Role of mast cells and protease-activated receptor-2 in cyclooxygenase-2 expression in urothelial cells

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Wang ZY, Wang P, Bjorling DE. Role of mast cells and protease-activated receptor-2 in cyclooxygenase-2 expression in urothelial cells. Am J Physiol Regul Integr Comp Physiol 297: R1127–R1135, 2009. First published August 12, 2009; doi:10.1152/ajpregu.00310.2009—Mast cells have been shown to play a role in development and persistence of various inflammatory bladder disorders. Mast cell-derived tryptase specifically activates protease-activated receptor-2 (PAR-2), and PAR-2 is known to be involved in inflammation. We investigated whether mast cells participate in increase of cyclooxygenase-2 (COX-2) protein abundance in urothelium/suburothelium of bladders of mice subsequent to cyclophosphamide (CYP)-induced bladder inflammation. We also used primary cultures of human urothelial cells to investigate cellular mechanisms underlying activation of PAR-2 resulting in increased COX-2 expression. We found that treatment of mice with CYP (150 mg/kg ip) increased COX-2 protein abundance in bladder urothelium/suburothelium 3, 6, and 24 h after CYP (P < 0.01), and increased COX-2 protein abundance was prevented by treatment of mice with the mast cell stabilizer sodium cromolyn (10 mg/kg ip) for 4 consecutive days before CYP treatment. Incubation of freshly isolated mouse urothelium/suburothelium with a selective PAR-2 agonist, 2-furoyl-LIGRLO-amide (3 μM), also increased COX-2 protein abundance (P < 0.05). We further demonstrated that 2-furoyl-LIGRLO-amide (3 μM) increased COX-2 mRNA expression and protein abundance in primary cultures of human urothelial cells (P < 0.01), and the effects of PAR-2 activation were mediated primarily by the ERK1/2 MAP kinase pathway. These data indicate that there are functional interactions among mast cells, PAR-2 activation, and increased expression of COX-2 in bladder inflammation.

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PAINFUL BLADDER SYNDROME/INTERSTITIAL CYSTITIS (PBS/IC) is a chronic inflammatory disorder characterized by frequency, urgency, and bladder pain (1, 19). Although the etiology and pathogenesis of PBS/IC remain unknown, substantial evidence indicates that mast cells play a significant role in development and persistence of PBS/IC in at least some patients (20, 33, 49). Mast cells are multifunctional immune cells that contain a variety of inflammatory mediators (7), and mast cell numbers have been reported to be increased in bladder biopsies from PBS/IC patients (32, 45, 56, 57). There is also a significant increase in urinary concentrations of tryptase (a marker of mast cell activation) in PBS/IC patients (5, 43, 57).

Protease-activated receptors (PARs) are a family of four G protein-coupled receptors (34, 51, 54). PARs are activated by cleavage of their extracellular NH2-terminal domain by proteolytic enzymes, and the exposed NH2-terminal sequence acts as a tethered ligand that binds and activates the receptors, initiating signaling cascades (34, 51, 54). Mast cell tryptase specifically activates PAR-2 (7, 21, 39, 44), and activation of PAR-2 induces release of proinflammatory neuropeptides, such as calcitonin gene-related peptide and substance P, from afferent neurons (52). A recent study demonstrated that activation of PAR-2 induced colonic inflammation in mice (8). PAR-2 has been shown to be present in bladders of various species (16, 17, 39), and inflammation increased expression of PAR-2 in rat bladders (17).

Several studies have shown that PAR-2 functionally interacts with cyclooxygenase-2 (COX-2) (13, 26). PAR-2-mediated relaxation of mouse tracheal was prevented by treatment by selective COX-2 inhibitors (26). Trypsin induces itching in mice by activating PAR-2, and this effect was selectively inhibited by the COX-2 inhibitor celecoxib (13). COX-2 is a key enzyme in conversion of arachidonic acid to proinflammatory prostanooids, such as prostaglandin E2 (PGE2) (38), and PGE2 plays an important role in hyperreactivity and pain associated with bladder inflammation (35).

