EFFECTS OF QUERCETIN ON CYTOKINES AND PULMONARY CAPILLARY CELLS IN PULMONARY ARTERIAL HYPERTENSION RATS

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Abstract

This study aimed to investigate the effects of quercetin in the treatment of pulmonary arterial hypertension (PAH) in a murine model. Thirty-six adult male rats were randomly divided into three groups: the control group (saline), monocrotaline (MCT) - induced PAH group (MCT group) and quercetin treatment group (prevention group). After modelling, the animals from prevention group received 100 mg/kg bw/day quercetin by gavage and the gavage for 20 days, while the animals from the other two groups received the same amount of 0.9% sodium chloride saline solution. The mean pulmonary artery pressure, right ventricular index and relative expression levels of HIF-1 (hypoxia-inducible factor 1), ET-1 (vascular endothelin-1), TGF-β1 (transforming growth factor-β1), VEGF (vascular endothelial growth factor), IL-1 (interleukin-1), IL-6 (interleukin-6) and TNF-α (tumour necrosis factor α) in lung tissues significantly increased in MCT group compared with the control group, 21 days after modelling. The levels of HGF (hepatocyte growth factor) and NAC (N-acetyl-L-cysteine) significantly increased compared with the control group. The treatment with quercetin significantly decreased the level of mean PAH, right ventricular index and relative expression levels of HIF-1, ET-1, TGF-β1, VEGF, IL-1, IL-6 and TNF-α in lung tissues compared with MCT group and significantly decreased the levels of HGF and NAC. In vitro experiment with PCEC (pulmonary capillary endothelial cells) from the three groups showed that in the MCT group the cell proliferation was significantly decreased and the apoptosis was significantly increased compared with the control group, while the quercetin treatment inhibited the MCT-induced cell apoptosis and promoted cell proliferation. In conclusion, quercetin can alleviate PAH by regulating the inflammatory cytokines, promoting cell proliferation and inhibition of cell apoptosis.

Keywords: quercetin, pulmonary arterial hypertension, pulmonary capillary endothelial cells, cytokines

Introduction

Pulmonary arterial hypertension (PAH) is a common and life-threatening disease. It is characterized by continuous and significant increase of pulmonary artery pressure, which eventually leads to right ventricular failure and death [1, 2]. The pathological mechanism of PAH development is complex and multifaceted, and its main feature is the continuous increase of distal pulmonary artery resistance. The common factors of increased resistance include pulmonary vasoconstriction, in situ thrombosis and pulmonary vascular reconstruction. Pulmonary vasoconstriction can lead to alveolar hypoxia and imbalance of vasoactive mediators such as chlorine monoxide and serotonin secretion. Long term contraction can promote in situ thrombosis and lead to vascular reconstruction. PAH is a key pathological link of many common diseases, such as chronic obstructive
Quercetin is a natural dietary flavonoid, spread in nature with low toxicity [6]. In the past 20 years, it has attracted researchers’ attention due to its wide range of pharmacological effects, including antioxidant, anti-inflammatory and anti-cancer [7-10]. For a variety of 33 tumour cell lines, quercetin can inhibit the proliferation of various types of tumour cells, induce their apoptosis, promote their cell cycle arrest, reduce tumour cell migration, and play an anti-tumour role [11]. Besides, some recent studies have shown that quercetin can effectively inhibit coronary and pulmonary vasoconstriction and reduce blood pressure [12]. It was also found that in the PAH rat model induced by MCT, the formation of PAH can be significantly reversed by quercetin [13], however, the key molecular regulatory mechanism is still unclear.

In this research, we investigated if quercetin can effectively reverse PAH using a murine model, the cytokines’ expression and the stimulation of pulmonary capillary endothelial cells (PCEC) proliferation.

