Elevated cell migration, invasion and tumorigenicity in human KB carcinoma cells transfected with COX-2 cDNA

KAZUKI TAKAOKA1, HIROMITSU KISHIMOTO1, EMI SEGAWA1, SUSUMU HASHITANI1, YUSUKE ZUSHI1, KAZUMA NOGUCHI1, KAZUNARI SAKURAI1,2 and MASAHIRO URADE1

Departments of 1Oral and Maxillofacial Surgery, 2Surgical Pathology, Hyogo College of Medicine, 1-1 Mukogawa-cho, Nishinomiya, Hyogo 663-8501, Japan

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Abstract. In order to investigate the involvement of cyclooxygenase (COX)-2 in cell growth and invasion of oral cancer, a human epidermoid carcinoma cell line KB minimally expressing COX-2 protein was transfected with COX-2 cDNA and these activities were compared with mock-transfected KB in vitro and in vivo. KB/COX-2 clones showed a similar growth rate in vitro compared to KB/neo clones, but demonstrated significantly increased PGE2 production, cell migration and invasion. These KB/COX-2 clones markedly expressed MMP-9, pro-MMP-2 and activated-MMP-2 as compared to KB/neo clones in gelatin zymography. Western blot analysis showed that expression of MT1-MMP, Rho and Rac 1 in KB/COX-2 clones were stronger than that in KB/neo clones, but expression of TIMP-1 and TIMP-2 were weaker in KB/COX-2 clones than in KB/neo clones. When these cells were inoculated subcutaneously into nude mice, tumorigenicity and tumor growth were significantly elevated in KB/COX-2 tumors than in KB/neo tumors, and the gelatinase activity was much stronger in KB/COX-2 tumor tissues than in KB/neo tumor tissues in film in situ zymography. These results indicated that overexpression of COX-2 promotes cell migration in vitro and increases tumorigenicity, tumor growth and invasion of human KB carcinoma cells via up-regulated MMP and Rho family small GTPases and down-regulated TIMP activities.

Introduction

Cyclooxygenase (COX) is the rate-limiting enzyme in the biosynthesis of prostaglandins from arachidonic acid. Two isoforms of the COX enzyme exist, COX-1 and COX-2, and they have been postulated to be target molecules for non-steroidal anti-inflammatory drugs (NSAIDs). COX-1 is constitutively expressed in most tissues and appears to be a housekeeping enzyme responsible for various physiological functions, such as cytoprotection in the stomach, vasodilation in the kidney, and production of pro-aggregatory prostanoid thromboxane by platelets (1). On the other hand, COX-2 is induced by stimuli such as mitogens, cytokines, growth factors and tumor promoters, and is up-regulated at the sites of inflammation and in various cancer tissues; colon (2), stomach (3), breast (4), lung (5), esophagus (6), pancreas (7), urinary bladder (8), prostate (9) and skin (10). Former studies concerning involvement of COX-2 in cancer pathogenesis elucidated that COX-2 inhibits apoptosis (11) and immune surveillance (12), promotes angiogenesis (13), increases cancer invasiveness and metastasis (14) and regulates cell differentiation (15). In head and neck carcinoma several investigators have shown that expression of COX-2 is up-regulated (16-18), but its significance in relation to tumor cell migration, invasion and metastasis is not fully understood. Therefore, we investigated the effect of COX-2 overexpression in head and neck cancer cells on cell growth and invasive potentials in vitro and in vivo.

In this study, human KB carcinoma cell line minimally expressing COX-2 protein expression was transfected with COX-2 cDNA, and isolated clones with high COX-2 expression were compared with mock-infected clones in cell growth and invasion in vitro and in transplantable tumor model in nude mice. Consequently, it was indicated that overexpression of COX-2 promotes cell migration in vitro and increases tumorigenicity, tumor growth and local invasion in nude mice, via up-regulated matrix metalloproteinase (MMP), Rho family small GTPases and down-regulated tissue inhibitor of metalloproteinase (TIMP) activities.