Chemical cystitis induced by intraperitoneal injection of cyclophosphamide (CYP) in rodents is one of the most commonly used experimental models of bladder inflammation (4, 14). CYP treatment of rats increases expression of PAR-2 (17) and COX-2, as well as production of PGE2, in the bladder (23). CYP-induced COX-2 expression in bladder appears to occur primarily in the urothelium (11, 28). In the present study, we examined the functional interactions among mast cells, PAR-2, and increased COX-2 expression in the bladders of mice treated with CYP. We also used primary cultures of human urothelial cells to investigate cellular mechanisms underlying activation of PAR-2 resulting in increased COX-2 expression.

MATERIALS AND METHODS

Animals. C57BL/6J male mice were used at 3–6 mo of age. Experiments were conducted in accordance with National Institutes of Health guidelines, and all protocols were reviewed and approved by the Animal Care and Use Committee of the University of Wisconsin.

Induction of cystitis. CYP (150 mg/kg) was injected intraperitoneally (ip), and this dose of CYP induced cystitis of moderate severity (61). Control mice received an equivalent volume of sterile saline (0.9%, ip) instead of CYP. At the termination of experiments, mice were deeply anesthetized with pentobarbital sodium (50 mg/kg ip) and perfused through a cannula inserted into the left ventricle with 0.9% saline. Bladders were removed, and urothelium/suburothelium was mechanically separated from detrusor using fine forceps as described previously (28). Tissues were stored at −80°C until analyzed.

Treatment with sodium cromolyn. Sodium cromolyn stabilizes mast cell granules, thereby preventing release of chemical mediators from mast cells (53). Mice were treated with sodium cromolyn (10 mg/kg ip, given at 9:00 AM) for 4 consecutive days (13, 42, 48). Two hours after the last injection of sodium cromolyn, mice were treated with CYP or saline as described previously.

Culture of freshly isolated urothelium/suburothelium. Normal, untreated mice were deeply anesthetized with pentobarbital sodium (50...
mg/kg ip), and bladders were removed and cut into two halves longitudinally. The urothelium/suburothelium was mechanically separated from detrusor using fine forceps as described previously (28). The urothelium/suburothelium preparations were incubated in phenol red-free Ham’s F12 medium (Invitrogen, Carlsbad, CA) supplemented with 2% fetal bovine serum at 37°C in a CO2 incubator. One half of urothelium/suburothelium from each bladder was used as control (vehicle treated), and the another half was treated with the selective PAR-2 agonist 2-furoyl-LIGRLO-amide (37). Before treatment, the medium was changed every 30 min for 2 h. Preparations were then treated with either vehicle (phosphate-buffered saline, PBS) or 2-furoyl-LIGRLO-amide (3 μM) for 3 h. Tissues were rinsed briefly in PBS and stored at −80°C until analyzed.

Isolation of protein from urothelium/suburothelium. Tissues were homogenized with T-PER tissue protein extraction reagent (Thermo Scientific, Rockford, IL) containing protease inhibitors (Roche, Indianapolis, IN). Supernatants were collected by centrifugation at 10,000 g for 15 min at 4°C. Protein concentrations were determined using the BCA protein assay kit (Thermo Scientific). Protein samples were mixed 1:1 with Laemmli sample buffer (Bio-Rad, Hercules, CA), placed in boiling water for 5 min, and stored at −20°C until analyzed.

Culture of human urothelial cells. Acquisition and use of primary human urothelial cells was reviewed and approved by the University of Wisconsin Institutional Review Board. Primary urothelial cells were derived from a bladder specimen of a patient undergoing cystoplasty in the absence of cancer, active infection, or other bladder disorders (55). Phenol red-free Ham’s F12 medium (Invitrogen) supplemented with 0.1 μg/ml hydrocortisone, 5 μg/ml transferrin, 100 μg/ml penicillin, 100 μg/ml streptomycin, and 2% fetal bovine serum was used. The cells were used between three and six passages (55). Urothelial cells were treated with the selective PAR-2 agonist 2-furoyl-LIGRLO-amide (3 μM) or PBS for various periods of time to investigate the effects of PAR-2 activation on expression of PAR-2 and COX-2. In additional experiments, cells were treated with U0126 (10 μM), a specific inhibitor of phosphorylation of extracellular signal-regulated kinase (ERK1/2) of the mitogen-activated protein kinases (MAPK) family to investigate the role of ERK1/2 MAP kinase pathway activation in PAR-2 activation-induced expression of COX-2.