Materials and Methods

**PAH animal model and experimental groups**

\[
\text{weight ratio} = \frac{\text{weight of right ventricles}}{\text{Weight of left ventricle and interventricular septum}}
\]

**PCEC (pulmonary capillary endothelial cells) collection and primary culture**

The fresh lung tissue was collected from the rats from each group, digested in 0.25% pancreatin (ChemeGen, USA) at 37°C for 5 minutes, and then the tissue was carefully cut in 2 - 3 mm segments from the edge of the lung leaf and cut it into 1 mm³ small pieces. Then the tissue was dissolved in 20 mL endothelial cell culture medium (ECM1001, ScienCell, USA), and 6 mL 0.25% collagenase IV (Roche, Shanghai, China) was added for digestion in 37°C incubator for 15 min. Then 6 mL of 1.0% neutral protease solution (Roche, Shanghai, China) was added and the digestion was continued for 15 min at 37°C in an incubator for 15 min. The complete digestion of cells was observed under the microscope. When the digestion was stopped, another 10 mL ECM medium was added and centrifuged at 4000 rpm for 2 - 5 minutes. After centrifugation, the supernatant was discarded and 3 mL serum (Thermo Fisher, USA) was added to induce cell aggregation and centrifuged at 100 rpm for 6 minutes. The supernatant was discarded and the precipitation was collected and mixed with 20 mL ECM and inoculate into culture bottle. After incubation at 37°C in a 5% CO₂ incubator for 2 - 4 hours, the cells and fragments that were not attached to the wall were removed, and the new ECM was added to continue the cell culture.

**MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) assay**

The PCECs were collected, adjusted to the cell concentration of 3 × 10^4 cells/mL, inoculated in a 96-well plate at a concentration of 100 µL/well and cultured at 37°C in a 5% CO₂ incubator for 24 hours. Then the original culture medium was discard and 100 µL ECM was added in each well. Each group of cells were worked in triplicate. After 24 hours, 48 hours and 72 hours of continuous culture the original culture medium was discarded and 100 µL ECM containing 5 mg/mL MTT (Sigma-Aldrich, USA) was added into each well. After 4 hours of continuous...
culture, the liquid from each well was discarded and 100 μL DMSO (Aladdin, Shanghai, China) was added and samples were assessed at 750 nm.

\[ \text{cell inhibition rate (％)} = \frac{1 - \text{experimental group}}{\text{control group}} \]

Hoechst staining
The PCECs were collected, adjusted to the cell concentration of 1 × 10^6 cells/mL, inoculated in a 96-well plate at a concentration of 2 mL/well, covered with a glass, and cultured at 37°C in a 5% CO₂ incubator for 24 hours. Then the medium from the plate was discarded and a new ECM medium was added and further cultured at 37°C in a 5% CO₂ incubator for 48 hours. Afterwards culture, the medium from each well was discarded and 0.5 mL PBS solution was added and kept for 10 - 20 min. The wells were washed with PBS and 1 mL Hoechst33342 staining solution was added to each well. The cell morphology was observed and photographed under an inverted fluorescence microscope (IXplore SpinSR Super Resolution Microscope System, Plympus, Beijing, China).

Western blot assay and cytokine detection
The lung tissues were collected, rinsed with PBS, added with the corresponding volume of lysate was added, and left on ice for 5 minutes. The lysates were mixed and the samples were centrifuged at 12000 rpm, 4°C, for 10 minutes. The supernatant obtained represent the total protein extract of the cell. The protein concentration was determined by bicinchoninic acid assay (BCA Protein Assay Kit, ab102536, Abcam, China). The protein concentration in each group was equalized by adding 1×SDS gel buffer and then the protein samples were diluted with 5× loading buffer (ThermoFisher, USA) and PBS, boiled for 5 min and denatured. The protein liquid after denaturation was added to 12% SDS-PAGE gel (SDS-PAGE Gel kit, Elabscience, Wuhan, China) and underwent 80 V constant pressure electrophoresis for 60 - 120 min. The SDS-PAGE gel was placed on the nitrocellulose membrane to carry out constant current transfer. 3% BSA (bovine serum albumin, ThermoFisher, USA) was used in the final step. The nitrocellulose membrane was washed with PBS and monoclonal anti-HIF-1, anti-ET-1, anti-TGF-β1, anti-VEGF, anti-IL-1, anti-IL-6, anti-TNF-α, anti-HGF and anti-NAC antibody (Abcam, China), was added at 1:500 concentration. The membrane was incubated at room temperature for 45 minutes. Then the nitrocellulose membrane was washed with PBS buffer, and the incubation with the second antibody (Anti-mouse IgG for IP (HRP), ab131368, Abcam, China) was started. The second antibody labelled with horseradish peroxidase was added at 1:1000 and incubated at room temperature for 30 minutes. Then, the PBS buffer was used to wash the nitrocellulose membrane and the ECL chemiluminescence agent (Proandylbio, Xi’an, China) was used for chemiluminescence. Kodak X (Coolfilm, China) was used to expose the nitrocellulose membrane for about three minutes. The X-ray film was scanned and the gray analysis of the strip was processed by Gel-Pro Analyzer software (Software, Germany). The gray value of each band was measured in three independent replicates.

