

Inhibition of Proinflammatory Tumor Necrosis Factor- α -Induced Inducible Nitric-Oxide Synthase by Xanthine-Based 7-[2-[4-(2-Chlorobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine (KMUP-1) and 7-[2-[4-(4-Nitrobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine (KMUP-3) in Rat Trachea: The Involvement of Soluble Guanylate Cyclase and Protein Kinase G

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ABSTRACT

In the study of anti-proinflammation by 7-[2-[4-(2-chlorobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine (KMUP-1) and 7-[2-[4-(4-nitrobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine (KMUP-3), exposure of rat tracheal smooth muscle cells (TSMCs) to tumor necrosis factor- α (TNF- α), a proinflammatory cytokine, increased the expression of inducible nitric-oxide synthase (iNOS) and NO production and decreased the expression of soluble guanylate cyclase α_1 (sGC α_1), soluble guanylate cyclase β_1 (sGC β_1), protein kinase G (PKG), and the release of cGMP in TSMCs. The cell-permeable cGMP analog 8-Br-cGMP, xanthine-based KMUP-1 and KMUP-3, and the phosphodiesterase 5 inhibitor zaprinast all inhibited TNF- α -induced increases of iNOS expression and NO levels and reversed TNF- α -induced decreases of sGC α_1 , sGC β_1 , and PKG expression. These results imply that cGMP enhancers could have anti-proinflammatory potential in TSMCs. TNF- α also increased

protein kinase A (PKA) expression and cAMP levels, cyclooxygenase-2 (COX-2) expression, and activated productions of prostaglandin (PG) E₂ and 6-keto-PGF₁ α (stable PGI₂ metabolite). Dexamethasone and *N*-[2-(cyclohexyloxy)-4-nitrophenyl]-methane sulfonamide (NS-398; a selective COX-2 inhibitor) attenuated TNF- α -induced expression of COX-2 and activated productions PGE₂ and PGI₂. However, KMUP-1 and KMUP-3 did not affect COX-2 activities and did not further enhance cAMP levels in the presence of TNF- α . It is suggested that TNF- α -induced increases of PKA expression and cAMP levels are mediated by releasing PGE₂ and PGI₂, the activation products of COX-2. In conclusion, xanthine-based KMUP-1 and KMUP-3 inhibit TNF- α -induced expression of iNOS in TSMCs, involving the sGC/cGMP/PKG expression pathway but without the involvement of COX-2.

Pro-inflammatory cytokines, including TNF- α , play an important role in regulating the tracheal smooth muscle contractility that is found in the asthmatic phenotype. TNF- α is in-

creased in the sputa of patients with bronchial asthma and present in the bronchoalveolar lavage fluid of symptomatic asthmatic patients (Renauld, 2001). As a member of these cytokines, TNF- α attracts and activates nonspecific inflammatory macrophages and neutrophils during infection and hypersensitivity induced by the inhalation of organic particles or fumes (Mohr, 2004; Mendez-Samperio et al., 2006).

Likewise, proinflammatory TNF- α , inducible nitric oxide-

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ABBREVIATIONS: TNF- α , tumor necrosis factor- α ; TSM, tracheal smooth muscle; COX-2, cyclooxygenase-2; iNOS, inducible nitric-oxide synthase; PDE, phosphodiesterase; PG, prostaglandin; PKA, protein kinase A; PKG, protein kinase G; KMUP-1, 7-[2-[4-(2-chlorobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine; KMUP-3, 7-[2-[4-(4-nitrobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine; sGC, soluble guanylate cyclase; TSMC, tracheal smooth muscle cell; NS-398, *N*-[2-(cyclohexyloxy)-4-nitrophenyl]-methane sulfonamide; IL, interleukin; TTBS, Tris-buffered saline/Tween 20; PBS, phosphate-buffered saline; PDEI, phosphodiesterase inhibitor; buffer A, Triton X-100 and bovine serum albumin in phosphate-buffered saline; CPT, chlorophenylthio; YC-1, 3-(5'-hydroxymethyl-2'-furyl)-1-benzylindazole; Bay-41-2272, 5-cyclopropyl-2-[1-(2-fluorobenzyl)-1*H*-pyrazolo[3,4-*b*]pyridin-3-yl]pyrimidin-4-ylamine; Rp-CPT-cAMPs, 8-(4-chlorophenylthio)adenosine-3',5'-cyclic monophosphorothioate, Rp-isomer; Rp-CPT-cGMPs, 8-(4-chlorophenylthio)guanosine-3',5'-cyclic monophosphorothioate, Rp-isomer.

synthase (iNOS), and cyclooxygenase-2 (COX-2) are coexpressed in pulmonary airway infection. Local production of TNF- α has been found to be regulated by iNOS and COX-2 and thus serves to orchestrate the inflammation pathway (Watkins et al., 1999). Enhanced COX-2 and iNOS expression by TNF- α can increase the production of cAMP and cGMP as a result of activated adenylate cyclase and guanylate cyclase, respectively. Because high-output cyclic nucleotide production in response to inflammation suppresses protein kinase G (PKG) expression, and cAMP analogs are more potent than cGMP analogs in reducing PKG mRNA expression, suggesting that PKA mediated the effects of cAMP and cGMP through cross-activation (Browner et al., 2004).

Xanthine-based phosphodiesterase inhibitors (PDEIs) have been used as bronchodilators because they can nonselectively inhibit PDE, resulting in increases of cAMP. Therefore, they have been used to treat bronchospastic diseases associated with immunoresponses (Giembycz, 2000; Caramori and Adcock, 2003). In contrast, xanthine-based 7-[2-[4-(2-chlorobenzene)piperazinyl] ethyl]-1,3-dimethylxanthine (KMUP-1), with endothelium- and epithelium-derived NO-enhancing activities, has been found to relax smooth muscle contractions by both activating sGC and inhibiting PDE, leading to the accumulation of cGMP (Wu et al., 2001, 2004; Lin et al., 2002). KMUP-1 has been shown to activate large-conductance Ca^{2+} -activated K^{+} channels in basilar artery myocytes via cAMP- and cGMP-dependent protein kinases (Wu et al., 2005). Furthermore, intratracheal 7-[2-[4-(4-nitrobenzene)piperazinyl]ethyl]-1,3-dimethylxanthine (KMUP-3) has been reported to increase the respiratory performance and to protect against TNF- α -induced airway resistance, involving the activation of sGC/cGMP/PKG (Lin et al., 2006). Previously, however, TNF- α was reported to modulate tracheal responses to G-protein-coupled receptor agonist and to act as an inflammatory cytokine in patients with asthma (Chen et al., 2003). Thus, it is still not known whether xanthine-based KMUP-1 and KMUP-3, chemically with imidazole (isoform of indazole) moiety, have anti-proinflammatory actions to modulate or attenuate TNF- α -induced iNOS and COX-2 expression, cGMP, and expression of PKG.

A nonxanthine sGC activator, YC-1, chemically with indazole moiety, exerts cGMP-dependent and -independent actions. The latter include the inhibition of PDE (Friebe et al., 1998) and untoward COX-2 expression in pulmonary epithelial cells (Chang et al., 2004). Thereafter, a more selective sGC activator, indazole-based Bay-41-2272, similar to YC-1, has also been reported to come with cGMP-dependent and -independent actions in vascular system (Teixeira et al., 2006). In addition, KMUP-1 and KMUP-3 were described previously to activate sGC in trachea (Wu et al., 2004; Lin et al., 2006). sGC activation by YC-1 and BAY-41-2272 has thus encouraged us to investigate the involvement of sGC/cGMP/PKG in associated anti-proinflammation by KMUP-1 and KMUP-3. The difficulty in using sGC activators is that they act nonselectively on several organs (Doggrell, 2005), which might limit their clinical potential, particularly in the pulmonary airway.