Materials and methods

Cell lines and cell culture. Human KB carcinoma cell line (19) derived from epidermoid carcinoma of the floor of the mouth was used in this study. KB cells were grown in Dulbecco’s modified Eagle’s MEM (DMEM, Nissui Pharmaceutical Co., Tokyo, Japan) supplemented with 10% fetal bovine serum (HyClone Laboratories, Logan, UT, USA) and...
4 mM L-glutamine as growth medium at 37% in a 5% CO₂ incubator. This cell line was routinely subcultured with EDTA-trypsin mixture. In preliminary experiment, we found that KB cells lacked or minimally possessed COX-2 protein in Western blot analysis.

**COX-2 cDNA transfection.** Cells were transfected with pSG5-COX-2 plasmid containing a full-length human COX-2 cDNA (20) (a gift of Dr R. Kulmacz, University of Texas Medical School, Houston, TX) and pcDNA3 containing a neomycin-resistant marker by using calcium phosphate method as described previously (21). Positive transfectants were selected in DMEM containing 500 μg/ml genetecin (Cosmo Bio Co., Tokyo, Japan) and separate colonies were isolated using cloning cylinders.

**Cell growth assay.** Cells were plated at 2.5x10⁴ cells/well in a 100 μl volume in 96-well plates and cultured in growth medium at 37°C. Cell growth was assessed by 3(4,5-dimethylthiazolyl) 1,2-2,5-diphonyl tetrazolium bromide (MTT) assay after 1, 3, 5 and 7 days of incubation, as described previously (22).

**Prostaglandin E₃ (PGE₃) immunoassay.** Cells were plated at 2x10⁴ cells/well in 1 ml volume in 24-well plates and grown at 37°C for 24 h in growth medium. The medium was discarded and 1 ml of serum-free fresh medium was added to each well, and then conditioned medium was collected after 15 min. PGE₃ assay was performed as indicated in a protocol of prostaglandin E₃ EIA kit (Cayman Chemical, Ann Arbor, MI, USA), as described previously (22).

**Wound healing assay.** Cells were plated at 7.5x10⁴ cells/dish in 60-mm fibronectin-coated dishes (Asahi Techno Glass Co., Tokyo, Japan). After 6-h incubation, wound healing assay was performed. Scratch wounds were created by scraping confluent cell monolayers with a sterile pipette tip. After 24-h incubation, cell migration was quantified by measuring from wound edge to edge.

**Invasion assay.** Cell invasion assay was carried out using BioCoat Matrigel Invasion Chambers (Becton Dickinson Labware, Bedford, MA, USA) consisting of transwell membrane filter inserts in a 24-well tissue culture plate. The transwell filter has 8 μm pore size membrane coated with matrigel. Five thousand cells were seeded in the upper chamber of the transwell with serum-free DMEM, and DMEM containing 10% FBS were added to the lower chamber. After 24-h incubation, non-invading cells were removed by wiping the upper side of the membrane, and invaded cells were fixed and stained with Diff-Quick kit (Kokusai-shiyaku Co., Kobe, Japan). The number of invaded cells per membrane was counted under a light microscope at x200 magnification in each four fields of triplicate membranes.