Statistical analysis
SPSS 21.0 software (IBM, USA) was employed. One way ANOVA was used for comparing the groups, Lenvene method was used for homogeneity of variance, Student-Newman-Keuls (SNK) method was used for comparison between two groups of data on the basis of homogeneity of variance. A value of p < 0.05 indicates significant differences.

Results and Discussion
Pulmonary artery pressure and right ventricular hypertrophy index
The PAP and RVHI significantly increased in the MCT group compared with the control group (Table I). The treatment with quercetin significantly decreased the level of PAP and RVHI compare with the MCT group, but without reaching the levels of the control group (Table I).

| Table I | Comparison of PAP and RVHI in rats of each group |
|---------|-------------------------------------------------|
|         | Control group | MCT group | Prevention group |
| PAP (mmHG) | 15.78 ± 2.59 | 44.13 ± 5.38 | 22.52 ± 6.21 |
| RVHI     | 0.231 ± 0.040 | 0.578 ± 0.120 | 0.289 ± 0.042 |

*p < 0.05 vs. the control group; *p < 0.05 vs. MCT the group

The relative expression levels of cytokine
According to the results of Western blot assay, the relative expression levels of HIF-1, ET-1, TGF-β1, VEGF, IL-1, IL-6 and TNF-α in lung tissues were significantly increased in the MCT group compared with the control group. The treatment with quercetin significantly increased the HIF-1, ET-1, TGF-β1, VEGF, IL-1, IL-6 and TNF-α levels compared with MCT group (Figure 1). The relative level of HGF and NAC were significantly decreased in the MCT group compared with the control group. The quercetin treatment reverses the MCT effect on HGF and NAC (Figure 1).

Proliferation and apoptosis of PCEC
PCEC in three groups of rats were cultured in the primary culture, and the proliferation and apoptosis of cells in different groups were compared. The MTT
and Hoechst staining assay showed that compared with the control group, the proliferation ability of PCEC in the MCT group was significantly decreased and apoptosis was significantly increased (Figure 2).

![Figure 1](image1.png)

**Figure 1.**
Western blot analysis of the expression of cytokines in lung rat tissues in each group

\*\* \( p < 0.01 \) vs. the control group; \#\# \( p < 0.01 \) vs. the MCT group

![Figure 2](image2.png)

**Figure 2.**
Detection of cell proliferation and apoptosis. A: Detection of cell proliferation using MTT assay. B: Quantitative results of Hoechst staining. C: Qualitative result of Hoechst staining

\* \( p < 0.05 \) vs. the control group; \# \( p < 0.05 \) vs. the MCT group

As a part of the human normal diet, quercetin has a long history. With the development of research on the biological and pharmacological activities of quercetin, more and more studies have shown that quercetin is beneficial to human health, including anti-cancer, anti-allergy, anti-inflammatory, anti-atherosclerosis and other effects [15-17]. Quercetin is a flavonoid with many bioactivities, high medicinal value and is widely distributed in vegetal products. In recent years, researchers focused on its properties and confirmed that it has a variety of biological effects.

