**EXPERIMENTAL STUDY**

**Nanoparticle-Mediated Targeting of Pitavastatin to Small Pulmonary Arteries and Leukocytes by Intravenous Administration Attenuates the Progression of Monocrotaline-Induced Established Pulmonary Arterial Hypertension in Rats**

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**Summary**

Statins are known to improve pulmonary arterial hypertension (PAH) by their anti-inflammatory and anti-proliferative effects in animal models. However, recent clinical studies have reported that clinically approved statin doses failed to improve clinical outcomes in patients with PAH. We therefore hypothesized that nanoparticle (NP)-mediated targeting of pitavastatin could attenuate the progression of established PAH.

We induced PAH by subcutaneously injecting monocrotaline (MCT) in Sprague-Dawley rats. On day 14 after the MCT injection, animals that displayed established PAH on echocardiography were included. On day 17, they were randomly assigned to the following 5 groups: daily intravenous administration of (1) vehicle, (2) fluorescein-isothiocyanate-NP, (3) pitavastatin, (4) pitavastatin-NP, or (5) oral sildenafil. Intravenous NP was selectively delivered to small pulmonary arteries and circulating CD11b-positive leukocytes. On day 21, pitavastatin-NP attenuated the progression of PAH at lower doses than pitavastatin alone. This was associated with the inhibition of monocyte-mediated inflammation, proliferation, and remodeling of the pulmonary arteries. Interestingly, sildenafil attenuated the development of PAH, but had no effects on inflammation or remodeling of the pulmonary arteries. In separate experiments, only treatment with pitavastatin-NP reduced the mortality rate at day 35.

NP-mediated targeting of pitavastatin to small pulmonary arteries and leukocytes attenuated the progression of established MCT-induced PAH and improved survival. Therapeutically, pitavastatin-NP was associated with anti-inflammatory and anti-proliferative effects on small pulmonary arteries, which was completely distinct from the vasodilatory effect of sildenafil. Pitavastatin-NP can be a novel therapeutic modality for PAH.

**Key words:** Translational study, Statin, Pulmonary hypertension

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models, which was associated with enhanced eNOS expression,9,10; activation of eNOS,11 and reduction of Rho- kinase expression in the lungs.9,12 However, recent clinical trials that tested the effect of statins on PAH with currently approved doses (atorvastatin 10 mg/day or simvas- tatin 80 mg/day) showed neutral results with regard to the clinical benefits for patients with PAH.14,15 These results suggested that the currently approved doses of statins were not sufficient to exert beneficial effects on PAH and that a modality to enhance their efficacy is warranted. Therefore, a drug delivery system (DDS) that facilitates the targeting of statins to the lung would have greater clinical implications.

Recently, we developed a bioabsorbable poly-lactic/ glycolic acid (PLGA)-nanoparticle (NP)-mediated DDS for pitavastatin16-22 and showed that direct intratracheal instillation of pitavastatin-NP attenuated the development of PAH, with reductions in inflammation and pulmonary artery remodeling in a rat model of monocrotaline (MCT)-induced PAH.20. Additionally, we reported that intravenously administered pitavastatin-NP was selectively delivered to tissues with enhanced vascular permeability by passive targeting and to tissues with inflammation by direct incorporation and delivery by circulating monocytes in a mouse model of atherosclerosis18,23 and myocardial infarction,24 and also in rat and porcine models of myocardial ischemia-reperfusion injury.16,17 Therefore, we hypothesized that intravenously administered pitavastatin-NP could be selectively delivered to small pulmonary arteries and inflammatory cells in the lung and exert its anti-inflammatory and anti-proliferative effects on PAH. In the present study, we tested the efficacy of intravenous administration of pitavastatin-NP in a preclinical rat model of PAH to verify our hypothesis and translate the therapeutic effect of pitavastatin-NP into clinical practice.