**Western blot analysis.** Cells were lysed in a lysis buffer composed of Mg²⁺- and Ca²⁺- free phosphate-buffered saline (PBS) containing 20 mM Tris-HCl, pH 8.0, 1% NP40, 150 mM NaCl, 1 mM EDTA, 10% glycerol, 0.1% β-mercaptoethanol, 0.5 mM dithiothreitol, and a mixture of proteinase inhibitors consisting of 1 mM phenylmethylsulfonyl fluoride, 10 μg/ml aprotinin, 5 μg/ml leupeptin, 5 mM benzamidine, 1 μg/ml pepstatin, 2 μg/ml antipain hydrochloride (Boehringer, Mannheim, Germany), 50 mM 4-(2-aminoethyl)-benzene-sulfonyl fluoride hydrochloride (Wako Pure Chemical Industries, Ltd., Osaka, Japan), 2 mM sodium orthovanadate (Sigma-Aldrich Co., St. Louis, MO, USA), and 20 U/ml urokinase (Mochida Pharmaceutical, Tokyo, Japan). The lysate containing 15 μg protein was electrophoresed in a 10-20% gradient SDS-PAGE mini gel (Bio-Rad, Chicago, IL, USA) and blotted onto a PVDF membrane using Multiphor II (Amersham Pharmacia Biotech, Buckinghamshire, UK) for 30 min. The blotted membrane was blocked with 5% skim milk in 10 mM Tris-HCl, pH 7.2, containing 150 mM NaCl and 0.5% Tween-20 and incubated with primary antibodies (0.1-1 μg/ml) described below at 4°C for 16 h. The membrane was then incubated with alkaline phosphatase-conjugated secondary antibodies (0.02 μg/ml) described below for 4 h at room temperature. The membrane was rinsed, treated with nitroblue tetrazolium (Sigma-Aldrich) and 5-bromo-4-chloro-3-indolyl phosphate (Sigma-Aldrich) to visualize protein bands. The primary antibodies used were goat polyclonal antibody against COX-2, COX-1, RhoA and actin, and rabbit polyclonal antibody against Rac1 and Cdc42 (Santa Cruz Biotechnology, Santa Cruz, CA, USA), and mouse monoclonal antibodies against human MT1-MMP, TIMP-1, TIMP-2 (Daichi Fine Chemical Co. Ltd., Toyama, Japan). The secondary antibodies used were anti-goat, -rabbit or -mouse IgGs conjugated with alkaline phosphatase (Santa Cruz Biotechnology). Actin was used as internal control.

**Gelatin zymography.** The culture media of cells in subconfluent condition were replaced with serum-free DMEM, and the cells were cultured for further 24 h. The culture supernatants were collected as conditioned medium after centrifugation at 150 x g for 5 min to remove the debris. Because the culture supernatants contained only a small amount of protein, they were concentrated using a Centricon concentrator (Centricon YM-10; Millipore Co., Bedford, MA, USA) by centrifugation at 5000 x g for 30 min. The protein concentrated (10 μg) was mixed with non-reducing sample buffer (62.5 mM Tris-HCl, pH 6.8, 10% glycerol, 0.00125% bromophenol blue and 2% SDS) and electrophoresed at 4°C on a 7.5% polyacrylamide gel containing 0.1% SDS and 0.1% gelatin. After electrophoresis, the gel was washed twice with rinsing buffer (50 mM Tris-HCl, pH 7.5, 2.5% Triton X-100, 5 mM CaCl₂, 1 μM ZnCl₂, 0.05% NaN₃) at room temperature for 1 h to remove SDS. Strips of the gel were incubated at 37°C for 24 h in the reaction buffer for assaying gelatinolytic activity (50 mM Tris-HCl, pH 7.5, 5 mM CaCl₂, 1 μM ZnCl₂, 0.05% NaN₃) and then stained with Coomassie brilliant blue, 50% methanol and 20% acetic acid, and then destained in 30% methanol and 5% acetic acid. The proteolytic activity appeared as clear bands on blue background.

**Tumor formation in nude mice.** Cells (10⁵ or 10⁶) were inoculated subcutaneously into the flanks of 5-week-old female nude mice (BALB/C nu/nu; Oriental Yeast Ltd., Osaka, Japan). Tumorigenicity and tumor growth at the inoculated sites were examined and tumor size was measured by using
calipers once a week. The relative tumor weight (mg) was calculated by using the formula: \(a^2 \times b/2\), where \(a\) is the width in mm and \(b\) is the length in mm, according to the method of Battelle Columbus Laboratories (23). To examine the local tumor invasion, the orthotopic inoculation of 10^5 cells to the oral floor of nude mice was also performed.

