at 4°C. After the supernatant of each sample had been evaporated with N$_2$ gas, the residue was resolved in assay buffer and used to determine the concentration of PGE$_2$. The concentration of PGE$_2$ was measured using a PGE$_2$ enzyme immunoassay kit (GE Healthcare, Little Chalfont, Buckinghamshire, UK). In separate experiments, the effect of indomethacin (2 mg/kg), SC560 (3 mg/kg), or rofecoxib (3 mg/kg) on the concentration of PGE$_2$ on day 3 or 6 after ulceration was examined. Each drug was given p.o. once daily for the first or last 3 days, starting 1 or 4 days after ulceration, respectively.

Analyses of COX-2 mRNA Expression by Reverse Transcription-Polymerase Chain Reaction. The animals were killed under deep ether anesthesia on days 1, 2, 3, 5, and 7 after the administration of indomethacin (10 mg/kg s.c.), and the small intestine was removed, frozen in liquid nitrogen, and stored at −80°C before use. Tissue samples were pooled from two to three rats for extraction of total RNA, which was prepared by a single-step acid phenol-chloroform extraction procedure using TRizol (Invitrogen, Gaithersburg, MD). Total RNA primed by a random hexa-deoxyribonucleotide was reverse-transcribed with the SuperScript preamplification kit (Invitrogen). The sequences of sense and antisense primers for rat COX-2 were 5'-TGATGACTGCTCAATCCCATG3'- and 5'-AATGTGAAAGTGCCTGCCACG-3', respectively, giving rise to a 702-base pair PCR product (Tso et al., 1985; Feng et al., 1993). For rat glyceraldehyde-3-phosphate dehydrogenase (G3PDH), a constitutively expressed gene, the sequence was 5'-GAACGG-GAGACTCAGGTCAAGGC-3' for the sense primer and 5'-TGGAGGTCCACACCTGGTCTG-3' for the antisense primer, giving rise to a 310-base pair PCR product (Feng et al., 1993). An aliquot of the reverse transcription reaction product served as a template in 35 cycles of PCR with 1 min of denaturation at 94°C, 0.5 min of annealing at 58°C, and 1 min of extension at 72°C on a thermal cycler. A portion of the PCR mixture was electrophoresed in 1.8% agarose gel in TAE buffer (40 mM Tris buffer, 2 mM EDTA, and 20 mM acetic acid, pH 8.1), and the gel was stained with ethidium bromide and photographed. The sequence of the PCR product was analyzed using the BLAST program (National Center for Biotechnology Information, Bethesda, MD).

Determination of Vascular Content Based on Carmine Incorporation. Vascular content was assessed based on the formation of vascular casts incorporating carmine, using a modification of the method of Kimura et al. (1986). The animals were given indomethacin (10 mg/kg s.c.) for the induction of intestinal lesions and killed 4 or 7 days later. Various COX inhibitors were given once daily for the first 3 days or the last 3 days. Under ether anesthesia, a cast was formed by the i.v. injection of 5% carmine red in 10% gelatin at 37°C in a volume of 1 ml/100 g body weight. The curcuses were chilled for 3 h at 4°C, and the damaged mucosa was dissected. After a thorough rinsing with phosphate buffer, pH 7.4, the tissue was weighed, minced with scissors, and dissolved in 3.5 N NaOH (5 ml/g tissue) at 37°C. The dye solution was then neutralized by adding 3 N HCl, centrifuged at 3000 rpm for 10 min at 4°C, and filtrated using a Millipore filter (0.45 μm). The dye content of 200-μl samples was
assayed spectrophotometrically using a 96-well plate reader (Hitachi, Mito, Japan) at 529 nm. The results were expressed as milligrams of dye content per gram of tissue.