PDE5 inhibitors with cGMP-increasing activity have proven to induce tracheal relaxation. One of them, sildenafil, was found to induce endothelial nitric-oxide synthase and delay preconditioning through an iNOS-dependent pathway

(Salloum et al., 2003). However, proinflammatory iNOS is undesirable when researching new and safe tracheal relaxants. In contrast, classic xanthine-based PDEIs, theophylline and pentoxifylline, reduce TNF- α and iNOS expression and improve indomethacin-induced enteropathy, indicating their anti-proinflammatory merits (Saud et al., 2005). Xanthine-based KMUP-1 and KMUP-3 have been found to have cGMP-enhancing activity and thus may provide cGMP-dependent anti-proinflammatory benefits.

There are two classes of compounds: 1) indazole-based YC-1, BAY-41-2272, and sildenafil; and 2) imidazole-based KMUP-1, KMUP-3 (Fig. 1), and 8-Br-cGMP, predominantly mimic the action of cGMP by regulating one to three components of an endothelial nitric-oxide synthase/sGC/PDE-mixed functional enzyme system. However, in the present study, a cGMP-dependent bronchodilator with anti-inflammatory activity is desired to suppress TNF- α -induced bronchospastic dysfunction. The goal of this study was to develop a xanthine-based sGC activator/cGMP enhancer with tracheal relaxant and anti-inflammatory properties but devoid of YC-1's COX-2 and sildenafil's iNOS expression activity. So far, in addition to corticosteroids and β_2 -adrenoceptor agonists, neither xanthine-based PDEIs nor sGC activators are inhaled to attenuate inflammatory bronchospastic dysfunction.

This study aimed to investigate the anti-proinflammatory effects of xanthine-based KMUP-1 and KMUP-3 on TNF- α -induced expression of iNOS/COX-2 and the involvement of sGC/cGMP/PKG in tracheal smooth muscle cells (TSMCs).

Materials and Methods

Animals. Male Wistar rats (250–300 g) were provided by the National Laboratory Animal Breeding and Research Center (Taipei, Taiwan) and housed under constant temperature and controlled illumination. Food and water were available ad libitum. This study was approved by the Animal Care and Use Committee of the Kaohsiung Medical University.

Cell Culture. Rats were injected intraperitoneally with a lethal dose of pentobarbital. The tracheas were excised and cut longitudinally through the cartilage. Using a dissecting microscope, TSM strips were dissected from the surrounding parenchyma. The epithelium was removed from the luminal surface, and bands of TSM were gently separated from the underlying connective tissue. The TSM strips were then chopped into small sections (1 mm³) and incubated in Hanks' balanced salt solution (138 mM NaCl, 4 mM NaHCO₃, 5 mM KCl, 0.3 mM KH₂PO₄, 0.3 mM Na₂HPO₄, and 1.0 mM glucose)

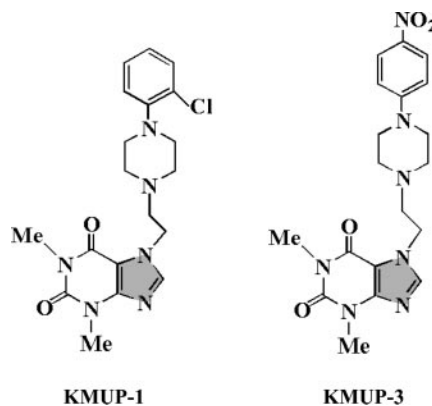


Fig. 1. Chemical structures of KMUP-1 and KMUP-3. The shadow ring indicates imidazole moiety.

with 0.05% elastase type IV and 0.2% collagenase type IV (Invitrogen, Carlsbad, CA) for 30 min at 37°C with gentle shaking. The solution of dissociated smooth muscle cells was centrifuged (6 min at 500g), and the pellet was resuspended in 1:1 Dulbecco's modified Eagle's medium/Ham's F-12 medium supplemented with 10% fetal bovine serum, 0.244% NaHCO₃, and 1% penicillin/streptomycin. Cells were cultured, with or without KMUP-1, KMUP-3, and other test agents, in 25-cm² flasks at 37°C in humidified air containing 5% CO₂. The medium was changed every 2 to 3 days. Confluent cells were passaged every third to fifth day onto 100-mm culture dishes after trypsinization. Experiments to measure levels of NO, cGMP, PGE₂, and PGI₂ were performed in cell cultures of 24-well plates with 10,000 cells per well. Determinations of iNOS, sGC, PKG, PKA, and COX-2 expressions were performed in cell cultures of 100-mm Petri dishes. The cells were used in passages 2 to 6.

Immunofluorescent Stain. According to our previous method (Wu et al., 2004), TSMCs (5 × 10³ cells/well) grown in 24-well plates on sterile 12-mm glass coverslips were rinsed twice with ice-cold phosphate-buffered saline (PBS) and fixed with fixing buffer (3% formaldehyde and 2% sucrose in PBS) for 5 min. Samples were washed with washing buffer (0.1 M glycine in PBS), and the cells were permeabilized for 20 min with permeabilization buffer (10% goat serum and 0.4% Triton X-100 in PBS). Samples were then washed with buffer A (0.2% Triton X-100 and 0.2% bovine serum albumin in PBS) and incubated with fluorescein isothiocyanate-conjugated monoclonal mouse anti-smooth muscle α -actin antibody for 2 h in a dark room. The resulting samples were washed with buffer A, aqueous fluorescent mounting media were added, and then coverslips were affixed to the slides. Finally, the coverslips were allowed to dry in the dark before viewing.

Protein Extraction and Western Blot Analysis. After incubation with or without KMUP-1, KMUP-3, and other test agents, the cells were washed rapidly with ice-cold PBS, incubated with protein extraction reagent (Pierce Biotechnology, Inc., Rockford, IL), and then cells were scraped from plates. The samples were sonicated for 10 s three times and centrifuged at 13,000 rpm at 4°C for 30 min. The protein concentrations of supernatants were determined by using bovine serum albumin as the standard. The cell extracts were then boiled in ratio of 4:1 with sample buffer (100 mM Tris, pH 6.8, 20% glycerol, 4% SDS, and 0.2% bromophenol blue). Electrophoresis was performed using 10% SDS-polyacrylamide gel electrophoresis and transferred to nitrocellulose membranes (Millipore Corporation, Billerica, MA). The membrane was blocked with Tris-buffered saline (20 mM Tris and 137 mM NaCl, pH 7.6) containing 0.1% Tween 20 (TTBS) and 5% nonfat milk at room temperature for 1 h, washed with TTBS, and then incubated overnight at 4°C in the appropriate primary antibody of COX-2, iNOS, PKA_{RI}, PKG_{1 α 1 β} , sGC α ₁, and sGC β ₁. The membranes were washed in TTBS before being incubated with horseradish peroxidase-conjugated antibody against mouse, goat, or rabbit IgG for 1 h. The membrane was then washed in TTBS and developed with the enhanced chemiluminescence for the detection of the specific antigen. The intensity of the bands was quantitated by densitometry.