For mRNA analysis, cells were plated in 12-well plates (~50,000 cells/well). After treatment, medium was removed and cells were washed with PBS. Total RNA was extracted with Trizol reagent (Invitrogen) and treated with DNase I (Invitrogen) to remove genomic DNA. First-strand cDNA was generated using a cDNA synthesis kit according to the manufacturer’s instructions (Invitrogen). For immunoblotting, cells were plated in six-well plates (~100,000 cells/well). After treatments, medium was removed and cells were washed with PBS. M-PER mammalian protein extraction reagent (Thermo Scientific) was added, and cell lysates were collected. Supernatant were collected by centrifugation at 10,000 g for 15 min at 4°C. Protein concentrations were determined using the BCA protein assay kit (Thermo Scientific). The protein samples were mixed 1:1 with Laemmli sample buffer (Bio-Rad), placed in boiling water for 5 min, and stored at −20°C until analyzed.

Immunoblotting analysis. Protein samples (20 μg/lane) were resolved on 10% SDS-polyacrylamide gels and transferred to Immobilon-P polyvinylidene difluoride membranes (Fisher Scientific, Itasca, IL). Membranes were blocked in 5% dry fat-free milk in 1x TBST (20 mM Tris·HCl, 137 mM NaCl, 0.05% Tween 20; pH 7.5). After being rinsed, membranes were incubated at 4°C overnight with the specific primary antibody. Membranes were then washed free of primary antibody and incubated for 1 h with appropriate secondary antibody conjugated to horseradish peroxidase at room temperature. Signals were revealed using a chemiluminescence detection reagent (Amersham, Arlington Heights, IL). Membranes were exposed to X-ray films, and films were developed. Membranes were then stripped and rebotted with a mouse anti-glyceraldehyde-3-phosphate dehydrogenase (GAPDH) antibody as a loading control. Images were scanned and quantified using NIH Image 1.62. The primary antibodies used for immunoblotting were polyclonal rabbit anti-murine COX-2 (1:1,000) and monoclonal anti-human COX-2 (1:2,000) antibodies (Cayman Chemical, Ann Arbor, MI); monoclonal anti-GAPDH (1:5,000) antibody (Abcam, Cambridge, MA); polyclonal goat anti-PAR-2 antibody (1:500, reacts with both human and murine PAR-2); polyclonal rabbit anti-phospho-ERK1/2 (1:1,000), phospho-p38 (1:1,000), and pan-ERK1/2 (1:1,000) antibodies (Cell Signaling, Danvers, MA). The secondary antibodies were goat anti-rabbit IgG, goat anti-mouse IgG, and bovine anti-goat IgG (all conjugated to horseradish peroxidase and used at 1:20,000 dilution) (Santa Cruz, Biotechnology, Santa Cruz, CA).

Semi-quantitative analysis of COX-2 mRNA in urothelial cells. Real-time PCR was performed using an ABI 7300 thermocycler (Applied Biosystems, Foster City, CA). Samples were amplified in duplicate using the following thermal cycling conditions: 94°C for 10 min, followed by 45 cycles of amplification at 94°C for 30 s and then 60°C for 1 min to allow denaturing and annealing-extension. Abundance of PCR product was determined semi-quantitatively using a standard curve for each sample. Expression of COX-2 was normalized to abundance of mRNA for S26, a constitutively expressed ribosomal protein (Sigma) according to the manufacturer’s instructions (Invitrogen). For immunohistochemistry, the sections were blocked with 10% normal goat serum for 1 h and then a polyclonal anti-uroplakin antibody (generously provided by Dr. Tung-Tien Sun, New York University School of Medicine; used at 1:1,500, diluted in PBS containing 0.1% BSA, 0.3% Triton-X 100) (63) was applied. Slides were kept in a humid chamber overnight at 4°C, and staining was revealed using secondary goat anti-rabbit IgG conjugated with FITC (1:1,000; Sigma), St. Louis, MO), Slides were rinsed and coverslipped with an anti-fading solution (Vector Laboratories, Burlingame, CA).