Preparation of Drugs. The drugs used were indomethacin (Sigma-Aldrich, St. Louis, MO); SC560 (Cayman Chemical, Ann Arbor, MI); rofecoxib (synthesized in our laboratory); PGE₂ and 16,16-dimethyl PGE₂ (Funakoshi, Tokyo, Japan); 17-phenyl PGE₂ (Nacalai Tesque, Kyoto, Japan); butapropro, NT-012, AE1-329, and AE3-208 (Ono Pharmaceutical Co. Ltd., Osaka, Japan); and carmine red (Wako Pure Chemicals, Osaka, Japan). All COX inhibitors were suspended in a hydroxy propyl cellulose solution (Wako Pure Chemicals). PGE₂ and other EP receptor ligands were dissolved in absolute ethanol and then diluted with saline to the desired concentration. All drugs were prepared immediately before use and administered p.o., s.c., or i.p. in a volume of 0.5 ml/100 g body weight or i.v. in a volume of 0.1 ml/100 g body weight.

Statistics. Data are presented as the mean ± S.E. of four to nine rats per group. Statistical analyses were performed using the two-tailed Dunnett’s multiple comparison test, and values of \( P < 0.05 \) were considered significant.

Results

Development and Healing of Indomethacin-Induced Intestinal Ulcers

Subcutaneous administration of indomethacin (10 mg/kg) in normally fed rats produced multiple hemorrhagic lesions in the small intestine, mainly in the jejunum and ileum, the ulcer score at 24 h being 226.3 ± 18.2 mm². The ulcers healed quite rapidly within 7 days, and the ulcer score on days 3 and 7 was reduced to approximately 41.6 and 27.8%, respectively, of the initial score observed 24 h after indomethacin treatment (Fig. 1). Histological observation showed that the damage on day 1 was deep in the mucosa, where the epithelial cells were totally denuded and severe edema was observed in the submucosa (Fig. 2B). Consistent with the macroscopical observation, the area of damage became smaller with time (Fig. 2, C–F), and the damaged portion on day 7 was surrounded by granulation tissue and covered with a newly formed thin epithelium (Fig. 2F).

Gene Expression of COX Isozymes and PGE₂ Content in Intestinal Mucosa after Administration of Indomethacin

Indomethacin (10 mg/kg p.o.) markedly decreased the mucosal PGE₂ content of the small intestine from 7.8 ± 1.3 ng/g tissue to less than 1.3 ng/g tissue within 3 h. However, the level of PGE₂ was restored 24 h after the administration of indomethacin and even increased significantly above normal on days 2 to 3, before gradually returning to normal on day 7 (Fig. 3). The peak value was 35.8 ± 8.1 ng/g tissue, approximately 4 times greater than normal levels.

Consistent with our previous article (Tanaka et al., 2002a), indomethacin at an ulcerogenic dose (10 mg/kg) up-regulated the expression of COX-2 mRNA in the intestinal mucosa when examined 24 h after the administration, although the expression was not detectable in the normal rat intestine (Fig. 4). The expression of COX-2 was observed in the mucosa during the first 3 days, with a peak response on day 2, and it gradually faded away thereafter. By contrast, both G3PDH and COX-1 mRNAs were observed in the intestinal mucosa before and after the administration of indomethacin.

Effects of Various COX Inhibitors on Healing of Intestinal Lesions and Mucosal PGE₂ Content

As shown in Fig. 1, the intestinal ulcers produced by indomethacin (10 mg/kg p.o.) healed with time, and the area of
we examined the effect of COX inhibitors on the healing of the intestinal lesions at two different phases, the first 3 days (early phase) and the last 3 days (late phase).

