**Pistacia weinmannifolia** ameliorates cigarette smoke and lipopolysaccharide-induced pulmonary inflammation by inhibiting interleukin-8 production and NF-κB activation

JAE-WON LEE1*, HYUNG WON RYU1*, SU UI LEE1, MIN-GU KIM1, OK-KYOUNG KWON1, MUN OK KIM1, TAE KYU OH2, JAE KYOUNG LEE2, TAE YOUNG KIM2, SANG WOO LEE3, SANGHO CHOI1, WAN-YI LI4, KYUNG-SEOP AHN1 and SEI-RYANG OH1

1Natural Medicine Research Center, Korea Research Institute of Bioscience and Biotechnology (KRIIBB), Cheongju, Chungcheongbuk-do 28116; 2BTC Corporation, Technology Development Center, Ansan, Gyeonggi-do 15588; 3International Biological Material Research Center, Korea Research Institute of Bioscience and Biotechnology (KRIIBB), Daejeon 34141, Republic of Korea; 4Institute of Medicinal Plants, Yunnan Academy of Agricultural Sciences, Kunming, Yunnan 650200, P.R. China

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**Abstract.** *Pistacia weinmannifolia* (PW) has been used in traditional Chinese medicine to treat headaches, dysentery, enteritis and influenza. However, PW has not been known for treating respiratory inflammatory diseases, including chronic obstructive pulmonary disease (COPD). The present *in vitro* analysis confirmed that PW root extract (PWRE) exerts anti-inflammatory effects in phorbol myristate acetate- or tumor necrosis factor α (TNF-α)-stimulated human lung epithelial NCI-H292 cells by attenuating the expression of interleukin (IL)-8, IL-6 and Mucin A5 (MUC5AC), which are closely associated with the pulmonary inflammatory response in the pathogenesis of COPD. Thus, the aim of the present study was to evaluate the protective effect of PWRE on pulmonary inflammation induced by cigarette smoke (CS) and lipopolysaccharide (LPS). Treatment with PWRE significantly reduced the quantity of neutrophils and the levels of inflammatory molecules and toxic molecules, including tumor TNF-α, IL-6, IL-8, monocyte chemoattractant protein-1, neutrophil elastase and reactive oxygen species, in the bronchoalveolar lavage fluid of mice with CS- and LPS-induced pulmonary inflammation. PWRE also attenuated the influx of inflammatory cells in the lung tissues. Furthermore, PWRE downregulated the activation of nuclear factor-κB and the expression of phosphodiesterase 4 in the lung tissues. Therefore, these findings suggest that PWRE may be a valuable adjuvant treatment for COPD.

**Introduction**

Chronic obstructive pulmonary disease (COPD) is a major health problem, and its incidence has continued to increase globally in the past decade (1,2). Airway inflammation, one of the main forms of COPD, is caused by cigarette smoking and its pathogenesis is closely associated with increased levels of inflammatory molecules (3). Bacterial respiratory tract infections are associated with the initial occurrence and exacerbation of the airway inflammatory response in COPD (4). The increased concentration of neutrophils is a unique characteristic and clinical indicator of COPD, and cell-derived toxic molecules, including reactive oxygen species (ROS) and neutrophil elastase (NE), cause oxidative stress and emphysema (5,6). The release of interleukin (IL)-8 by bronchial epithelial cells leads to an increase in neutrophil and macrophage chemotaxis (7). Macrophage activation has been reported to cause the airway inflammatory response via the release of inflammatory cytokines and chemokines, including IL-6, tumor necrosis factor-α (TNF-α) and monocyte chemoattractant protein-1 (MCP-1) (8,9). It is well understood that the activation of nuclear factor-κB (NF-κB) leads to the release of inflammatory cytokines and chemokines, including IL-6, IL-8, TNF-α and MCP-1 (10-12). Natural products and their derivatives have been used for the treatment of various diseases, including infectious diseases, and have been reported to serve an important role in the prevention of inflammatory diseases (13-15). It is well known that Traditional Chinese Medicine exerts valuable therapeutic effects on pulmonary diseases (16). *Pistacia weinmannifolia*
(PW) is a plant predominantly distributed in the Yunnan province of China and is used as a herbal remedy for treating dysentery, enteritis, influenza and lung cancer (17,18). The major metabolites of PW have been reported to possess a wide variety of biological activities, including antioxidant effects and histamine-release inhibition (19). However, to the best of our knowledge, its protective effect against cigarette smoke (CS)- and lipopolysaccharides (LPS)-induced pulmonary inflammation remains to be elucidated in a mouse model. Based on results of previous studies (17-19) and an observation from the present in vitro study, which demonstrated the anti-inflammatory effects of PW root extract (PWRE), the protective effect of PWRE against CS- and LPS-induced pulmonary inflammation was examined.