Cyclic Nucleotide Radioimmunoassay. Intracellular concentrations of cAMP and cGMP in TSMC were measured according to our previous reports (Wu et al., 2004; Lin et al., 2006). In brief, cells were grown in 24-well plates (10⁵ cells/well) with or without KMUP-1, KMUP-3, and other test agents. At confluence, monolayer cells were washed with PBS and then incubated with isoproterenol, KMUP-1, KMUP-3, and zaprinast (10 μ M) for 20 min at 37°C in medium and further incubated in the absence or presence of TNF- α (100 ng/ml) for 9 h. The reaction was terminated by replacement of medium with 1 ml of ice-cold 1 N hydrochloric acid. Cell suspensions were sonicated and then centrifuged at 2500 g for 15 min at 4°C. Then, the supernatants were lyophilized, and the concentrations of cAMP and cGMP of each sample were measured using cGMP-¹²⁵I and cAMP-¹²⁵I radioimmunoassay kits (GE Healthcare, Little Chalfont, Buckinghamshire, UK).

Prostaglandin Radioimmunoactivity. After treatment with cGMP-enhancing KMUP-1 and KMUP-3, the culture media were collected. Concentrations of PGE₂ and 6-keto-PGF_{1 α} (stable metabolite of PGI₂) in the culture medium were measured using PGE₂ and 6-keto-PGF_{1 α} EIA assay kits (Cayman Chemical Co., Ann Arbor, MI). All samples were run in duplicate.

Measurement of Nitrite/Nitrate. After treatment with KMUP-1 and KMUP-3, the culture media were collected. Both nitrite and nitrate (stable breakdown product of NO) were measured in cell culture media. All samples were analyzed using nitrite/nitrate colorimetric assay kits (Cayman Chemical Co.) and run in duplicate.

Drugs and Chemicals. KMUP-1 and KMUP-3 were synthesized in this laboratory. 8-Br-cGMP, dexamethasone, indomethacin, isoproterenol, NS-398, Rp-8-CPT-cAMPS, Rp-8-CPT-cGMPS, TNF- α , and zaprinast were all purchased from Sigma-Aldrich (St. Louis, MO). COX-2, iNOS, sGC α ₁, and sGC β ₁ antibodies were obtained from Santa Cruz Biotechnology (Santa Cruz, CA). PKA_{RI} and PKG_{1 α 1 β} antibodies were purchased from Calbiochem (San Diego, CA). All drugs and reagents were dissolved in distilled water unless otherwise noted. KMUP-1 and KMUP-3 (10⁻² M) were dissolved in 10% absolute alcohol, 10% propylene glycol, and 2% 1 N HCl. Dexamethasone, indomethacin, and zaprinast (10⁻² M) were dissolved in dimethyl sulfoxide. Serial dilutions were made by distilled water.

Statistical Analysis. All data are expressed as the mean \pm S.E. Statistical differences were determined by independent and paired Student's *t* test in unpaired and paired samples, respectively. Whenever a control group was compared with more than one treated group, the one-way or two-way analysis of variance was used. When the analysis of variance manifested a statistical difference, results were further analyzed with Dunnett's or Tukey test. A probability value (*P* value) less than 0.05 was considered to be significant. Analysis of the data and plotting of the figures were done with the aid of SigmaPlot software (version 8.0) and SigmaStat (version 2.03; both from Systat Software, Point Richmond, CA) run on an IBM-compatible computer.

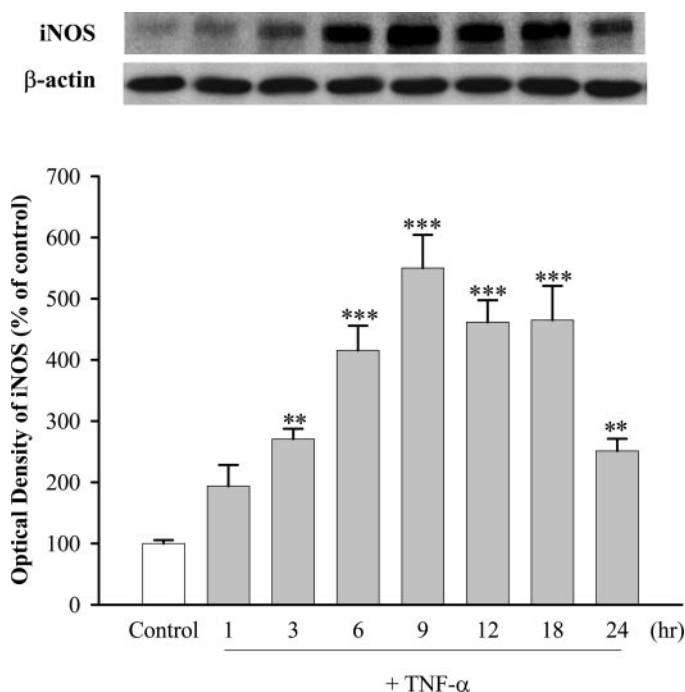


Fig. 2. TNF- α modulated the expression of iNOS in cultured TSMCs. Cells treated with TNF- α (100 ng/ml) for 1, 3, 6, 9, 12, 18, and 24 h. Values are means \pm S.E., *n* = 6. **, *P* < 0.01, ***, *P* < 0.001 versus control.

Results

iNOS and sGC Expression. To investigate the effects of TNF- α on iNOS and sGC protein, TSMCs were incubated with TNF- α (100 ng/ml) for 1, 3, 6, 9, 12, 18, and 24 h, and the levels of iNOS and sGC subunit proteins were measured by immunoblotting. As shown in Fig. 2, exposure of TSMCs to TNF- α increased iNOS protein expression in a time-dependent manner, with the maximum level evident at 9 h. KMUP-1 and KMUP-3 (0.01–100 μ M; Fig. 3A) and the PDE5 inhibitor zaprinast and exogenous 8-Br-cGMP (10 μ M; Fig. 3B) inhibited TNF- α -induced increases of iNOS expression to a similar extent (Fig. 3). In contrast, TNF- α decreased the expression of sGC α_1 and sGC β_1 proteins in a time-dependent manner, with the maximal inhibition achieved at 9 h (Fig. 4). These inhibitions were reversed by KMUP-1, KMUP-3, zaprinast, and 8-Br-cGMP at 10 μ M (Fig. 5).

PKG and PKA Expression. TNF- α (100 ng/ml) time-dependently increased PKA within 24 h and significant decreased the expression of PKG protein between 6 and 9 h