The Early Phase. The area of intestinal lesions was reduced within 4 days to approximately one-half of that observed 24 h after the administration of indomethacin (10 mg/kg p.o.), the value being 82.6 ± 15.8 mm². When the animals were given indomethacin (2 mg/kg p.o.) once daily for the first 3 days after ulceration, the healing was impaired significantly, the value being 243.2 ± 27.1 mm² (Fig. 6). Daily administration of rofecoxib but not SC560 (3 mg/kg p.o.) for the first 3 days also significantly impaired the healing; the effect was almost the same as that of indomethacin. The healing was also significantly impaired by the combined administration of SC560 and rofecoxib. The impaired healing caused by indomethacin was antagonized by dmPGE₂ (20 μg/kg p.o.) given twice daily for 3 days, together with indomethacin. Consistent with the macroscopical observation, the lesions healed with newly formed epithelial cells on day 7 after ulceration (Fig. 7). In contrast, the damage in the animals given indomethacin for the first 3 days was still deep in the mucosa, without granulation tissue in the surrounding area. dmPGE₂ given together with indomethacin, however, restored the healing of these ulcers as revealed by histological examination, and the denuded area was covered with newly epithelial cells. On the other hand, the level of mucosal PGE₂ on day 4 was significantly greater than normal (Fig. 8). Indomethacin given for the first 3 days markedly reduced PGE₂ content to less than 5% of the control level. Likewise, a significant reduction in PGE₂ content was observed when rofecoxib but not SC560 was given for the first 3 days. Certainly, the combined administration of SC560 and rofecoxib markedly decreased the level of PGE₂ as effectively as did indomethacin.

![Fig. 4. Mucosal expression of COX-1 and COX-2 mRNAs after the induction of intestinal lesions in rats. The animals were given indomethacin (10 mg/kg s.c.) and killed various days (1, 2, 3, 5, and 7 days) after the administration. The expression of COX-1 and COX-2 mRNAs was analyzed by reverse transcription-PCR. Note that the expression of COX-2 mRNA was up-regulated in the intestinal mucosa after ulceration, the intensity being greatest during the first 3 days and gradually fading thereafter. N, normal rat (without indomethacin treatment).](image)

![Fig. 5. Effect of COX inhibitors on healing of indomethacin-induced small intestinal lesions in rats. The animals were given indomethacin (10 mg/kg s.c.) and killed 7 days later. Indomethacin (2 mg/kg), SC560 (3 mg/kg), rofecoxib (3 mg/kg; ROF), or SC560 plus rofecoxib was given p.o. once daily for 6 days, starting from 1 day after ulceration. Data are presented as the mean ± S.E. for eight rats. *, significant difference from control at P < 0.05.](image)

![Fig. 6. Effect of COX inhibitors on the early stage healing of indomethacin-induced small intestinal lesions in rats. The animals were given indomethacin (10 mg/kg s.c.) and killed 7 days later. Indomethacin (2 mg/kg), SC560 (3 mg/kg), rofecoxib (3 mg/kg; ROF), or SC560 plus rofecoxib was given p.o. once daily for 3 days, starting from 1 day after the induction of lesions. dmPGE₂ (20 μg/kg p.o.) was given twice daily for 3 days 30 min before and 6 h after indomethacin (2 mg/kg). Data are presented as the mean ± S.E. for seven to nine rats. Significant difference from control (+) and from vehicle (#) at P < 0.05.](image)
The Late Phase. On day 7 after ulceration, the area of lesions became much smaller, the value being 25.3 ± 11.6 mm², approximately 10% of that observed 24 h after the administration of indomethacin (10 mg/kg) (Fig. 9). When the animals were given various COX inhibitors once daily for the last 3 days, the healing of lesions was not affected by rofecoxib (3 mg/kg) alone but significantly impaired by either indomethacin (2 mg/kg), SC560 (3 mg/kg), or SC560 plus rofecoxib. The area of lesions in the latter three groups was roughly 150–200 mm², approximately 10 times greater than that of the control, whereas the rofecoxib-treated animals it was 28.4 ± 3.8 mm², almost equivalent to that of control animals. Levels of mucosal PGE₂ on day 7 remained slightly higher than normal (Fig. 10). Indomethacin (2 mg/kg) given for the last 3 days significantly decreased PGE₂ content to approximately one-third of that in the control. SC560 (3 mg/kg) or SC560 plus rofecoxib (3 mg/kg) also significantly reduced PGE₂ content, similar to indomethacin, whereas rofecoxib alone had no effect.