Methods

Preparation of PLGA nanoparticles: PLGA nanoparticles incorporating fluorescein-isothiocyanate (FITC; Dojin Chemical, Tokyo) (FITC-NP) or pitavastatin (Kowa Pharmaceutical Co. Ltd., Tokyo) (pitavastatin-NP) were prepared by an emulsion solvent diffusion method as described previously.16,20,23-31

Experimental animal models and protocols: All experiments were reviewed and approved by the Committee of Ethics of Animal Experiments, Kyushu University Graduate School of Medical Sciences. Male Sprague-Dawley (SD) rats (Charles River, Yokohama; weight 200-230 g) were subcutaneously injected with 60 mg/kg of MCT (Sigma-Aldrich, MO), which induces severe PAH in 3 weeks.20 Pulmonary artery acceleration time/cycle length (PAAT/CL) was measured by echocardiography (Vevo 2100 ultrasound system; Visual Sonic, Toronto) in each animal at day 14 after the MCT injection and was used as a noninvasive estimation of pulmonary artery systolic pressure.20 Animals with PAAT/CL < 0.140 were assumed to have developed PAH and were included in the study. Experimental protocols are shown in Figures 1-3.

Echocardiographic measurement of RV and pulmonary artery hemodynamics: Transthoracic 2-D and pulsed-wave Doppler echo were obtained with a 30-MHz transducer. Echocardiographic images were obtained as described previously.20 Pulsed-wave Doppler was used to measure PAAT. During echocardiography, the animals were anesthetized with isoflurane (1%, AbbVie, IL).

Flow cytometry of blood, lung, and bronchiolar alveolar lavage fluid (BALF): Peripheral blood was lysed with VersaLyse lysing solution (Becton Dickinson Bioscience, CA) for 15 minutes at room temperature. The lungs were isolated and digested with a cocktail of 450 U/mL collagenase type I, 125 U/mL collagenase type XI, 50 U/mL DNase I, and 60 U/mL hyaluronidase (Sigma-Aldrich) in phosphate-buffered saline (PBS) containing 20 mM Hepes at 37°C. BALF was collected in cold PBS. The cell suspension was centrifuged at 300 × g for 5 minutes at 4°C. After blocking the Fc receptor with anti-CD16 antibody (Becton Dickinson Bioscience), cell suspensions were incubated with anti-CD11b antibody (Becton Dickinson Bioscience) or an isotype control for 1 hour at 4°C. Finally, they were incubated with 7-AAD (Becton Dickinson Bioscience) for 5 minutes at room temperature and analyzed with Gallios (Beckman Coulter, CA).

Direct hemodynamic measurement: The animals were
Figure 2. Experimental protocol for treatment. Scheme showing the experimental protocol for treatment.

anesthetized with isoflurane (1%), and polyethylene catheters (PE10, Becton Dickinson, NJ; internal diameter, 0.28 mm) were inserted into the right ventricle through the right jugular vein as described previously. RV systolic pressure (RVSP) was measured with a polygraph system (PowerLab, AD Instruments, Dunedin, New Zealand).

Assessment of RV hypertrophy: The RV wall was dissected from the left ventricle and the ventricular septum. The right and left ventricles plus the ventricular septum were weighed, and the degree of right ventricular hypertrophy (RVH) was quantified as RV weight/[left ventricle plus ventricular septum weights].

Histopathological and immunohistochemical analyses of the lungs: For paraffin sections, lungs were inflated with 1% formalin plus 0.5% agarose at 20 cm H2O pressure and fixed in 10% formalin overnight. Then, they were embedded in paraffin and cut into 5-μm sections. Sections were incubated overnight at 4°C with the following primary antibodies: anti-α-smooth muscle actin (SMA) antibody (dilution 1:500; clone 1A4, Dako, MI), anti-CD68 antibody (dilution 1:100; AbD Serotec, NC), anti-NF-κB antibody (dilution 1:1,000; p65 subunit, clone 12H11, Merck Millipore, MA), and anti-PCNA antibody (dilution 1:400; Dako), followed by incubation with biotin-conjugated secondary antibodies.

The degree of remodeling in small peripheral pulmonary arteries was assessed by counting the number of SMA-positive pulmonary arteries by a blinded observer, as described previously. To quantify the degrees of monocyte infiltration and NF-κB activity in the lung, a blinded observer counted the number of ED-1-positive or NF-κB-positive cells in 5 high-power fields. To quantify the degree of vascular cellular proliferation, a blinded observer counted the number of PCNA-positive cells in 10 intra-acinar arteries with diameters of 25-50 μm in each rat.