Human urothelial cells were also stained for keratin to confirm their urothelial origin (6, 50). Cells were grown in Lab-Tek slide chambers (NUNC, Rochester, NY) for 24 h and were rinsed with PBS and fixed in cold acetone. They were rinsed and blocked with 10% normal goat serum. The monoclonal anti-AE1/AE3 keratin antibody (generously provided by Dr. Tung-Tien Sun, New York University School of Medicine; used at 1:1,500, diluted in PBS containing 0.1% BSA, 0.3% Triton-X 100) (63) was applied. Slides were kept in a humid chamber overnight at 4°C, and staining was revealed using secondary goat anti-rabbit IgG conjugated with FITC (1:1,000; Sigma), St. Louis, MO), Slides were rinsed and coverslipped with an anti-fading solution (Vector Laboratories). Slides were examined with a Nikon Eclipse microscope, and digital images were captured. Negative staining controls were prepared using normal mouse IgG instead of the specific antibody.

Reagents. Sodium cromolyn (dissolved with 0.9% saline) was purchased from Sigma. U0126 (dissolved with DMSO at 10 mM as the stock solution and diluted in PBS to desired concentration) and the selective PAR-2 agonist 2-furoyl-LIGRLO-amide (dissolved in PBS) (37) were obtained from EMD Chemicals (Gibbstown, NJ).

Statistical analysis. Data are arithmetic means ± SE. Data were analyzed using one-way ANOVA followed by Tukey’s post hoc multiple comparison test (GraphPad Prism, San Diego, CA). Student’s t-tests were also used when only two means were compared. A P value <0.05 was considered indicative of significant differences.

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[Correction: The above text contains a series of errors and inconsistencies that make it difficult to understand. It appears to be a mix of experimental results and procedural descriptions, possibly related to a specific research study.]
RESULTS

COX-2 protein abundance was increased in urothelium/suburothelium after induction of cystitis. Histological appearance of urothelium/suburothelium revealed that contamination of detrusor was very limited if there was any (Fig. 1A). Superficial urothelial cells were also stained strongly with a specific uroplakin antibody (Fig. 1B). COX-2 protein varies between 70 and 74 kDa on denaturing acrylamide gel, probably related to differences in glycosylated forms (28). Abundance of COX-2 protein in urothelium/suburothelium from normal control mice was relatively low, and this was consistent with previous reports (11, 28). As shown in Fig. 1C, the abundance of COX-2 protein in the urothelium/suburothelium was significantly increased 3 (16-fold), 6 (36-fold), and 24 h (32-fold) after treatment with CYP (150 mg/kg; n = 5, P < 0.01 vs. control at all time points examined), and increased COX-2 protein appeared to reach a maximum 6 h after CYP treatment.

PAR-2 protein is ~65 kDa on denaturing acrylamide gel (36). PAR-2 was present in urothelium/suburothelium from normal, control mice. As shown in Fig. 1D, abundance of PAR-2 protein remained unchanged 3 h (P > 0.05 vs. control), but was signifi-

Fig. 1. A: histological appearance (hematoxylin and eosin staining) of urothelium/suburothelium reveals that the contamination of detrusor was very limited. Scale bar, 50 μm. B: strong staining with a specific uroplakin antibody is shown in superficial urothelial cells. C: treatment of mice with cyclophosphamide (CYP; 150 mg/kg ip) for 3, 6, and 24 h induced an increase in cyclooxygenase-2 (COX-2) protein abundance in the bladder urothelium/suburothelium, respectively. D: similarly, treatment of mice with CYP induced an increase (6 and 24 h) in protease-activated receptor-2 (PAR-2) protein abundance in the bladder urothelium/suburothelium. Control mice received saline (ip). Relative COX-2 or PAR-2 protein abundance was analyzed by immunoblotting. Values determined by densitometry were normalized to those of GAPDH (loading control) in each sample. Data are means ± SE; n = 4–5. **P < 0.01 vs. control.
significantly increased 6 (4-fold, \( P < 0.01 \) vs. control) and 24 h (5-fold, \( P < 0.01 \) vs. control), after CYP treatment. These results demonstrate that bladder inflammation increased abundance of COX-2 and PAR-2 protein in the urothelium/suburothelium.

Treatment with sodium cromolyn inhibited increased COX-2 protein abundance. Mice were treated with sodium cromolyn (10 mg/kg) for 4 days to stabilize mast cells. Since the increase of COX-2 protein abundance reached a maximum 6 h after treatment with CYP, we examined COX-2 protein abundance 3 and 6 h after CYP treatment. As shown in Fig. 2, A and B, treatment with sodium cromolyn inhibited increased COX-2 abundance by 83 (\( n = 5, P < 0.01 \) vs. CYP-treated group) and 35% (\( n = 5, P < 0.05 \) vs. CYP-treated group) 3 and 6 h after CYP treatment, respectively. These results indicate that mast cells are involved in increased COX-2 protein abundance in response to CYP-induced bladder inflammation.