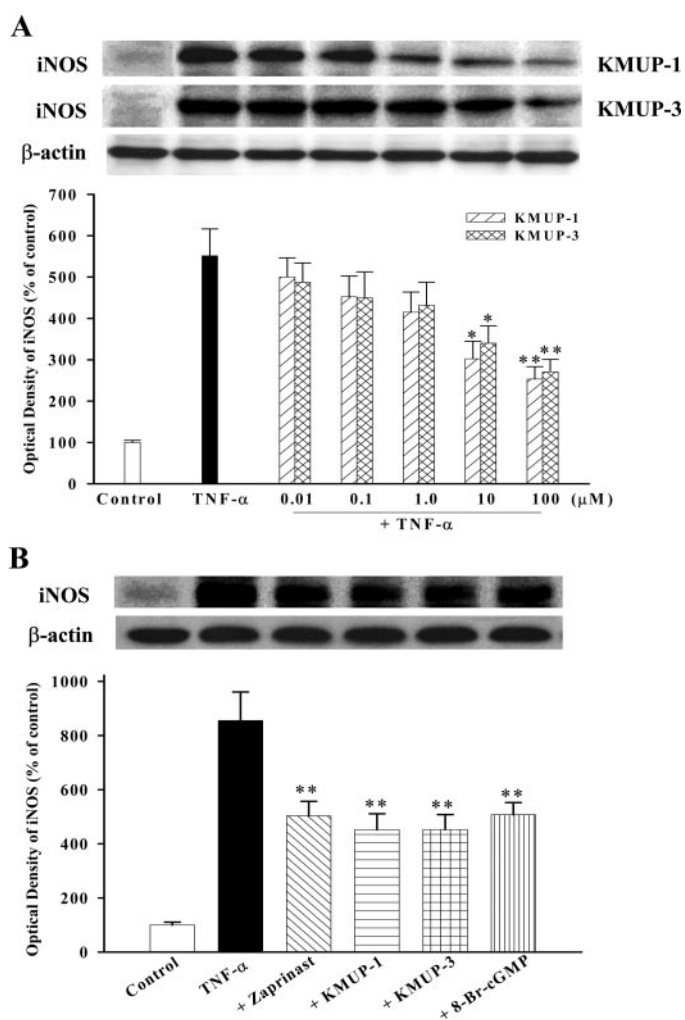


Fig. 3. Effects of KMUP-1, KMUP-3, and various agents on TNF- α -induced iNOS expression in cultured TSMCs. A, concentration-dependent effects of KMUP-1 and KMUP-3 (0.01–100 μ M) on TNF- α (100 ng/ml)-induced iNOS expression after 9-h incubation. *, $P < 0.05$; **, $P < 0.01$ versus TNF- α -treated cells. B, cells pretreated with zaprinast, KMUP-1, KMUP-3, and 8-Br-cGMP (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 9 h. Values are means \pm S.E., $n = 6$. **, $P < 0.01$ versus TNF- α -treated cells.

(Fig. 6). The increased expression of PKA protein by TNF- α was not further increased by KMUP-1 and KMUP-3 (0.1–100 μ M); however, decreases of PKG protein expression were reversed by both KMUP-1 and KMUP-3 (Figs. 7 and 8). Zaprinast and 8-Br-cGMP at 10 μ M also reversed TNF- α -induced decreases of PKG protein (Fig. 8A), whereas 8-Br-cAMP and zaprinast at 10 μ M, similar to KMUP-1 and

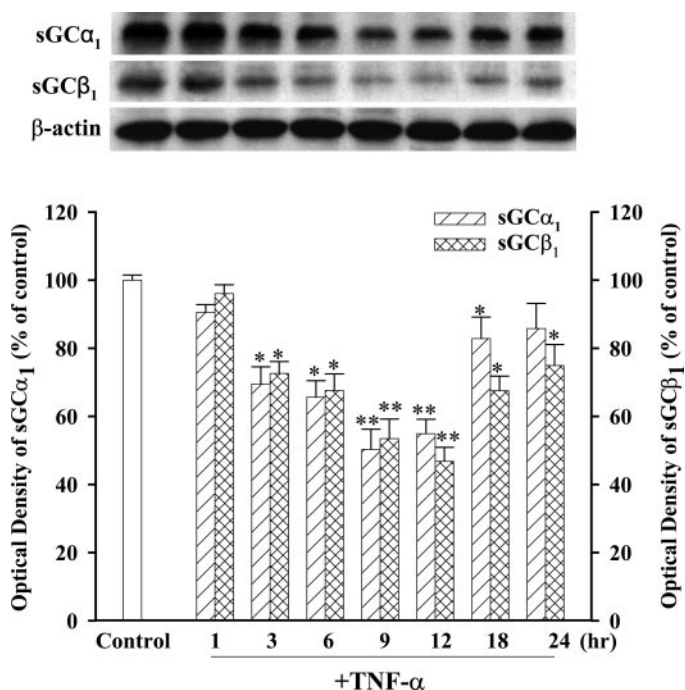


Fig. 4. TNF- α modulated the expression of sGC α_1 and sGC β_1 in cultured TSMCs. Cells treated with TNF- α (100 ng/ml) for 1, 3, 6, 9, 12, 18, and 24 h. Values are means \pm S.E., $n = 6$. *, $P < 0.05$; **, $P < 0.01$ versus control.

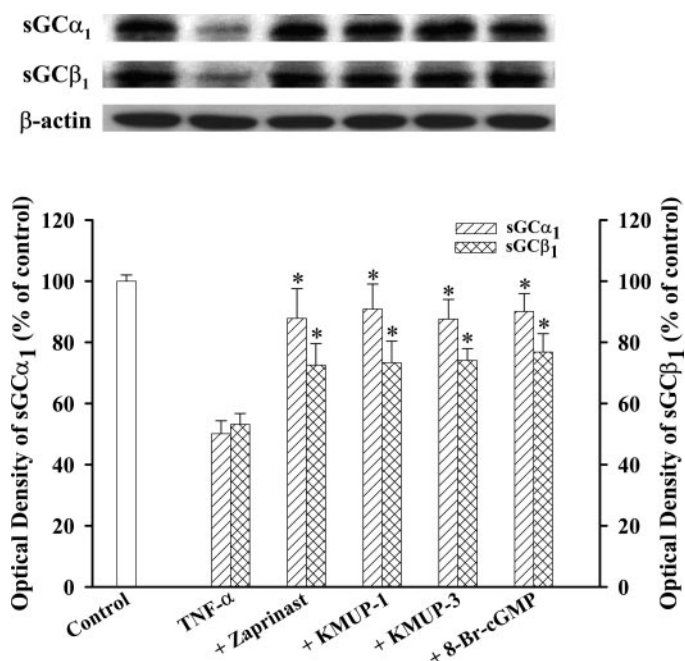


Fig. 5. Effects of KMUP-1, KMUP-3, and various agents on TNF- α -inhibited sGC α_1 or sGC β_1 expression in cultured TSMCs. Cells pretreated with zaprinast, KMUP-1, KMUP-3, and 8-Br-cGMP (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 9 h. Values are means \pm S.E., $n = 6$. *, $P < 0.05$ versus TNF- α -treated cells.

KMUP-3, did not affect TNF- α -induced increases of PKA protein (Fig. 8B). The addition of the PKG inhibitor Rp-8-CPT-cGMPs had no effect on TNF- α -induced decreases of PKG expression. In contrast, the PKA inhibitor Rp-8-CPT-cAMPS reversed the capacity of TNF- α to down-regulate PKG expression (Fig. 8C). These results suggest that the activation of PKA by TNF- α down-regulates PKG expression in rat TSMCs.

COX-2 Expression. In the absence of TNF- α in rat TSMCs, the expression of COX-2 showed time-dependent decreases during incubation for 24 h. In the presence of TNF- α (100 ng/ml), the expressions of COX-2 were limited to moderate decreases after 6 h and were sustained for 24 h compared with non-TNF- α challenge groups (Fig. 9A). It was clear that not KMUP-1 and KMUP-3 (10 μ M) but dexamethasone (1 μ M) and COX-2 inhibitor NS-398 (10 μ M) can significantly inhibit TNF- α -induced COX-2 expression in TSMCs (Fig. 9B).

Cyclic Nucleotide Levels. Intracellular cGMP production was decreased, reaching minimal production at 9 h in TSMCs in the presence of TNF- α (100 ng/ml). However, in the presence of KMUP-1, KMUP-3, and zaprinast (10 μ M), cGMP was reversed to the basal level (Fig. 10A). In the absence of TNF- α , the use of KMUP-1, KMUP-3, and isoproterenol (10 μ M) in the culture of TSMCs significantly increased the production of cAMP compared with the control group. However, in the presence of TNF- α , KMUP-1, and KMUP-3 could not further increase the production of cAMP compared with the control group (Fig. 10B).