Recently, it has been reported that quercetin can significantly alleviate pulmonary hypertension induced by MCT [18], but its potential molecular mechanism is still poorly understood. In this study, we found that quercetin can reverse pulmonary hypertension induced by MCT, which is consistent with previous results. The main pathological changes observed in PAH are pulmonary vasoconstriction, \textit{in situ} thrombosis and pulmonary vascular remodelling [19]. The results of this study showed that the injury of endothelial cells is the starting point in PAH, and the enhancement of
pulmonary vasoconstriction and the reconstruction of pulmonary vascular structure can be considered as the main characteristics of vascular changes after the injury of endothelial cells. The dysfunction of endothelial cells leads to the abnormal proliferation of intima and smooth muscle cells, the decrease of apoptosis, and finally the reconstruction of pulmonary vessels. Pulmonary vascular endothelial cells are considered to be the early mechanism of idiopathic PAH, which is accompanied by endothelial cell proliferation and apoptosis resistance.

The relative expression levels of HIF-1, ET-1, TGF-β1, VEGF, IL-1, IL-6 and TNF-α in lung tissues were significantly higher in the MCT group than in the control group, however, after the administration of quercetin the upregulation of inflammatory cytokines is reversed. The levels of HGF and NAC were decreased in the MCT group and significantly increased in the quercetin prevention group. Transcription factor NAC is an anti-apoptotic factor that modulates the expression of TNF-α and ICAM-1 (intercellular adhesion molecule 1). HGF regulates cell growth, cell motility, and morphogenesis by activating a tyrosine kinase signalling cascade after binding to the proto-oncogenic c-Met receptor, and acts as a multi-functional cytokine on cells of mainly epithelial origin [20]. Its ability to stimulate mitogenesis, cell motility and matrix invasion gives it a central role in angiogenesis, tumorigenesis, and tissue regeneration. HGF is a potent mitogen for mature parenchymal hepatocyte cells, it seems to be a hepatotrophic factor, and acts as a growth factor for a broad spectrum of tissues and cell types [20]. Previous studies have been shown that quercetin has directly reverses vasostriction caused by various endogenous factors (norepinephrine, ET-1, TXA2), protein kinase (PKC) activator and depolarizing factor (such as KCl) [21]. The molecular mechanism of this effect is not completely clear. It may be related to the multiple effects of PKC and myosin light chain kinase. The effect of PKC on resistance vessels is higher than on volume vessels. Quercetin at low concentration can inhibit the release of ET-1 and the transcription of its precursor by inhibiting tyrosine kinase in bovine aortic endothelial cells. Quercetin can inhibit the migration and proliferation of endothelial cells, mainly by blocking DNA synthesis to make it stagnate in cell cycle G0/G1, and can reduce the level of vascular endothelial growth factor by expressing cyclin inhibitory kinase 27 [21]. According to Hsieh et al. quercetin can inhibit the proliferation of pulmonary artery endothelial cells and induce apoptosis [22]. It acts by inhibition of cell proliferation through blocking the S-phase and G2-phase of cell cycle and upregulating p53 and p21 expression and the apoptotic Bax gene that induce apoptosis [22]. Toll-like receptor (TLR) plays an important role in the process of inflammation [23]. As an important transmembrane protein in the immune system, TLR can recognize various pathogenic related molecules and is an important barrier for the body to resist infectious diseases [24]. Among them, TLR4 is the earliest TLR that can mediate the body's response to pathogens. When the body is stimulated by relevant pathogenic molecules, TLR4 can induce the release of inflammatory cytokines such as IL-1 β, IL-6, TNF by activating nuclear factor-κB (NF-κB) [25-27].

Our in vitro experiments showed that compared with the control group, the proliferation ability of PCEC in MCT group was significantly decreased and apoptosis was significantly increased, while quercetin could inhibit the apoptosis of PCEC and promote cell proliferation. The results suggest that quercetin may also inhibit the expression of related cytokines and regulate cells through TLR4 mediated inflammatory signalling pathway.

Conclusions

Our study showed that quercetin had a therapeutic effect on PAH. It is suggested that the therapeutic effect of quercetin on PAH may be related to the regulation of cytokines, the promotion of cell proliferation and the inhibition of apoptosis. In addition, quercetin may also play a role in the treatment of PAH by inhibiting platelet aggregation and improving the function of vascular endothelial cells.

This study suggests that quercetin has a therapeutic effect on MCT induced PAH in rats and further clinical studies should investigate these findings also in humans in order to provide a new therapeutic alternative for PAH.

Conflict of interest

The authors declare no conflict of interest.

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