Treatment with PAR-2 agonist increased COX-2 protein abundance in isolated urothelium/suburothelium. Freshly isolated mouse urothelium/suburothelium was treated with the PAR-2 agonist 2-furoyl-LIGRLO-amide (3 \( \mu M \)) for 3 h. The relative COX-2 protein abundance was 0.1 \( \pm 0.02 \) in the control group and 0.72 \( \pm 0.19 \) in the PAR-2 agonist-treated group. 2-Furoyl-LIGRLO-amide significantly increased COX-2 protein abundance (7-fold; \( n = 6, P < 0.01 \)) 3 and 6 h after CYP treatment. As shown in Fig. 2, A and B, treatment with sodium cromolyn inhibited increased COX-2 protein abundance by 83 (\( n = 5, P < 0.01 \) vs. CYP-treated group) and 35% (\( n = 5, P < 0.05 \) vs. CYP-treated group) 3 and 6 h after CYP treatment, respectively. These results indicate that mast cells are involved in increased COX-2 protein abundance in response to CYP-induced bladder inflammation.

Activation of PAR-2 increased COX-2 expression, and this effect was mediated by the ERK1/2 kinase pathway in cultured primary human urothelial cells. As shown in Fig. 3A, urothelial cells were stained positively with the antibody against AE1/AE3 keratin, indicating that these cells retain their epithelial characteristics in culture (6, 50). The use of normal mouse IgG instead of primary keratin antibody abolished specific staining (Fig. 3B). PAR-2 was present in urothelial cells (Fig. 3C), and PAR-2 protein abundance was unaffected by treatment with the selective PAR-2 agonist 2-furoyl-LIGRLO-amide (3 \( \mu M \)) for 6 h (\( n = 4, P > 0.05 \)) (Fig. 3C).

2-Furoyl-LIGRLO-amide (3 \( \mu M \)) significantly increased COX-2 mRNA expression after 30 min and 1 and 2 h of treatment (\( n = 6, P < 0.01 \) vs. control; Fig. 4A). Three and six hours after treatment with 2-furoyl-LIGRLO-amide, COX-2 mRNA expression was not different from that observed in controls (\( n = 6, P > 0.05 \) vs. control) (Fig. 4A).

2-Furoyl-LIGRLO-amide (3 \( \mu M \)) also induced phosphorylation of ERK1/2 in a time-dependent manner (5–60 min), and this effect was abolished by pretreatment with U0126 (10 \( \mu M \)), an inhibitor of ERK1/2 phosphorylation, applied 15 min before 2-furoyl-LIGRLO-amide (Fig. 4B). Treatment with U0126 (10 \( \mu M \)) before exposure to 2-furoyl-LIGRLO-amide (3 \( \mu M \)) abolished increased COX-2 mRNA expression 30 min after treatment with 2-furoyl-LIGRLO-amide (\( n = 6, P < 0.01 \) vs. PAR-2 agonist-treated group) (Fig. 4C).

Similarly, treatment with 2-furoyl-LIGRLO-amide (3 \( \mu M \)) for 6 h increased COX-2 protein abundance by 61% (\( n = 6, P < 0.01 \) vs. control) (Fig. 4D). U0126 also inhibited PAR-2 agonist-induced increased COX-2 protein by 70% (\( n = 6, P < 0.05 \) vs. PAR-2 agonist-treated group) (Fig. 4D). These results indicate that activation of PAR-2 increased COX-2 mRNA expression and protein abundance, and this effect of PAR-2 activation was mediated by the ERK1/2 MAP kinase pathway. Treatment of cells with the PAR-2 agonist failed to induce
phosphorylation of p38 MAP kinase (data not shown), and p38 MAP kinase therefore does not appear to play a role in this process.

DISCUSSION

In the present study, we found that 1) treatment of mice with CYP increased COX-2 protein abundance in bladder urothelium/suburothelium; 2) mast cells were involved in inflammation-induced increase of COX-2 protein abundance; and 3) activation of PAR-2 increased COX-2 protein abundance in mouse urothelium/suburothelium. 4) Similarly, activation of PAR-2 increased COX-2 expression in cultured primary human urothelial cells; and 5) the effect of PAR-2 activation on COX-2 expression in cultured urothelial cells was mediated by the ERK1/2 MAP kinase pathway.