For OCT-embedded sections, the lungs were inflated with a mixture of 1:1 OCT compound (Sakura Finetchnical Co. Ltd., Tokyo) and PBS at 20 cm H2O pressure and embedded in OCT compound at −80°C. Sections of 10 μm were cut and incubated with the following antibodies: anti-SMA antibody (dilution 1:200; Thermo Scientific, MA) and anti-von Willebrand factor (vWF) antibody (dilution 1:3,000, Dako), followed by incubation with Alexa Fluor 555-conjugated secondary antibody (anti-rabbit IgG Alexa Fluor 555 antibody, dilution 1:1,000, Invitrogen, MA). The nuclei were counterstained with DAPI. To quantify the fluorescence intensity of FITC, the mean fluorescence intensity of 20 randomly selected high-power fields was measured by ImageJ.

Ex-vivo angiography of pulmonary artery by Microfil: Fasudil hydrochloride hydrate (10 mg/kg) (Eril; Asahi Kasei Pharmaceutical Corporation, Tokyo) and heparin (100 units/100 g body weight) were injected into the femoral vein. Diluted heparin sodium (5 units/mL, 20 mL) was injected via the right ventricle, and the heart and lungs were extracted. Polyethylene tubing (PE50, Becton Dickinson; internal diameter, 0.53 mm) was advanced to the main pulmonary artery and secured in place using 5-0 silk sutures. A small incision was made in the left atrial appendage, and 2 mL of diluted heparin sodium was pumped at 2 mL/minute using a continuous syringe pump.
A silicone polymer casting compound, MV-yellow Microfil (Flow Tech, MA), was mixed with a medium-viscosity diluent in a 5:4 (diluent:compound) volume ratio and added to 5% (by volume) of the curing agent. This mixed silicone polymer casting material was pumped through the catheter at 0.1 mL/minute until the polymer was uniformly visible on the lung surface.33) The lungs were inflated with 1% formalin plus 0.5% agarose at 20 cm H2O pressure. After complete polymerization at 4°C for 24 hours, the heart and lungs were fixed in 10% formalin. The lungs were sequentially bathed in serial concentrations of ethanol. The tissues were then placed in methyl salicylate and photographed with a stereoscopic microscope.34)

Measurements of cytokine, chemokine, and growth factor levels in the lung by multiplex immunoassay: Cytokine, chemokine, and growth factor levels in the lung were measured with a cytokine magnetic-based assay: Bio-Plex Pro Rat Cytokine 24-Plex Panel (Bio-Rad, CA), according to the manufacturer’s instructions. A heat map showing the means of the z-scores (z = (x-mean)/SD) was used only to show the trend of cytokine/chemokine and growth factor expressions in each group.

Statistical analysis: The data are presented as the means ± standard deviation (SD) and were analyzed using the Prism Software program (Graph Pad Software, San Diego, CA). The statistical analysis of differences in 2 groups was performed using the t-test. The statistical analysis of differences between more than 3 groups was performed using the one-way analysis of variance (ANOVA) followed by Dunnett’s tests. The bar graphs in Figure 5B and 9C were compared using two-way ANOVA, followed by Tukey’s multiple comparison tests. The survival curves were described with the Kaplan-Meier method, and the survival rates between the treatment groups were compared with the log-rank test. P values < 0.05 were considered statistically significant.

Results

Localization of intravenously administered FITC-NP in the lung and CD-11b-positive leukocytes in rats with MCT-induced PAH: We evaluated the localization of NP in the lung after the daily administration of FITC or FITC-NP in both MCT-induced PAH rats and normal rats. In MCT-induced PAH rats, strong FITC signals were observed in the vascular walls of the SMA-positive vessels and the bronchial epithelium in FITC-NP-injected animals, but only faint signals were detected in FITC-alone-injected animals (Figure 4). FITC signals co-localized with SMA (Figure 5), but not with vWF (Figure 6). In normal rats, less significant FITC signals were observed in FITC or FITC-NP-injected animals (Figures 4, 5).

Flow cytometry showed that animals treated with FITC-NP had significantly stronger FITC signals in CD11b-positive leukocytes in the blood, lung, and BALF than those treated with FITC alone (Figure 7).