Materials and methods

Preparation of PWRE. PW roots were collected in the Yunnan province of China in August 2017 and identified by Dr. Sangwoo Lee (International Biological Material Research Center, Korea Research Institute of Bioscience and Biotechnology, South Korea). A voucher specimen (D180305001) of this raw material was deposited at the International Biological Material Research Center (http://www.ibmrc.re.kr/) of the Korea Research Institute of Bioscience and Biotechnology (Chungbuk, Korea). The drug substance from the PW roots was produced by a processing method described in the International Conference on Harmonisation and Ministry of Food and Drug Safety guidelines (http://www.mfds.go.kr/eng/wppe/m_17/de0110081001.do). The collected roots were dried immediately after sampling and then ground to a powder and stored at a temperature lower than -20°C prior to extraction. The raw materials were then packed in laminated bags, delivered to Korea and stored at -25°C for 3 months until further analysis and biological testing. The extracts of PW roots were provided by the BTC Corporation (Korea Good Manufacturing Practice; cat. no. BTC-PWE-180118). The powdered samples were extracted twice with 50% ethanol at 80°C and concentrated in vacuo at 60°C. The product was dried in a freeze dryer to produce dried extracts (~19% yield).

Ultra-high performance liquid chromatography-quadrupole time-of-flight mass spectrometry (UPLC-QToF-MS). UPLC-QToF-MS (Micromass Q-Tof Premier™, Waters Corporation, Milford, MA, USA) analyses were performed with an ACQUITY BEH C18 chromatography column (2.1x100 mm; 1.7 µm; Waters Corporation) with a linear gradient (0 min, 1% B; 0-1.0 min, 1-5% B; 1.0-10.0 min, 5-30% B; 10.0-17.0 min, 30-60% B; 17.0-17.1 min, 60-100% B; 17.1-19.0 min, 100% B; 19.0-20.0 min, back to 1% B) of acetonitrile/water (HPLC-grade; Merck KGaA). The injection volume was 2 µl. The QToF-MS was operated in negative ion mode with the following conditions: Source temperature, 110°C; desolvation temperature, 350°C; capillary voltage, 2,300 V; and cone voltage, 50 V. Leucine-enkephalin (400 pg/ml) was used as the reference compound (m/z 554.2615 in the negative mode) at a flow rate of 0.4 ml/min. Accurate mass, MS/MS and elemental composition were calculated using the MassLynx software (Waters Corporation) incorporated into the instrument.

Cell preparation and culture. Human lung epithelial NCI-H292 cells were purchased from the American Type Culture Collection (catalog no. CRL-1848; Manassas, VA, USA). Cells were cultured in RPMI-1640 medium (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS; HyClone; GE Healthcare, Chicago, IL, USA), and 100 U/ml penicillin and 100 µg/ml streptomycin (HyClone; GE Healthcare) at 37°C under a humidified 5% CO₂ atmosphere.

For the enzyme-linked immunosorbent assay (ELISA) to assess IL-8, IL-6 and MUC5AC production, NCI-H292 cells were seeded in 24-well plates at a density of 1x10⁵ cells for 16 h. The cells were then transferred to reduced-serum medium (0.1% FBS). After a 16-h incubation period, the cells were treated with PWRE for 2 h prior to adding inflammatory stimuli, including 100 nM phorbol myristate acetate (PMA; Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) or 20 ng/ml TNF-α (PeproTech, Inc., Rocky Hill, NJ, USA) based on the previous reports (20-23). After the addition of the stimuli, the cells were further incubated for 12 h at 37°C under a humidified 5% CO₂ atmosphere.