Although urothelium has historically been viewed as a simple barrier separating the bladder wall from urine, increasing evidence also suggests that the urothelium plays a critical role in physiological and pathophysiological processes in the bladder (2, 59). It has been demonstrated that urothelial cells have the capacity to secrete a variety of signaling molecules such as PGE₂, nerve growth factor, nitric oxide, and cytokines (59). These cells also express many signaling molecules similar to those in sensory afferent neurons, including purinergic receptors, vanilloid channels, and tyrosine kinase receptors (2, 59). Afferent nerve fibers within the bladder wall are located in close proximity to urothelial cells, suggesting that urothelial cells could be a target of neurotransmitters released from afferent nerves. Conversely, chemical mediators derived from urothelial cells could significantly influence the function of afferent nerve fibers (2, 59). Thus interaction between urothelial cells and afferent nerves may play a crucial role in the physiology and pathology of normal and pathological bladder function.

CYP is an antineoplastic alkylating agent commonly used to treat cancer patients, and an undesirable side effect of CYP is hemorrhagic cystitis (4, 14, 29, 61). CYP is metabolized by the liver to acrolein, and accumulated acrolein in urine causes cystitis (4, 14). CYP-induced cystitis has been used by many investigators to study mechanisms underlying bladder inflammation and associated visceral pain (4, 23, 29, 58, 61). COX-2 is an inducible enzyme that is upregulated during inflammation (11, 23, 38, 62); several studies have reported increased expression of COX-2 in rat bladders after treatment with CYP, and increased COX-2 protein appears to be primarily localized in the urothelium (11, 28). Furthermore, increased COX-2 protein in the bladder is accompanied by enhanced production of prostaglandins, particularly PGE₂ (23), and PGE₂ has been shown to cause bladder hyperreactivity by sensitizing afferent nerves (24, 36). In fact, increased bladder contractions induced by CYP, surgery, or lipopolysaccharide were reversed by a selective COX-2 inhibitor in rats (23, 30). These studies clearly demonstrated that COX-2 expression and subsequent increased PGE₂ release play a role in altered bladder functions in response to inflammation. Our study provides further evidence that treatment of mice with CYP increases COX-2 protein abundance in the urothelium/suburothelium, and a CYP-induced increase of COX-2 protein abundance is at least partly mediated by mast cells.
Mast cells are multifunctional immune cells that develop from a specific bone marrow progenitor and migrate into tissue perivascular spaces (25). Previous studies have shown an increase in mast cell numbers and activation in bladders of PBS/IC patients (56). Interestingly, mast cells isolated from bladders of PBS/IC patients are more responsive to various stimuli and release increased amounts of histamine (20). Histamine and tryptase levels are increased in the urine of PBS/IC patients (49). Together, these studies suggest that mast cells play an important role in the pathogenesis of PBS/IC (32, 33, 57). Mast cells contain granule-stored, presynthesized proinflammatory molecules and rapidly secrete mediators upon allergen exposure (22, 25). Mast cells are generally activated by specific antigens through cross-linking of IgE and high-affinity surface receptors (22, 25). However, mast cells also can be activated by nonimmunologic stimuli such as bacteria.

![Graph A](image)