**Histological and immunohistochemical studies.** Tissue specimens were fixed in 10% formalin, embedded in paraffin, and cut into 4-μm-thick sections according to conventional procedures. Histological examination was performed by H&E stain. For immunohistochemical studies, a modification of the streptavidin-biotin-peroxidase complex (SABC) method (24) was employed by using rabbit polyclonal antibody against human COX-2 (IBL, Gunma, Japan) as described previously (25). Non-immunized rabbit serum was used as negative control.

**Film in situ zymography.** To examine the localization of gelatinase activity in tumor tissues, film in situ zymography was performed. MMP in situ Zymo-Film (Fuji Photo Film Co., Ltd. Tokyo, Japan) coated with a gelatin base emulsion was used to detect the gelatinase activity in the underlying tissue. Also, MMP-PT in situ Zymo-Film coated with 1,10-phenanthroline (an MMP inhibitor) was used as the control. A 4-μm-thick frozen section was placed onto the film, incubated at 37˚C for 12 h, and then stained with 1% amido black (Nacalai, Kyoto, Japan). Gelatinolysis was detected as the disappearance of amido black staining.

**Statistical analysis.** Statistical analysis was done by using the Student's t-test. Differences were considered significant when the p-value was <0.05.

**Results**

**Isolation of cell clones overexpressing COX-2 protein.** Overexpression of COX-2 protein was seen in two clones among 5 separate clones transfected with COX-2 cDNA by Western blot. We isolated and designated these two clones as KB/COX-2 clone 1 (C1) and KB/COX-2 clone 4 (C4), and also isolated two neomycin-transfected clones, KB/neo clone 1 (N1) and KB/neo clone 2 (N2) as controls. KB/COX-2 clones showed about 3- to 4-fold increase of COX-2 protein expression and slightly increase of COX-1 protein expression as compared to KB/neo clones (Fig. 1). There was no apparent
morphological change between KB/COX-2 and KB/neo clones. In cell growth curve, KB/COX-2 and KB/neo clones showed a similar growth rate and there was no significant difference between them (Fig. 2). However, PGE2 production was much more in KB/COX-2 clones than KB/neo clone (N1, 5.2±1.2 pg/10^3 cells; N2, 4.3±1.3 pg/10^3 cells; C1, 7.3±1.8 pg/10^3 cells; C4, 17.1±2.8 pg/10^3 cells).

Increased cell migration in KB/COX-2 clones. To investigate the effect of COX-2 overexpression on tumor cell migration, wound healing assay and Matrigel invasion assay were employed. KB/COX-2 clones showed increased cell migration and filled scratch wound faster than did KB-neo clones (Fig. 3a). In addition, KB/COX-2 clones demonstrated a significantly increased ability to migrate through Matrigel invasion chambers as compared to KB/neo clones (Fig. 3b). Thus, COX-2 plays an important role in promoting cell migration of human KB carcinoma cells.

Elevated activity of metalloproteinases and Rho family small GTPases in KB/COX-2 clones. When gelatin zymography was performed using conditioned media from the cell clones, 92-, 72- and 62-kd bands exhibiting gelatin-degrading activity were detected. The 92-, 72-, and 62-kd bands were consistent with MMP-9, pro-MMP-2, and activated-MMP-2, respectively. These bands were thicker in KB/COX-2 clones than in KB/neo clones, indicating elevated activity of MMP-9, pro-MMP-2 and activated-MMP-2 in KB/COX-2 clones (Fig. 4a). As activated-MMP-2 band was markedly seen in KB/COX-2 clones, we examined the levels of MT1-MMP, which is known to cleave pro-MMP-2 to yield activated-MMP-2. As a result, expression of MT1-MMP was stronger in KB/COX-2 clones than in KB/neo clones by Western blot analysis (Fig. 4b). Since the activity of MMP-2 is controlled in at least two ways of proenzyme activation and inhibition of activity by tissue inhibitors of metalloproteinases (TIMPs), levels of TIMP-1 and TIMP-2 were examined, and found that expression of TIMP-1 and TIMP-2 was weaker in KB/COX-2 clones than in KB/neo clones (Fig. 4b). Rho family small GTPases such as RhoA, Rac1 and Cdc42 involved in cell migration and