NO Levels Measured by Nitrite/Nitrate. TSMCs were incubated with TNF- α for 9 h. Exposure of TSMCs to TNF- α (100 ng/ml) led to the accumulation of nitrite and nitrate in the culture medium, indicating the release of NO. 8-Br-cGMP, KMUP-1, KMUP-3, and zaprinast (10 μ M) all inhibited

ited TNF- α -induced production of nitrite/nitrate, which represented the NO levels (Fig. 11).

PGs and COX-2 Activities. Incubation of TSMCs with TNF- α (100 ng/ml) increased the releases of PGs, PGE₂, and 6-keto-PGF₁ α , a PGI₂ stable metabolite. The greatest increases that occurred were measured at 24 h. Dexamethasone (1 μ M) and NS-398 (10 μ M), a selective COX-2 inhibitor, significantly inhibited TNF- α -induced PGE₂ and 6-keto-PGF₁ α formation. However, KMUP-1 and KMUP-3 (10 μ M) did not inhibit the production of PGs, resulting from the activation of COX-2 (Fig. 12).

Discussion

Bronchodilation and immunosuppression or anti-inflammation activities are important in drug therapy for inflam-

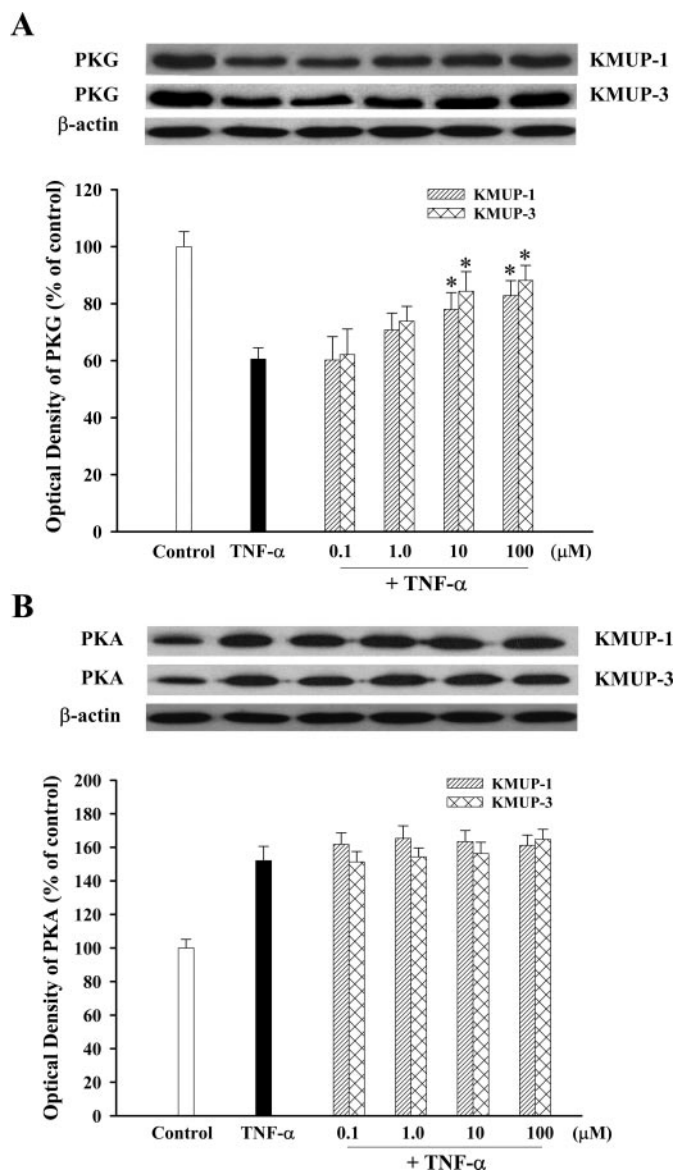


Fig. 7. Effects of KMUP-1 and KMUP-3 on TNF- α -modulated PKG or PKA expression in cultured TSMCs. A, concentration-dependent effects of KMUP-1 and KMUP-3 (0.1–100 μ M) on TNF- α (100 ng/ml)-regulated PKG expression after 9-h incubation. *, $P < 0.05$ versus TNF- α -treated cells. B, KMUP-1 and KMUP-3 (0.1–100 μ M) on TNF- α (100 ng/ml)-regulated PKA expression after 24-h incubation. Values are means \pm S.E., $n = 6$.

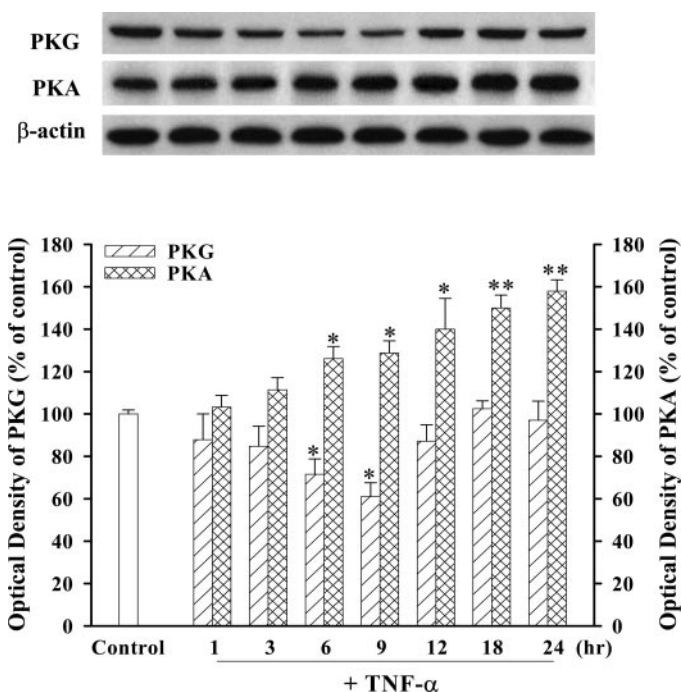


Fig. 6. TNF- α modulated the expression of PKG and PKA in cultured TSMCs. Cells treated with TNF- α (100 ng/ml) for 1, 3, 6, 9, 12, 18, and 24 h. Values are means \pm S.E., $n = 6$. *, $P < 0.05$ versus control.