Cell viability assay. NCI-H292 cells were seeded and cultured in 96-well plates in RPMI-1640 medium at a density of 5x10⁴ cells/well for 16 h. The medium was subsequently replaced with a reduced-serum medium (0.1% FBS). After a 16-h incubation, the cells were cultured with the corresponding concentrations (1, 2.5, 5 and 10 µg/ml) of PWRE in the presence or absence of inflammatory stimuli, including PMA (100 nM) or TNF-α (20 ng/ml) for 24 h at 37°C under a humidified 5% CO₂ atmosphere. Cell growth was measured in triplicate using a Cell Counting Kit-8 (Dojindo Molecular Technologies, Inc., Rockville, MD, USA), according to the manufacturer’s protocol. The absorbance at 450 nm was measured with a BioTek microplate reader and calculated as a percentage of the control value for each condition.

Mouse model of pulmonary inflammation induced by CS and LPS. In mice, the inhalation of CS and the administration of LPS accelerates the airway inflammatory response similar to what is seen in patients with COPD (2,3,24). Therefore, CS exposure and LPS administration were achieved using a modification of the methods described by Lee et al (25). Briefly, six-week-old C57BL/6N male mice (n=24; body weight, 18-20 g) were obtained from Koatech Co. Ltd. (Pyeongtaek-si, Korea), quarantined and acclimated to a specific pathogen-free atmosphere. Cell growth was measured in triplicate using a Cell Counting Kit-8 (Dojindo Molecular Technologies, Inc., Rockville, MD, USA), according to the manufacturer’s protocol. The absorbance at 450 nm was measured with a BioTek microplate reader and calculated as a percentage of the control value for each condition.
(ROF) were orally administered for six consecutive days prior to CS exposure. LPS (5 µg dissolved in 30 µl distilled water) was intranasally instilled on day 5. The experimental procedures of this study were performed in accordance with the procedures approved by the Institutional Animal Care and Use Committee of the Korea Research Institute of Bioscience and Biotechnology (Chungbuk, Korea; KRIBB-AEC-18054) and in compliance with the National Institutes of Health guidelines for the Care and Use of Laboratory Animals and Korean national laws for animal welfare.

Measurement of inflammatory cell count. To distinguish the different cells, bronchoalveolar lavage fluid (BALF) was collected as described previously (26). Briefly, mice were anesthetized with Zoletil 50® (30-50 mg/kg IP; Virbac, Co., Seoul, Korea) and Xylazine (5-10 mg/kg IP; Bayer Korea, Seoul, Korea) 24 h after the intranasal administration of LPS. The anesthetic conditions were based on previous studies (27,28), which provide information on the usefulness of the combined use of Zoletil with Xylazine. This combined administration adequately maintained sedation and analgesia. Toxic signs, including dyspnea and vomiting, were not observed in this condition. BALF was collected by tracheal cannulation and infusion with 0.7 ml ice-cold PBS twice (total volume, 1.4 ml). To count the inflammatory cells, 0.1 ml BALF was centrifuged onto a slide using a Cytospin (Hanil Science Industrial, Seoul, Korea) at 246 x g for 5 min at room temperature, and the slides were dried and stained using the Diff-Quik® staining reagent (cat. no. 38721; Sysmex Corporation, Ltd., Japan), according to the manufacturer's protocol. Finally, inflammatory cells were counted under a light microscope (magnification, x400), with ≥4 fields counter per BALF sample.

Determination of inflammatory cytokines and toxic molecules. The levels of IL-6, IL-8, TNF-α and MCP-1 in the BALF were measured using ELISA kits, according to the manufacturers' protocols. The IL-6 (cat. no. 555240) and TNF-α (cat. no. 558534) ELISA kits were purchased from BD Biosciences, the IL-8 kit (cat. no. MBS261967) was from MyBioSource, Inc. (San Diego, CA, USA), and the MCP-1 kit (cat. no. DY479) was obtained from (R&D Systems, Inc., Minneapolis, MN, USA). To evaluate the levels of IL-6, IL-8 and MUC5AC in the human NCI-H292 cells, the presence of cytokines in the supernatant was analyzed using human IL-6 and IL-8 ELISA kits (BD Biosciences, Franklin Lakes, NJ, USA), according to the manufacturer's protocols. Human MUC5AC protein levels were measured by modified ELISA assay using anti-MUCSAC (Abcam, Cambridge, MA, USA), as previously described (29). The absorbance was measured at 450 nm using a Spark™ 10 M multimode microplate reader (Tecan Science and Technology Co., Ltd., Shanghai, China). To measure the intracellular ROS activity, the BALF cells were incubated with 20 µM 2',7'-dichlorofluorescein diacetate (Sigma-Aldrich; Merck KGaA) for 10 min. at 37°C. The results were observed using a fluorescence plate reader (488 nm excitation and 525 nm emission). The level of NE activity was monitored using N-succinyl-(Ala)3-p-nitroanilide (Sigma-Aldrich; Merck KGaA) at 37°C for 90 min. The absorbance was read at 405 nm using an Epoch Microplate Spectrophotometer (BioTek Instruments, Inc., Winooski, VT, USA) and used to calculate the percentage of the control for each condition.