Effects of pitavastatin-NP on the development of PAH in the rat model of MCT-induced PAH: MCT injections induced severe PAH, which was characterized by elevated RVSP (Figure 8A), development of RVH (Figure 8B), and pulmonary artery remodeling (Figure 8C, D). On day 14, PAAT/CL measured using echocardiography was > 0.140 in 9 of 108 animals, indicating low pulmonary artery systolic pressure, and these animals were excluded. On day 17, the included 99 animals were randomized into 4 groups and received intravenous treatment with vehicle, FITC-NP, pitavastatin alone, or pitavastatin-NP in several doses. Treatment with pitavastatin-NP containing 1 or 3 mg/kg of pitavastatin significantly attenuated the elevation of RVSP, the development of RVH, and pulmonary artery remodeling compared with the control nanoparticle treatment (FITC-NP) and with the same dose of pitavastatin alone. Treatment with pitavastatin alone at the dose of 30 mg/kg, but not at doses of 1, 3, or 10 mg/kg, also attenuated the development of RVH and pulmonary artery remodeling compared with vehicle treatment. These data
Figure 4. Delivery of FITC-NP to the lung. Representative bright field and fluorescence microscopic views of lung sections in normal rats or MCT-induced PAH rats after intravenous injection of FITC or FITC-NP. Fluorescence views are immunostained with smooth muscle actin (SMA; red). The nuclei are counterstained with DAPI. Scale bar: 1 cm.

Figure 5. Magnified view of Figure 4. A: Representative magnified view of lung sections. Scale bar: 100 μm. B: Bar graph showing the quantitative measurement of FITC fluorescence intensity of lung sections shown in A. n = 20 each.
suggested that pitavastatin-NP had approximately 30 times higher efficacy than pitavastatin alone in attenuating the development of PAH induced by the MCT injection.

We also performed ex-vivo angiography of the pulmonary arteries with Microfil to assess the effect of pitavastatin-NP on the development of small pulmonary artery stenosis and obstruction 3 weeks after the MCT injection (Figure 9A). The degree of small pulmonary artery stenosis and obstruction was quantified by counting the number of small vessels branching from the 2nd, 3rd, or 4th branch of the main pulmonary artery (Figure 9B). The number of patent vessels branching from the 3rd or 4th branch was significantly reduced in the vehicle and pitavastatin-alone (3 mg/kg) groups, but it was preserved in the pitavastatin-NP (3 mg/kg of pitavastatin) group (Figure 9C).

**Effects of pitavastatin-NP on inflammation in rats with MCT-induced PAH:** Treatment with pitavastatin-NP containing 1 or 3 mg/kg of pitavastatin significantly attenuated the infiltration of CD68-positive monocytes, and the increase in NF-κB activity induced by MCT injection compared with the control nanoparticle treatment (FITC-NP) and with the same dose of pitavastatin alone. Also, treatment with pitavastatin alone at the dose of 30 mg/kg, but not at doses of 1, 3, or 10 mg/kg, also attenuated the increase in NF-κB activity compared with vehicle treatment (Figure 10A, B). These data suggested that NP-mediated targeting of pitavastatin to the lung by intravenous injection is more effective in blocking the inflammatory pathway and monocyte infiltration than pitavastatin alone.

We also performed a multiplex immunoassay to measure cytokines, chemokines, and growth factors simultaneously. The MCT injection induced a significant elevation in the cytokine/chemokine levels of the lung associated with monocytes/macrophages, such as MCP-1, MIP-1α, GRO/KC, and IL-1α. On the other hand, the MCT injection suppressed cytokines/chemokines associated with lymphocytes, such as IL-2, IL-4, IL-7, IL-10, IL-18, and IFN-γ. Pitavastatin-NP attenuated the elevation of monocyte/macrophage-associated cytokines/chemokines, such as MIP-1α, MIP-3α, GRO/KC, M-CSF, and IL-1α, compared to the vehicle, but it had no influence on lymphocyte-associated cytokines/chemokines (Figure 11).