Effects of Various COX Inhibitors on Vascular Content

Effects of various COX inhibitors on vascular content were evaluated by measuring the amount of carmine incorporated in the vascular casts. The amount of carmine in the control group was 560.2 ± 21.3 μg/g tissue when determined 4 days after ulceration (Fig. 11A). Both indomethacin (2 mg/kg) and SC560 (3 mg/kg), ROF (3 mg/kg), or SC560 plus rofecoxib was given p.o. once daily for 3 days, starting from 4 days after the ulceration, and the animals were killed 3 h after the last administration. Data are presented as the mean ± S.E. for five to eight rats. *, significant difference from control at P < 0.05.
not shown). Indomethacin apparently suppressed angiogenesis in the ulcerated mucosa, whereas dmPGE2 given together with indomethacin restored the angiogenic response. On the other hand, the amount of carmine on day 7 was less than that observed on day 4, the value being $135.7 \pm 14.2 \mu g/g$ tissue (Fig. 11B). Indomethacin (2 mg/kg), given for the last 3 days, significantly decreased the content to $79.1 \pm 12.8 \mu g/g$ tissue, approximately one-half the control value. Likewise, the amount of carmine on day 7 was also significantly decreased by SC560 (3 mg/kg) but not rofecoxib (3 mg/kg) given for the last 3 days.

Effects of Various EP Agonists and Antagonists on the Healing of Intestinal Lesions

The healing of intestinal lesions was found to be markedly impaired when the production of PGE2 was suppressed by COX inhibitors. To further investigate which EP receptor is involved in the healing of these lesions, we examined the effect of various EP agonists on the impaired healing caused by indomethacin. As shown in Fig. 12A, the healing was markedly impaired by indomethacin (2 mg/kg) given for 6 days, the area of lesions being $243.1 \pm 29.4$ mm$^2$. This effect was significantly antagonized by either PGE2 (1 mg/kg) or the EP4 agonist AE1-329 (0.1 mg/kg) given twice daily for 6 days, and the area of lesions in both cases decreased to the control value. Other prostanoids, such as 17-phenyl PGE2 (EP1 agonist), butaprost (EP2 agonist), and NT-012 (EP3 agonist), did not affect the deleterious influence of indomethacin on healing.

To confirm the involvement of EP4 receptors in the healing of intestinal ulcers, we also examined the effect of AE3-208, the EP4 antagonist, on the healing in comparison with that of indomethacin. Daily administration of indomethacin (2 mg/kg p.o.) for 6 days markedly impaired the healing of intestinal ulcers, the area of lesions on day 7 being $118.4 \pm 8.7$ mm$^2$, approximately 5 times greater than that in the control (Fig. 12B). Likewise, AE3-208 (3 mg/kg i.p.) given for 6 days also impaired the healing of ulcers, as effectively as indomethacin, and the area of lesions ($117.9 \pm 23.5$ mm$^2$) was equivalent to that of the indomethacin-treated animals.

Discussion

It has been recognized for many years that NSAIDs hamper the spontaneous healing of peptic ulcers in experimental
animals and humans, whereas exogenously administered PGE\(_2\) accelerates the healing process (Wang et al., 1989; Levi et al., 1990; Schmassmann et al., 1995; Ukawa et al., 1998). Recent studies demonstrated an up-regulation of COX-2 expression, concomitant with an increase in the production of endogenous PGs, in the stomach after ulceration and suggested that COX-2/PGs play an important role in promoting the healing of gastric ulcers (Mizuno et al., 1997; Ukawa et al., 1998). However, it remains unknown whether the same is true for healing of intestinal ulcers. The present study showed for the first time that PGs are actively involved in the healing of indomethacin-induced intestinal ulcers, but the COX isozyme responsible for the production of PGs differs depending on the stage of the healing; COX-2 in the early stage and COX-1 in the late stage. In addition, it was suggested that the healing promoting action of PGs occurs through stimulation of the angiogenic response, probably mediated by the activation of EP4 receptors (Tanaka et al., 2005).