Western blot analysis. The lung tissues were homogenized in lysis buffer (cat. no. c3228; Sigma-Aldrich; Merck KGaA) containing phosphatase and protease inhibitors (cat. nos. 04906837001 and 11836153001, respectively; Roche Diagnostics, Basel, Switzerland). The Pierce BCA Protein assay kit (cat. no. 23225; Thermo Fisher Scientific, Inc.) was used to determine the protein concentration. Equal quantities of proteins (50 µg/lane) were separated by 10-15% SDS-PAGE and the proteins were then transferred onto PVDF membranes (EMD Millipore, Billerica, MA, USA). The membranes were blocked in 5% skim milk for 1 h and probed overnight at 4°C with primary antibodies. The primary antibodies and dilutions were as follows: Anti-phosphorylated (p)-NF-κB p65 (cat. no. 3033; 1:1,000; Cell Signaling Technology, Inc.,Danvers, MA, USA), anti-IκB-α (cat. no. 2859; 1:1,000; Cell Signaling Technology, Inc.), anti-β-actin (cat. no. 4967; 1:1,000; Cell Signaling Technology, Inc.), anti-NF-κB p65 (cat. no. sc-372; 1:1,000; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), anti-p-IκB-α (cat. no. MA5-15087; 1:1,000; Invitrogen; Thermo Fisher Scientific, Inc.) and anti-PDE4B (cat. no. ab170939; 1:1,000; Abcam). The membranes were washed with TBS and 0.05% Tween-20 (TBST), and incubated with HRP-conjugated secondary antibodies (cat. no. 111-035-003; 1:2,000; Jackson ImmunoResearch Laboratories, Inc., West Grove, PA, USA) at room temperature for 2 h. Finally, the membranes were washed with TBST and developed with an enhanced chemiluminescence kit (Thermo Fisher Scientific, Inc.). All bands were visualized by the LAS-4000 luminescent image analyzer (Fujifilm, Tokyo, Japan) and quantified by densitometry using Fuji Multi Gauge software version 3.0 (Fuji photo film Co, Ltd. Japan).

Histological analysis. The lung tissues were removed from the mice 24 h after the last oral administration of PWRE and fixed in 10% (v/v) neutral buffered formalin solution (BBC Biochemical, Mount Vernon, WA, USA) at room temperature for 48 h. For histological evaluation, the lung tissues were embedded in paraffin and sectioned at a thickness of 4 µm using a rotary microtome. Subsequently, the lung sections were stained with a hematoxylin (cat. no. 3580; BBC Biochemical Inc.) and eosin (cat. no. 6766007; Thermo Fisher Scientific Inc.) solutions at room temperature for 30 sec each. The sections were visualized using a light microscope (magnification, x100) to estimate the influx of inflammatory cells. The degree of inflammatory cell influx in each group was examined by three independent observers using the following semi-quantitative scope: 0, no influx; 1, low influx; 2, moderate influx; 3, large influx; and 4, severe influx.