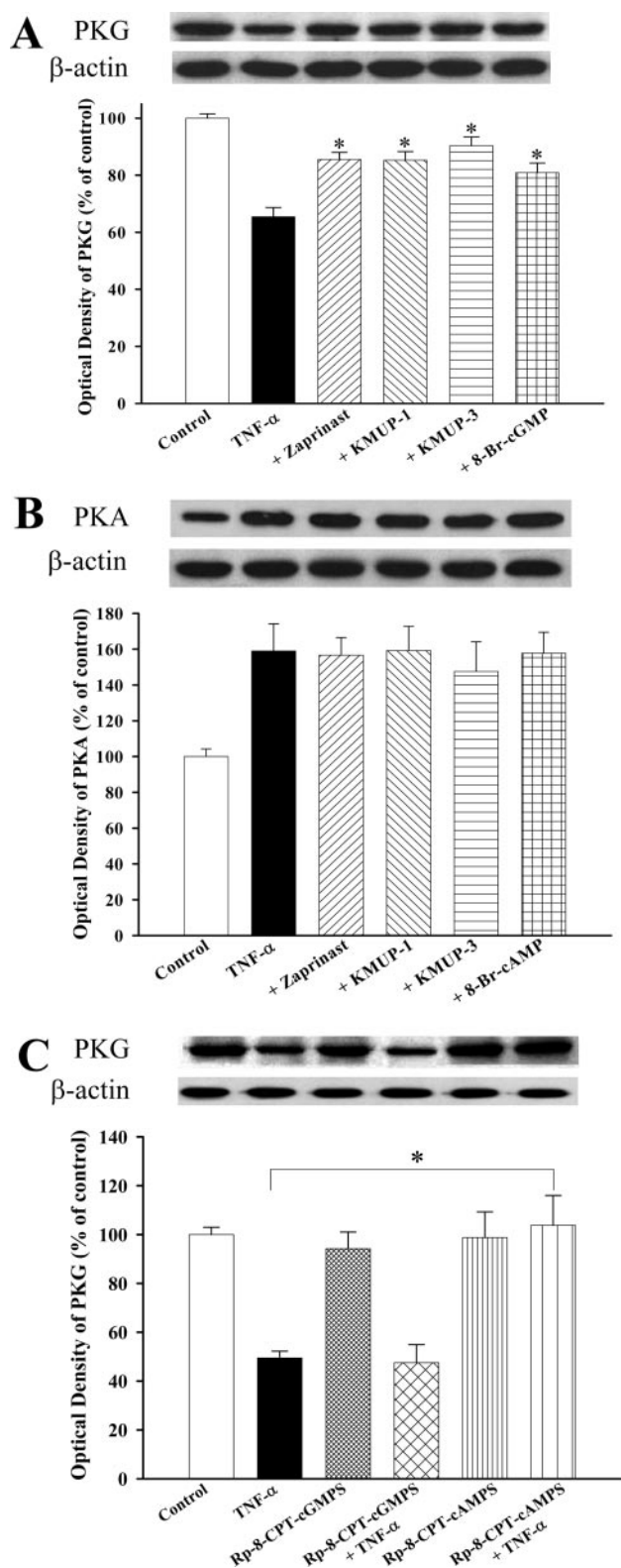


Fig. 8. Effects of KMUP-1, KMUP-3, and various agents on TNF- α -modulated PKG or PKA expression in cultured TSMCs. A, cells pretreated with zaprinast, KMUP-1, KMUP-3, and 8-Br-cGMP (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 9 h. *, $P < 0.05$ versus TNF- α -treated cells. B, cells pretreated with zaprinast, KMUP-1, KMUP-3, and 8-Br-cAMP (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 24 h. C, cells pretreated with Rp-8-CPT-cGMPs and Rp-8-CPT-cAMPS (1 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 9 h. *, $P < 0.05$ versus TNF- α -treated cells. Values are means \pm S.E., $n = 6$.

matory bronchospastic dysfunction. Two xanthine- or imidazole-based derivatives, KMUP-1 and KMUP-3, have been found to increase the expression of cGMP and PKG expression (Wu et al., 2004; Lin et al., 2006), which might make them useful in the treatment of pulmonary airway dysfunctions. In comparison, the more previously discovered indazole derivatives YC-1 and BAY-41-2272 have been shown to activate sGC by a binding manner, leading to increases of cGMP (Ko et al., 1994; Chang et al., 2004; Balashova et al., 2005). The chemical relationship between KMUP-1/KMUP-3 and YC-1/BAY-41-2272 were thus noted for their sGC activation, PKG expression, and cGMP-enhancing activities.

Exogenous TNF- α in vivo has been shown to induce bron-

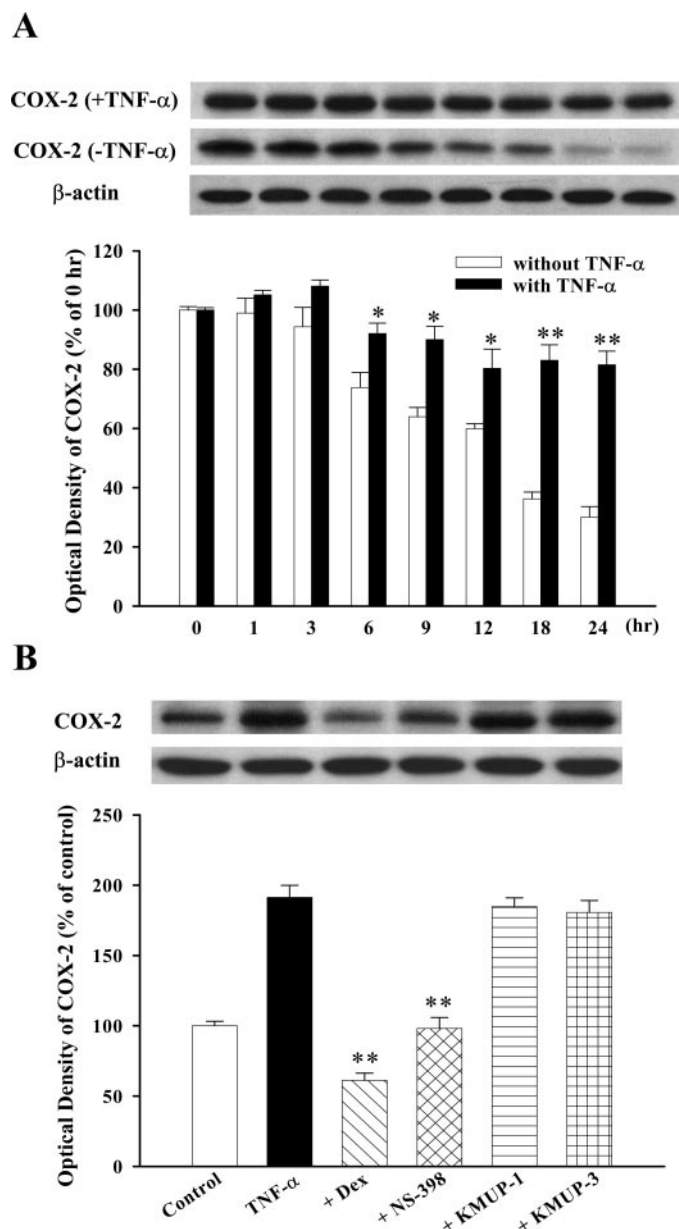


Fig. 9. TNF- α modulated the expression of COX-2 in cultured TSMCs. A, cells treated with TNF- α (100 ng/ml) for 0, 1, 3, 6, 9, 12, 18, and 24 h. *, $P < 0.05$; **, $P < 0.01$ versus respective time without TNF- α . B, cells pretreated with dexamethasone (Dex, 1 μ M), NS-398 (10 μ M), KMUP-1 (10 μ M), and KMUP-3 (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 24 h. Values are means \pm S.E., $n = 6$. **, $P < 0.01$ versus TNF- α -treated cells.

chial hyper-responsiveness (Hakonarson et al., 1996). In this study, TNF- α induced the expression of PKG and cGMP and increased the expression of iNOS and NO. In theory, iNOS would be expected to increase the production of NO and cGMP (Giembycz, 2000; Lincoln et al., 2001). However, cytokines have been found in one study to decrease sGC subunit mRNA and protein levels and reduce NO-stimulated sGC enzyme activity (Takata et al., 2001). This reduced generation of cGMP in cytokine-treated cells could be caused by a decrease in sGC, an increase in cGMP phosphodiesterase activity, or an endogenous reaction between NO and O₂⁻, leading to the formation of ONOO⁻. This oxidation reaction would affect the activity of sGC (Weber et al., 2001; Yan et al., 2003). In the present study, the time of onset and the duration of the induction of iNOS expression by TNF- α were

similar to those of sGC subunit protein levels. Thus, we further investigated the effects of KMUP-1 and KMUP-3 on TNF- α -induced iNOS and associated sGC expression.