**Fig. 4.** A: treatment of primary cultures of human urothelial cells with a selective PAR-2 agonist (3 μM) altered expression of COX-2 mRNA with significant increase at 30 min and at 1 and 2 h. Data are means ± SE; n = 6. **P < 0.01 vs. control. B: PAR-2 agonist (3 μM) induced phosphorylation of ERK1/2, which was abolished by U0126 (10 μM), a selective ERK1/2 inhibitor. A representative example of the results of 3 independently performed experiments is shown. C: treatment with U0126 (10 μM) prevented PAR-2 agonist-induced increase in expression of COX-2 mRNA at 1 h. Data are means ± SE; n = 6. **P < 0.01 vs. control. ##P < 0.01 vs. PAR-2 agonist-treated group. D: treatment of PAR-2 agonist (3 μM) for 6 h also increased COX-2 protein abundance, and this effect of PAR-2 was prevented by U0126 (10 μM). Data are means ± SE; n = 4. **P < 0.01 vs. control. #P < 0.05 vs. PAR-2 agonist-treated group.
including p38, ERK and c-Jun NH2-terminal kinase (9, 18), MAPK consist of three major families of protein kinases, that mediate signal transduction from the cell surface to the cell. Therefore, our results reveal abundance in urothelium. In addition, activation of PAR-2 is demonstrated that activation of PAR-2 increases COX-2 protein abundance are consistent with these findings. We further demonstrated that both ERK1/2 and p38 inhibitors (26). These studies suggest that both ERK1/2 and p38 inhibitors (27). Similarly, PAR-2-mediated relaxation and pain (12, 52, 54). Whereas other PARs are primarily activated by thrombin, PAR-2 is specifically activated by mast cell tryptase (51). Nakahara et al. (40) reported that tryptase and selective PAR-2 agonist induced contractions of isolated rat bladder strips, and the contractions were prevented by removal of the urothelium. Furthermore, the contractions were inhibited by indomethacin, a nonselective COX inhibitor (40). In another study by the same group, activation of PAR-2-induced detrusor contractions and release of PGE2 from urothelium were enhanced in rats treated with CYP (41). These results strongly suggest that 1) activation of PAR-2 increases release of prostaglandins from urothelium; 2) released prostaglandins cause detrusor contractions; and 3) PAR-2 mediated contractions are augmented during bladder inflammation. Our observations that PAR-2 was present in the urothelium and that treatment with CYP increased PAR-2 and COX-2 protein abundance are consistence with these findings. We further demonstrated that activation of PAR-2 increased COX-2 protein abundance in urothelium. In addition, activation of PAR-2 increased COX-2 mRNA expression and protein abundance in cultured human urothelial cells. Therefore, our results reveal that there are functional interactions between PAR-2 activation and COX-2 expression.

The MAPK are a group of protein serine/threonine kinases that mediate signal transduction from the cell surface to the nucleus in response to a variety of extracellular stimuli (9). The MAPK consist of three major families of protein kinases, including p38, ERK and c-Jun NH2-terminal kinase (9, 18), and activity of MAPK depends on their phosphorylation status (9, 18). Interestingly, treatment of rats with CYP induced phosphorylation of ERK1/2 in bladder (46), suggesting ERK1/2 kinase is involved in regulating bladder function. In human lung-derived A549 epithelial cells, activation of PAR-2 increased expression of COX-2 and enhanced release of PGE2 (27), and these effects were inhibited by both selective ERK1/2 and p38 inhibitors (27). Similarly, PAR-2-mediated relaxation of isolated mouse tracheal was prevented by selective ERK1/2 and p38 inhibitors (26). These studies suggest that both ERK1/2 and p38 pathways are involved in PAR-2-mediated expression of COX-2. In the present study, a PAR-2 agonist induced phosphorylation of ERK1/2, but not p38, in urothelial cells. Furthermore, increased COX-2 mRNA expression and protein abundance following PAR-2 activation were prevented by the selective ERK1/2 phosphorylation inhibitor U0126 (26).

Therefore, our results indicate that PAR-2-induced expression of COX-2 is primarily mediated by the ERK1/2 MAP kinase pathway in urothelial cells. These results also suggest that participation of MAP kinase pathways in PAR-2 signaling varies among different tissues or species.

**Perspectives and Significance**

Our findings demonstrate that COX-2 protein abundance is increased in mouse urothelium/suburothelium after induction of cystitis by CYP treatment and that the increase of COX-2 protein abundance is at least partially mediated by mast cells. We further demonstrate that activation of PAR-2 increases expression of COX-2 in primary cultures of human urothelial cells and mouse urothelium/suburothelium. Furthermore, the effects of PAR-2 activation in urothelial cells appear to be mediated primarily by the ERK1/2 MAP kinase pathway. These data indicate that there are functional interactions among mast cells, PAR-2 activation, and increased expression of COX-2. The results of the current study also raise intriguing questions about the convergent role of PAR-2 and COX-2 in bladder inflammatory diseases and suggest that both PAR-2 and COX-2 are likely therapeutic targets for the treatments of patients with PBS/IC. It should be noted that bladder inflammation is a complicated process and that other inflammatory mediators, such as cytokinins, neuropeptides and neurotrophic factors, also may contribute to regulating COX-2 expression, and this must be the focus of future studies.

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