Statistical analysis. Data are presented as the mean ± standard deviation. Statistical significance was analyzed by a two-tailed Student's t-test for comparisons between two groups. One-way ANOVA followed by Dunnett's multiple comparison test was used to analyze differences between multiple groups. Data were analyzed using SPSS 20.0 (IBM Corp., Armonk, NY, USA). P<0.05 was considered to indicate a statistically significant difference.
Results

Extract of PWRE. The high potency of the *P. weinmannifolia* roots indicated the presence of compounds that may be therapeutic against COPD. Therefore, the major compounds of the extracts were expected to exhibit inhibitory activities. Following the analysis, the major compounds were characterized using mass data. As presented in Fig. 1, it was revealed that the deprotonated molecule [M-H]⁻ was the most abundant for two compounds: m/z 197 for ethyl gallate (C₉H₁₀O₅) and m/z 939 for penta-O-galloyl-glucose (C₄₁H₃₂O₂₆).

Effects of PWRE on PMA- or TNF-α-stimulated NCI-H292 cells. PMA and TNF-α are potent pro-inflammatory stimuli that are used extensively to induce secretagogue effects on the human respiratory epithelium, and they are a good in vitro model for evaluating the inflammatory response (30). NCI-H292 cells, a human pulmonary mucoepidermoid carcinoma cell line, are often used for the purpose of elucidating the inflammatory response involved in airway inflammatory gene expression and mucin secretion (31). Thus, the present study investigated whether PWRE suppresses the inflammatory response in PMA- or TNF-α-stimulated human lung epithelial NCI-H292 cells. Prior to this investigation, the current study evaluated the cytotoxicity of PWRE in the presence or absence of PMA or TNF-α in NCI-H292 cells and identified that PWRE demonstrates no cytotoxicity at any of the concentrations used (Fig. 2A and E). A concentration <10 µg/ml was used in the subsequent experiments. As presented in Fig. 2B-D, PMA significantly enhanced IL-8, IL-6 and MUC5AC secretion in NCI-H292 cells. This increase in IL-8, IL-6 and MUC5AC secretion was significantly suppressed by treatment with PWRE. Furthermore, PWRE suppressed the expression of TNF-α-induced IL-8 and IL-6 in NCI-H292 cells (Fig. 2F and G). These results indicate that PWRE effectively inhibited the PMA- or TNF-α-induced inflammatory response in NCI-H292 cells.

Effects of PWRE on the quantity of inflammatory cells in the BALF of the CS- and LPS-induced pulmonary inflammation animal model. Pulmonary inflammation was achieved by CS exposure and LPS administration (Fig. 3). A significant increase in inflammatory cells, including neutrophils and macrophages, was detected in the BALF of the CS+LPS group compared with the NC group (Fig. 4A and B). However, the level of neutrophils was significantly reduced in the PWRE-treated mice compared with the CS+LPS group (*P*<0.01). The level of macrophages was also decreased to the level of the ROF in the
PWRE-treated mice (P<0.01). In total, the inhibitory effects of 15 mg/kg PWRE were similar to those of 5 mg/kg ROF, which was used as a positive control.

**Effects of PWRE on inflammatory cytokines and chemokines in the BALF of the CS- and LPS-induced pulmonary inflammation animal model.** The concentration of inflammatory molecules in the BALF was measured by ELISA. As presented in Fig. 5, the levels of TNF-\(\alpha\), IL-6, IL-8 and MCP-1 were significantly higher in the CS+LPS group compared with the NC group, whereas a significant reduction of these molecules was detected in the PWRE-treated group (Fig. 5A-D). Compared with the CS+LPS group, the inhibition levels of ROF against the inflammatory molecules were 53.50 (TNF-\(\alpha\)), 68.98 (IL-6), 49.46 (IL-8) and 53.61% (MCP-1; P<0.01), while for PWRE, they were 62.21 (TNF-\(\alpha\)), 72.09 (IL-6), 49.46 (IL-8) and 75.53% (MCP-1; P<0.05).
Effects of PWRE on the levels of toxic molecules in the BALF of the CS- and LPS-induced pulmonary inflammation animal model. The present results demonstrated the inhibitory effect of PWRE on neutrophil influx (Fig. 4A). Based on this result, the inhibitory activity of PWRE on the neutrophil-derived toxic molecules NE and ROS was evaluated. As presented in Fig. 6A, a significant increase in NE activity was observed in the CS+LPS group compared with the NC group. However, PWRE-treatment led to a significant reduction in this activity down to the same level.
as that observed following ROF-treatment. Similarly, the increased production of ROS was also significantly reversed by PWRE-treatment (Fig. 6B).