Figure 4. Gelatin zymography and Western blot for MMPs and TIMPs in KB/COX-2 and KB/neo clones. (a) Gelatin zymography was performed to detect MMP-2 and MMP-9 using concentrated conditioned media of cell clones as described in the Materials and methods. (b) Expressions of MT1-MMP, TIMP-1, TIMP-2, RhoA, Rac1 and Cdc42 were examined by Western blot using monoclonal or polyclonal antibodies against those proteins.

Figure 5. Growth curves of tumors produced by KB/COX-2 (C4) and KB/neo (N1) clones in nude mice and immunohistochemical staining of COX-2. (a) Cells (10^6) were inoculated subcutaneously into the flanks of five 5-week-old female mice (BALB/C nu/nu). Tumors developed at the inoculated sites were measured by using calipers once a week. The relative tumor weight was determined, according to the method of Battelle Columbus Laboratories (23). ● N1; ■ C4. *p<0.05. (b) Immunohistochemical staining was performed by a modification of SABC method using anti-human COX-2 rabbit antiserum (IBL, Gunma, Japan). Original magnification, x200.
invasive phenotypes were next examined. Expression of RhoA and Rac1 protein were increased more in KB/COX-2 clones than in KB/neo clones by western blot analysis (Fig. 4b), but there was no difference in Cdc42.

Increased tumorigenicity of KB/COX-2 cells and local invasiveness of their tumors. The tumorigenicity and tumor growth rate in nude mice of KB/COX-2 clone 4 (C4) and KB-neo clone 1 (N1), which were most different in COX-2 expression, were examined by subcutaneous inoculation of cells. When 10^5 cells were inoculated subcutaneously in the flanks of six nude mice, C4 produced tumors in all mice, but N1 produced no tumors. Inoculation of 10^6 cells produced tumors in all five mice in both cell types. Tumors produced by C4 grew faster than those by N1 (Fig. 5a) and expressed COX-2 immunohistochemically (Fig. 5b). Film in situ zymography revealed that the gelatinase activity was much stronger in C4 tumor tissues than in N1 tumor tissues (Fig. 6a). This activity was diminished by the presence of 1,10-phenanthroline, an MMP inhibitor (Fig. 6b). To examine the local tumor invasiveness, both cell types were orthotopically inoculated to the oral floor. Consequently, C4 tumors showed invasive character with mandibular bone resorption, whereas N1 tumors showed demarcated margins (Fig. 7).

Discussion
Overexpression of COX-2 causes excess production of prostaglandins, and induces an increase of cell proliferation and decrease of apoptosis, mostly mediated by PGE2 and its receptor EP1-4 (26). Many studies have indicated significant involvement of COX-2 and PGE2 in carcinogenesis and progression for a variety of cancer (27-29). Furthermore, expression of COX-2 has been reported to contribute to malignant phenotypes such as cancer cell migration, invasion and metastasis (14,30,31). Although several studies have reported increased expression of COX-2 in head and neck carcinoma (16-18), it is not fully understood how COX-2...
contributes to malignant phenotype in squamous cell carcinoma in the majority of head and neck malignancies. Therefore, the present study was designed to investigate the effect of COX-2 overexpression on cancer cell migration, tumorigenicity and invasion, which are fundamental features of malignant phenotype, using the human epidermoid carcinoma KB cell line transfected with COX-2 cDNA. As a result, KB/COX-2 clones showed elevated PGE$_2$ production as compared to KB.neo clones, although cell growth rate of these clones was very similar. Nevertheless there was no significant difference in cell growth, cell migration and invasion in KB/COX-2 clones were significantly increased as compared to KB.neo clones, as demonstrated by wound healing assay and Matrigel invasion assay. Reorganization of the actin cytoskeleton of the cell is the primary mechanism of cell motility and is essential for most types of cell migration. Actin reorganization is regulated by Rho family small GTPases as Rho, Rac and Cdc42 (32). Slice et al (33) investigated that overexpression of constitutively active mutants of Rho and Rac, but not Cdc42 induced transcription from the COX-2 promoter. In this study, Western blot analysis showed that expression of RhoA and Rac1 but not of Cdc42 in KB/COX-2 clones were stronger than that in KB.neo clones, supporting the data of Slice et al (33).