First, we confirmed that indomethacin at a dose of 10 mg/kg decreased mucosal PGE\(_2\) content within 3 h and caused severe hemorrhagic lesions in the small intestine, mainly in the jejunum and ileum, within 24 h. Consistent with a previous study dealing with the healing of gastric lesions induced by nectrotizing agents (Takeuchi et al., 1994), these lesions healed rapidly and were partially re-epithelialized within 7 days with granulation in the damaged portion. Indomethacin at the ulcerogenic dose up-regulated COX-2 expression in the intestinal mucosa, with a peak reached 2 days after the administration, and the PGE\(_2\) content showed a 4- to 5-fold increase over the control level, followed by a gradual return to normal. These results suggested an important role for COX-2/PGE\(_2\) in healing of intestinal ulcers, similar to that of gastric ulcers. Indeed, Mizuno et al. (1997) reported that induction of ulcers in the stomach was followed by potent expression of both COX-2 mRNA and protein, and the inhibition of PG production by the selective COX-2 inhibitor delayed the healing of gastric ulcers. The up-regulation of COX-2 expression was observed even in stomachs acutely damaged with 0.2 N HCl or ischemia-reperfusion (Kishimoto et al., 1997; Sawoaka et al., 1997). We also found in this study that the healing of intestinal ulcers was markedly impaired by indomethacin as well as rofecoxib given for 6 days after ulceration was induced, confirming the involvement of COX-2/PGs in the healing process. Interestingly, healing was similarly impaired when the animals were given SC560, a selective COX-1 inhibitor. These results suggest that COX-1/PGs are also involved in the healing of intestinal ulcers, in addition to COX-2/PGs. Brzozowski et al. (2001) showed that the selective COX-1 inhibitor resveratrol significantly delayed the healing of acetic acid-induced gastric ulcers in rats. Blickslager et al. (2002) also reported that the repair of intestinal epithelial damage after irradiation was significantly retarded in COX-1 knockout mice.

As shown in the present study, the up-regulation of COX-2 expression was observed during the early stage of the healing process, up to 3 days after ulceration was induced. Then, we examined the effect of selective COX inhibitors on the healing of intestinal ulcers by administering these drugs for the first 3 days or the last 3 days. As expected, the healing was impaired by both indomethacin and rofecoxib but not SC560 given for the initial 3 days, whereas it was delayed by both indomethacin and SC560 but not rofecoxib given for the final 3 days. Certainly, the combined administration of SC560 plus rofecoxib impaired the healing, similar to indomethacin, irrespective of whether they were given for the first or last 3 days. These results clearly indicate that endogenous PGs are actively involved in the healing of intestinal ulcers, yet the COX isozyme responsible for the production of PGs differs depending on the stage of the healing process; COX-2 in the early stage and COX-1 in the late stage. This idea was supported by the findings that the increased production of PGE\(_2\) on day 3 after ulceration was significantly inhibited by rofecoxib but not SC560, whereas the PGE\(_2\) content on day 7 was decreased by SC560 but not rofecoxib. Although the expression of COX-1 remained unchanged throughout the experimental period, it was observed that SC560, a selective COX-1 inhibitor, delayed healing in the late phase but not the early phase. In the early phase of healing, PGE\(_2\) production is mainly governed by COX-2 activity because of the up-regulation of COX-2, whereas in the late phase PGE\(_2\) production is mainly due to COX-1 activity because COX-2 expression subsided. That is why SC560, even if inhibiting COX-1 activity, did not reduce PGE\(_2\) content sufficiently for impairing the healing of intestinal lesions.