**Effects of PWRE on inflammatory cell influx in the lungs of the CS- and LPS-induced pulmonary animal model.** As the present study confirmed the regulatory effects of PWRE on neutrophil recruitment and MCP-1 production (Figs. 4A and 5D), it was next examined whether the inflammatory cell recruitment into the lungs by exposure to CS and LPS is attenuated by PWRE. Hematoxylin and eosin staining was used to examine histological changes in the lungs. As presented in Fig. 7, in the lungs of the CS+LPS group, the degree of inflammatory cell influx was significantly higher compared with that of the NC group. Notably, the degree of influx was significantly reduced in the lungs of the PWRE group.

**Effects of PWRE on the activation of NF-κB in the lungs of the CS- and LPS-induced pulmonary inflammation animal model.** Under inflammatory conditions, the activation of NF-κB leads to the release of inflammatory molecules, including IL-6, IL-8 and MCP-1 (32-34). The present study confirmed the effect of PWRE on the production of these molecules (Fig. 5). Therefore, it was next evaluated whether PWRE regulates the activation of NF-κB using western blot analysis. As presented...
in Fig. 8, the level of NF-κB p65 phosphorylation was significantly upregulated in the lung tissues of the CS+LPS group compared with the NC group. However, PWRE-treatment significantly reduced this level in the PWRE group compared with the CS+LPS group. To further clarify the relevance of the NF-κB signaling pathway, the current study next focused on the molecular aspects, including IκB-α, which is a key signaling molecule in the NF-κB pathway (3,26). It is well established that IκB activation and degradation lead to NF-κB activation (35,36). As presented in Fig. 8, the level of IκB-α phosphorylation was significantly upregulated in the lung tissues of the CS+LPS group compared with the NC group, whereas this level was effectively inhibited in the lungs of the PWRE group.

Effects of PWRE on PDE4 expression in the lungs of the CS- and LPS-induced pulmonary inflammation animal model. It is well known that PDE4 serves an important role in inflammation (37) and that the PDE4 inhibitor ROF ameliorates airway inflammation induced by CS and LPS (2,3). The appropriated concentration of ROF was determined with reference to the previous studies (38-42). The present study confirmed that PWRE exerts an anti-inflammatory effect by suppressing the influx of inflammatory cells and the production of inflammatory molecules (Figs. 4-7). These effects of PWRE were similar to those of ROF. Based on these results, the current study further investigated the regulatory effect of PWRE on CS- and LPS-induced PDE4 expression. As presented in Fig. 9, the expression of PDE4B was significantly increased in the lung tissues of the CS+LPS group compared with those of the NC group. However, this increase was significantly reduced in the PWRE group compared to the CS+LPS group. Notably, the inhibitory effect of PWRE on PDE4 expression was similar to those in the ROF group. This result was similar to those of a previous study (37).

Discussion

The results of the present in vitro study demonstrated that PWRE inhibits the release of IL-6, IL-8 and MUC5AC, which are important parameters in COPD pathology. In the in vivo study, PWRE inhibited not only inflammatory cell recruitment but also inflammatory molecules. In addition, the levels of toxic molecules were downregulated with PWRE-treatment. These inhibitory activities were accompanied by NF-κB inactivation.

Airway epithelial cell-derived inflammatory molecules, including IL-6, IL-8 and MUC5AC, have been associated with the pathogenesis of COPD (43). IL-6 is an inflammatory cytokine that is produced by airway epithelial cells and...
macrophages, and its increased level is closely associated with infection and the inflammatory response (10,44,45). IL-8 is recognized as an important neutrophil chemoattractant and MUC5AC inducer, and its level is notably upregulated in bronchial epithelial cells from patients with COPD (46). The increased levels of MUC5AC in respiratory epithelial cells lead to lung function decline (10,47). This suggests that controlling IL-6, IL-8 and MUC5AC levels may be an important element in the prevention and treatment of COPD. Therefore, the present study evaluated the inhibitory effect of PWRE on these molecules in activated H292 airway epithelial cells. It was confirmed that PWRE could control the levels of IL-6, IL-8, MCP-1 and MUC5AC in PMA- or TNF-α-stimulated airway epithelial cells (Fig. 2). Therefore, the protective effect of PWRE on CS- and LPS-induced pulmonary inflammation was subsequently investigated.