Cancer cell invasion has been reported to be promoted by the activation of matrix metalloproteinases (MMPs), especially MMP-2 and MMP-9. MMPs belong to a family of enzymes that degrade the extracellular matrix (ECM) components and are known to contribute to tissue repair, tumor invasion and metastasis. Among them, MMP-2 is secreted as inactive proenzyme, pro-MMP-2. The activation of pro-MMP-2 is thought to take place on the cell surface by MT1-MMP. This process requires both active MT1-MMP and TIMP-2. The amino terminal of TIMP-2 binds to the catalytic domain of MT1-MMP forming a receptor, and the carboxyl terminal of TIMP-2 then interacts with the haemopexin domain of MMP-2 forming a ternary complex. The pro-MMP-2 localized on cell surface is cleaved by another molecule of active MT1-MMP. Intimate coordination of these proteins is therefore required to regulate the proteolytic activity of MMP-2. Several reports indicate that increased levels of MMP-2 correlate with invasive properties of certain tumor cell types (34-36). Tsuji et al (14) showed that overexpressing COX-2 increases MMP-2 expression/activity and cellular invasiveness in colon cancer cells. Thus, increased migration and invasiveness correlate with enhanced expression and activation of MMP-2. On the other hand, it has been shown that MMP-9 might play a more important role than MMP-2 in the invasive and metastatic potentials of squamous cell carcinoma of the head and neck (37-39). The correlation between increased TIMP-1 and TIMP-2 levels and decreased aggressiveness of tumors was found in some studies (40-43), but others have reported that TIMPs are up-regulated during invasion in head and neck carcinoma (44-46). Therefore, the expression of TIMPs in relation to the biological behavior of head and neck carcinoma is still controversial. In our study, KB/COX-2 clones with increased invasive activity demonstrated the activation of MT1-MMP, MMP-2 and MMP-9 and the inhibition of TIMP-1 and TIMP-2 expression. Since Philip et al (34) provided evidence that gelatinolytic activity is increased by inducing MT1-MMP expression via activation of the NF-kB pathway related to COX-2 transcription, the NF-kB pathway in KB/COX-2 clones is now under investigation.

We next examined the tumorigenicity of KB/COX-2 cells showing increased migration and elevated MMP activity in vitro and the invasive potential of tumors formed in nude mice. KB/COX-2 cells showed increased tumorigenicity and tumor growth rate although the growth of these cells was similar to that of KB.neo cells in vitro. In a recent experiment, we found that KB/COX-2 clone 4 (C4) produced higher amount of VEGF than did KB.neo clone 1 (N1) (data not shown). Therefore, it was speculated that tumor angiogenesis stimulated tumorigenicity and tumor growth in KB/COX-2 clone as compared to KB.neo clone. Furthermore, KB/COX-2 tumors demonstrated strong invasiveness with mandibular bone resorption in orthotopic inoculation. This finding was supported by the result of the film in situ zymography that the gelatinase activity was expressed stronger in KB/COX-2 tumor tissues than in KB.neo tumor tissues.

In summary, our results demonstrate that overexpression of COX-2 elevated cell migration in vitro and tumorigenicity and local tumor invasion in vivo via up-regulated MMP and Rho family small GTPases and down-regulated TIMP activities in human KB carcinoma cell line; hence it is strongly suggested that COX-2 regulating malignant phenotype could become a potent therapeutic target for head and neck carcinoma.

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