It should be noted that in the present study, following the administration of indomethacin or SC560 for the last 3 days, the ulcer score on day 7 was greater than that in the control on day 4. Similar results were obtained in the animals given indomethacin or rofecoxib for the first 3 days after ulceration was induced. These results suggest that such treatments not only impaired the healing but also exacerbated the ulcers. A deficiency of PGs due to inhibition of COX-1 causes a decrease in the secretion of mucus and an increase in intestinal motility, resulting in the facilitation of enterobacterial invasion and inflammatory responses (Tanaka et al., 2002b). Likewise, PGs produced by COX-2 have been shown to reduce inflammatory responses through inhibition of iNOS activity as well as neutrophil activation (Tanaka et al., 2002b). Thus, it is possible that inhibition of COX-1 for the last 3 days or COX-2 for the first 3 days causes an exacerbation of the intestinal damage, in addition to an interruption of the healing process. The present study demonstrated the importance of both COX-1 and COX-2 in the healing process of intestinal lesions and further showed that COX-2 is mainly involved in the early stage of healing, whereas COX-1 is involved in the late stage. Thus, healing of intestinal lesions was impaired when either COX-1 or COX-2 activity was inhibited by SC560 or rofecoxib, respectively. This is in contrast to the pathogenesis of NSAID-induced intestinal lesions. It is known that the intestinal ulcerogenic properties of NSAIDs are not accounted for solely by inhibition of COX-1 and require inhibition of COX-2 as well (Tanaka et al., 2002a). Inhibition of COX-1, despite causing intestinal hypermotility, bacterial invasion, and inducible nitric-oxide synthase expression, up-regulates the expression of COX-2, and the PGE\(_2\) produced by COX-2 counteracts deleterious events and maintains the mucosal integrity (Takeuchi et al., 2002; Tanaka et al., 2002b).

The healing of tissues involves multiple steps, including the formation of granulation tissue, contraction of the ulcerated tissue, and re-epithelialization, and these processes are regulated by growth factors, such as vascular endothelial growth factor (VEGF), basic fibroblast growth factor (bFGF), and other cytokines produced locally by regenerating cells.
are involved in the healing of intestinal ulcers through inter-
possible that several growth factors such as VEGF and bFGF
cells through p38 mitogen-activated protein kinase. It is thus
nous PGs produced not only by COX-2 but also by COX-1 and
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COX-1 and COX-2 are important for the regulation of angiogenesis. Blik-
s also observed that the expression of VEGF in the ulcerated
mosa was suppressed by the EP4 antagonist as well as rofecoxib, suggesting that PGE2 increases VEGF expression via EP4 receptors (Tanaka et al., 2005). In the present study, we observed that the EP4 antagonist AE3-208 impaired the healing of intestinal ulcers, similar to indomethacin or rofe-
oxib and that the EP4 agonist AE1-329 significantly antag-
onized the deleterious effect of indomethacin on the healing
These results suggest that PGE2 produced by both COX-1 and COX-2 promoted the healing of intestinal ulcers through the stimulation of angiogenesis mediated by EP4 receptors. However, Amano et al. (2003) showed that angiogenesis in a sponge implantation model was markedly sup-
pressed in mice lacking EP3 receptors with reduced expres-
sion of VEGF, suggesting the significance of PGE2/EP3
ceptor signaling in angiogenesis. Houchen et al. (2003) reported that PGE2 protected the intestinal mucosa against radiation-induced damage mediated by EP2 receptors through the suppression of apoptosis. At present, these discrep-
crepancies remain unexplained, yet they may be due to dif-
the present findings, it is concluded that endogenous
PGs produced by both COX-1 and COX-2 are involved in the healing of intestinal ulcers, but the COX isozyme responsible for the production of PGs differs depending on the phase of the healing process, COX-2 at the early phase and COX-1 at the late phase. In addition, PGE2 produced by COX-1 and COX-2 contributes to the healing of intestinal ulcers through stimulation of the angiogenic response, probably mediated by the activation of EP4 receptors. The precise mechanism by which PGE2 stimulates angiogenesis should be clarified. Fi-
ally, because the doses of various COX inhibitors given in this study are translated to those that are relevant to hu-
mans, the use of COX-1/COX-2 inhibitors needs to be redis-
cussed in patients with ulcers depending on the stage of healing

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References
287:G306–G310.
duodenal ulcers is not impaired by indomethacin or rofecoxib, the selective COX-2
Bliklager AT, Timmel DN, Young KM, Campbell NB, Little D, and Argenzio RA (2002) Recovery of ischemic injured porcine ileum: evidence for a contributory role of
Bliklager AT, Timmel DN, Young KM, Campbell NB, Little D, and Argenzio RA (2002) Recovery of ischemic injured porcine ileum: evidence for a contributory role of


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