The persistent pulmonary inflammatory response is a prominent feature of COPD (3). Macrophages and neutrophils have a central role in lung inflammation and destruction (48,49). Macrophages are the initiators of the inflammatory response, and these cell-derived inflammatory cytokines and chemokines promote a hyper-inflammatory immune response (2,50). A previous study reported that the levels of TNF-α were higher in macrophages in patients with COPD compared with nonsmokers without COPD (4). It is understood that COPD is associated with high levels of IL-6 (51), and the influx of macrophages and the levels of IL-6 are upregulated in the lungs of COPD animal models (52,53). In response to CS exposure, macrophage-derived IL-8 and MCP-1 cause neutrophil recruitment (34,54,55). The high level of neutrophil influx is understood to be a typical characteristic of COPD (56). Neutrophils release a protease that causes damage to the endothelial cells of the airway and accelerates lung destruction (57). Neutrophil-derived ROS lead to oxidative stress in the pathogenesis of lung inflammation and are consistently observed in patients with COPD and COPD animal models (58). Therefore, the regulation of macrophage/neutrophil influx and these cell-derived molecules could be a valuable therapeutic approach in COPD. In the present study, it was identified that PWRE reduces not only the number of macrophages but also their cell-derived inflammatory cytokines and chemokines (Figs. 4B and 5).

The current results also demonstrated that PWRE inhibits neutrophil recruitment and neutrophil-derived toxic molecules (Figs. 4A and 6). In the histological analysis, PWRE regulated the influx of inflammatory cells into the lungs (Fig. 7). Based on the presents results, it can be suggested that the inhibition of inflammatory cell recruitment and inflammatory molecules, including IL-6, IL-8 and MCP-1, by PWRE could eventually lead to the amelioration of the pulmonary inflammatory response. These effects of PWRE were similar to those of ROF and therefore are valuable.

NF-κB activation is closely associated with pulmonary inflammation, and this association has been demonstrated in the lung tissues of COPD animal models (59). In recent studies, approaches to reduce the activation of NF-κB have been attempted in in vivo studies, and improvement of pulmonary inflammation was confirmed with NF-κB inactivation (60,61). Therefore, the current study evaluated the regulatory effect of PWRE on NF-κB activation in mouse models with CS- and LPS-induced pulmonary inflammation. As presented in Fig. 8, PWRE regulated the levels of IκB phosphorylation, IκB degradation and NF-κB phosphorylation. Therefore, it is expected that PWRE exerts protective effects against the inflammatory response in pulmonary inflammatory disease, including COPD.

PDE4 is also closely associated with the inflammatory response (37). In the present study, PWRE ameliorated pulmonary inflammation, and notably, the inhibitory effects of PDE4 expression by PWRE were similar to those of ROF, the positive control. This result is valuable, and therefore, PWRE is expected to exhibit a protective effect against the inflammatory response in pulmonary inflammatory diseases, including COPD.

In conclusion, to the best of our knowledge, for the first time, the present study confirmed that PWRE effectively inhibits inflammatory cell recruitment, toxic molecules and inflammatory mediators. These effects were in accordance with the inhibition of NF-κB. Overall, the current results demonstrated that PWRE exerts a protective role in pulmonary inflammatory response. These findings suggest that PWRE is a potential anti-inflammatory adjuvant treatment in inflammatory diseases, including COPD.

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Availability of data and materials

All data generated and/or analyzed during the present study are included in this published article.

Authors' contributions

JWL performed the in vivo experiments and wrote the manuscript. HWR contributed to the extraction of P. weinmannifolia roots and contributed to the interpretation of the results. SUL performed the in vitro experiments and contributed to the interpretation of the results. TKO, JKL and TYK contributed to the acquisition of data. MGK, OKK, MOK, SWL, SC, WYL and KSA made substantial contributions to the conception and design of the study, acquisition of data, and the analysis and interpretation of data. SRO designed the study and was involved in revising it critically for important intellectual content. All authors discussed the results and approved the final version of the manuscript.

Ethics approval and consent to participate

All experiments were approved by the Institutional Animal Care and Use Committee of the Korea Research Institute of Bioscience and Biotechnology (Cheongju-si, Korea).
Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interest.

References