Exogenous cytokine, such as interleukin-1 β in pulmonary epithelial cells, has been reported to increase the release of PGE₂ through the activation of COX-2 (Chang et al., 2004). In this study, exogenous TNF- α also caused significant increases in the release of PGE₂ and PGI₂, production of cAMP, and expression of PKA in TSMCs (Figs. 8, 10B, and 12). These results seemed to confirm that TNF- α is able to elicit the PGE₂ formation via the activation of COX-2 in TSMCs. KMUP-1 and KMUP-3 cannot further increase PKA and cAMP in the presence of TNF- α (Figs. 8B and 10B) and implicate that they might predominantly have the ability to protect against TNF- α -induced inhibition of PKG (Fig. 13).

Inflammatory response is under the control of cAMP,

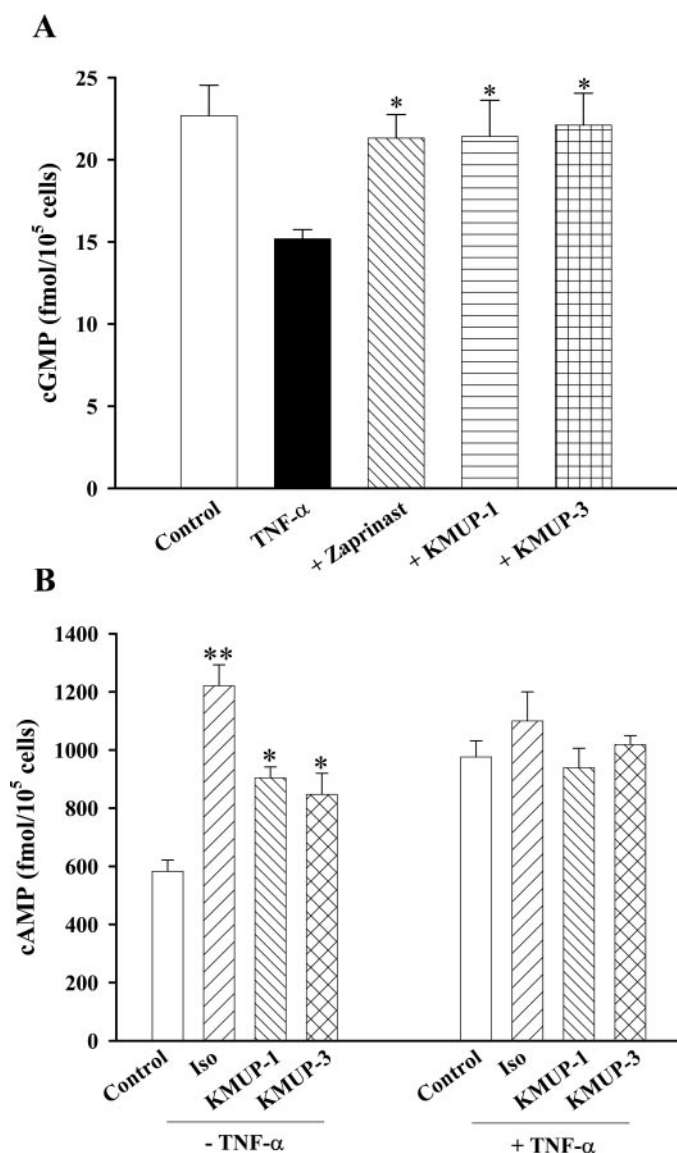


Fig. 10. Effects of KMUP-1, KMUP-3, and various agents on TNF- α -induced cGMP or cAMP production in cultured TSMCs. A, cells pretreated with zaprinast, KMUP-1, and KMUP-3 (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 9 h. *, $P < 0.05$ versus TNF- α -treated cells. B, cells pretreated with isoproterenol (Iso), KMUP-1, and KMUP-3 (10 μ M) for 20 min and further incubated with and without TNF- α (100 ng/ml) for 24 h. Values are means \pm S.E., $n = 3$, each conducted in triplicate. *, $P < 0.05$; **, $P < 0.01$ versus control.

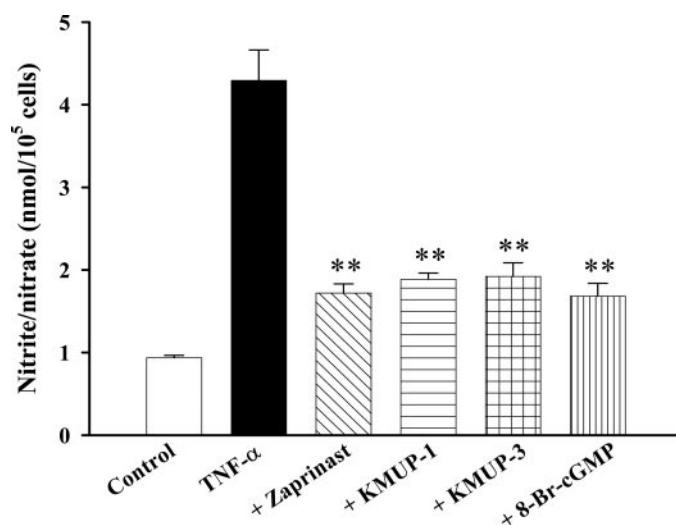


Fig. 11. Effects of KMUP-1, KMUP-3, and various agents on TNF- α -induced nitrite/nitrate production in cultured TSMCs. Cells pretreated with zaprinast, KMUP-1, KMUP-3, and 8-Br-cGMP (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 9 h. Values are means \pm S.E., $n = 3$, each conducted in triplicate. **, $P < 0.01$ versus TNF- α -treated cells.

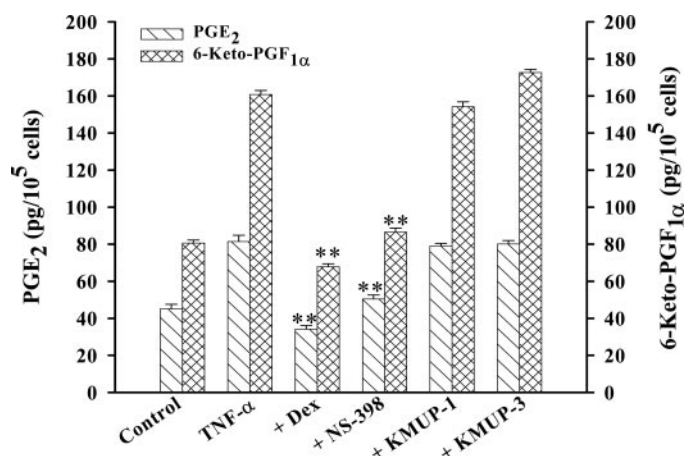


Fig. 12. Effects of KMUP-1, KMUP-3, and various agents on TNF- α -induced PGE₂ or 6-keto-PGF₁ α (PGI₂ stable metabolite) production in cultured TSMCs. Cells pretreated with dexamethasone (Dex, 1 μ M), NS-398 (10 μ M), KMUP-1 (10 μ M), and KMUP-3 (10 μ M) for 20 min and further incubated with TNF- α (100 ng/ml) for 24 h. Values are means \pm S.E., $n = 3$, each conducted in triplicate. **, $P < 0.01$ versus TNF- α -treated cells.

which is partly regulated by PDE. The role of PDE4 inhibition in anti-inflammation and on COX-2 expression has been described in non-TSMCs (Juergens et al., 1999; Jimenez et al., 2004). PDEIs suppressed IL-1-induced NO release and iNOS mRNA expression (Geng et al., 1998; Beshay and Prud'homme, 2001). Pentoxifylline also affected NO synthesis by interferon- γ (Samardzic et al., 2000). TNF- α -induced iNOS expressions were reduced by amrinone in cardiomyocyte, and such effects were increased by milrinone, a PDE3 and PDE4 inhibitor (Chanani et al., 2002). On the other hand, 8-Br-cGMP augmented NO synthesis in IL-1 β -stimulated articular chondrocytes and cardiac myocytes (Geng et al., 1998). The increase of cGMP also augmented IL-1-induced iNOS expression in vascular smooth muscle cells (Inoue et al., 1995). In addition, 8-Br-cGMP inhibited iNOS expression and NO production in LPS-activated macrophages (Pang and Hout, 1997). We observed that 8-Br-cGMP attenuated the expression of iNOS in TSMCs. These evidences suggest that the effects of cGMP on iNOS seem to be cell- and probably stimulus-dependent.

We investigated whether the effects of TNF- α on TSMCs could be prevented by cGMP enhancers. As shown in Fig. 10A, pretreatment with KMUP-1, KMUP-3 and zaprinast all protected against TNF- α -induced decreases of cGMP. We further investigated whether exposure of TSMCs to TNF- α could affect the expression of PKG protein by cGMP enhancers. As expected, incubation of TSMCs with TNF- α decreased the level of PKG, which was reversed by treatment with KMUP-1 and KMUP-3 (Fig. 7A).

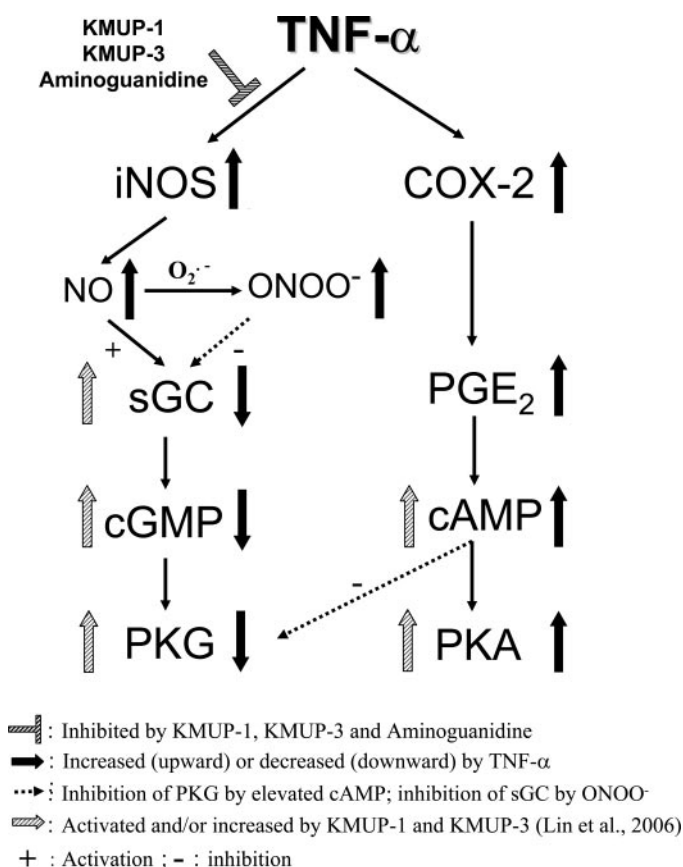


Fig. 13. Proposed mechanisms of action of KMUP-1 and KMUP-3 on TNF- α -induced inflammatory responses in rat tracheal smooth muscle cells.

In certain physiological and pathophysiological circumstances, cAMP may act through PKG, showing their cross-interaction. Although PKG is relatively specific for cGMP over cAMP, the basal cAMP concentration in smooth muscle is typically 5 to 6 times higher than the concentration of cGMP, which allows cross-activation when cAMP is elevated. Conversely, in some instances, PKA may be cross-activated by cGMP (Shabb, 2001). In the presence of TNF- α , the expression of PKG decreased to the minimum at 9 h and then returned to basal level; in contrast, cAMP increased gradually and time-dependently. Thus, we have further confirmed previous findings (Browner et al., 2004) that elevation of cAMP but not cGMP by pro-inflammatory mediators mediates the PKG down-regulation in rat TSMCs (Figs. 8C and 13). Here, KMUP-1 and KMUP-3 were suggested to inhibit iNOS and associated ONOO $^-$ formation, similar to that by aminoguanidine (Ruetten and Thiemeermann, 1996), and to increase cGMP by activating sGC and inhibiting PDE (Lin et al., 2006). It is noteworthy that both elevated cAMP- and TNF- α -induced decreases of PKG could be prevented or reversed by KMUP-1 and KMUP-3 (Fig. 13).

COX-2 is up-regulated by inflammatory stimuli such as cytokines, including TNF- α , and it contributes to the pathogenesis of inflammation (Laporte et al., 1998). Although COX-2 is generally believed to be inducible, there is evidence that COX-2 is expressed constitutively and the expression is low under physiological conditions (Baber et al., 2003). In this study, the expression of COX-2 protein without TNF- α challenge sharply decreased in a time-dependent manner. In contrast, in the presence of TNF- α , the expression of COX-2 was limited to moderate decreases. However, at a respective incubation time of TSMCs, the expression of COX-2 by TNF- α was significantly different from that of nontreatment (Fig. 9A). Accompanied with TNF- α -induced COX-2 expression in TSMCs, we further investigated whether exposure of TSMCs to TNF- α could elicit the production of PGE $_2$ and PGI $_2$. As expected, incubation of TSMCs with TNF- α increased PGE $_2$ and 6-keto-PGF $_{1\alpha}$ (PGI $_2$ stable metabolite). Preincubation of TSMCs with NS-398 (a selective COX-2 inhibitor) caused a marked reduction in TNF- α -induced PGs, the products of activated COX-2. However, KMUP-1 and KMUP-3 had no significant effects on the expression of COX-2 and the production of PGE $_2$ and PGI $_2$ after TNF- α challenge.

In summary, our results indicate that TNF- α significantly increases iNOS expression and NO formation in TSMCs. Thus, overproduction of NO and accompanied oxidative reaction were suggested to decrease sGC expression, cGMP production and PKG protein levels (Weber et al., 2001; Munzel et al., 2003; Yan et al., 2003). TNF- α also induces expression of COX-2 and productions of PGE $_2$ and PGI $_2$, leading to increases in cAMP and PKA. KMUP-1, KMUP-3, and 8-Br-cGMP protect against TNF- α -induced increases in expression of iNOS and PKA and production of NO. They further protect against decreases in expression of sGC and PKG in this study. These findings suggest that they might possess iNOS inhibition-associated anti-proinflammatory properties, unrelated to inhibition on COX-2 expression, in TSMCs. KMUP-1 and KMUP-3 did not induce the iNOS expression activity, which was able to be shown by sildenafil (Salloum et al., 2003). Unlike YC-1 in pulmonary epithelial cells (Chang et al., 2004), KMUP-1 and KMUP-3 were not found to induce

the activation of COX-2 in TSMCs. Similar to that with β_2 -adrenergic agonist, pretreatment of TSMCs with KMUP-1 and KMUP-3 had no significant effects on TNF- α -induced expression of COX-2 and production of PGE₂ and PGI₂ (Shore, 2002). However, elevation of cAMP by PGE₂ can desensitize the airway response to β_2 -adrenergic agonist by inhibiting cGMP and PKG (Laporte et al., 1998). We thus suggest that KMUP-1 and KMUP-3 might reverse this desensitization via enhancing cGMP, sGC activation, and PKG expression.

In conclusion, in the presence of TNF- α in TSM, KMUP-1 and KMUP-3 might modulate the cross-action between PKA and PKG by activating sGC/cGMP/PKG pathway without the involvement of COX-2. KMUP-1 and KMUP-3 might reduce cytokine-induced proinflammation and limit the risk of further worsening of pulmonary dysfunction.

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