**Comparison of the effects of pitavastatin-NP with PDE5 inhibitor sildenafil on MCT-induced PAH:** To compare the effects of pitavastatin-NP with the currently approved vasodilatory therapy, we administered the PDE5
Effect of pitavastatin-NP on the survival rate of rats with MCT-induced PAH: Finally, we examined the effect of pitavastatin-NP on the survival rate of rats with MCT-induced PAH on day 35. Compared with treatment with vehicle, FITC-NP, or sildenafil, treatment with pitavastatin-NP significantly improved the survival rate: 18%, 25%, 30%, and 61% in the vehicle- (n = 34), sildenafil- (n = 28), FITC-NP- (n = 37), and pitavastatin-NP-treated groups (n = 23), respectively (Figure 14).

Discussion

The major findings in this study are as follows: (1) FITC-NP was selectively delivered to the small pulmonary arteries and inflammatory cells in the lung after intravenous administration in a rat model of MCT-induced PAH; (2) pitavastatin-NP attenuated the progression of established PAH and improved survival; and (3) the beneficial effects of pitavastatin-NP were associated with anti-inflammatory and anti-proliferative effects, which were mechanisms distinct from those of the PDE5 inhibitor, sildenafil.

PLGA-NP is known to distribute and accumulate in tissues with enhanced vascular permeability after intravenous administration. As vascular permeability is enhanced in PAH, intravenous PLGA-NP is probably delivered to the pulmonary vascular wall by passive targeting. In the present study, significant FITC signals were noted in the SMA-positive small pulmonary artery walls after the daily administration of FITC-NP. Additionally, NP was found to be incorporated into CD11b-positive inflammatory cells in the peripheral blood, lung, and BALF. As alveolar macrophages in the BALF are known to be derived largely from circulating monocytes, these data suggested that intravenously administered PLGA-NP was incorporated by circulating monocytes and delivered to the alveolar space in the lung. These results indicated the benefit of intravenously administered PLGA-NP as a promising DDS in PAH.

Inflammation has been recently considered as a key pathophysiological factor that contributes to remodeling in PAH. Infiltrates of inflammatory cells to the lung and also inflammatory cytokines mediate the pathological features in patients with PAH. For example, IL-6 is reported to promote the development and progression of pulmonary vascular remodeling by activating anti-apoptotic and pro-proliferative pathways. Although several anti-inflammatory therapies, such as dexamethasone, FK506, IL-1 receptor antagonist, leukotriene A4 hydrolase, or CCR5 antagonist, have been investigated in animals, none of them have been clinically approved for the treatment of PAH. MCP-1 and its receptor CCR2 pathway, a key regulator of monocyte recruitment, have been shown to be involved in the pathogenesis of inflammation in cardiovascular diseases, including PAH. Plasma and lung tissue MCP-1 levels are reported to be increased in patients with PAH, which decreased after treatment with epoprostenol. In addition, treatment with anti-MCP-1 antibody or transfection with a dominant negative inhibitor of MCP-1 attenuated MCT-induced
PAH in the MCT-induced PAH rat model. These results showed that monocyte-mediated inflammation mediated by the MCP-1/CCR2 pathway can be a promising target in the treatment of PAH.

Statins are known to inhibit the activation of NF-κB and directly activate Akt, which are both key regulators of MCP-1.\textsuperscript{53,54} Previously, we showed that pitavastatin-NP inhibited the MCP-1 induced chemotaxis of monocytes \textit{in vitro}\textsuperscript{18} and also inhibited the expression of MCP-1 in the ischemic myocardium of the ischemia-reperfusion injury rat model \textit{in vivo}\textsuperscript{17} compared with pitavastatin alone. We also observed that the cardioprotective effect of pitavastatin-NP on ischemia-reperfusion injury in wild-type mice was blunted in CCR2-deficient mice (unpublished observation in our laboratory). In the present study, pitavastatin-NP significantly reduced the infiltration of CD68-positive monocytes/macrophages in the lungs of rats with MCT-induced PAH. Additionally, the result of the multiplex immunoassay suggested that pitavastatin-NP attenuated the elevation of cytokines/chemokines associated with monocytes/macrophages induced by MCT. Additionally, the fact that cytokines/chemokines associated with lymphocytes were significantly downregulated supports previous reports that showed that lymphocyte accumulation, such as T cells and NK cells, is attenuated in the lungs of MCT-induced PAH rats.\textsuperscript{55,56} Collectively, these results suggest that the therapeutic effect of pitavastatin-NP was mediated by inhibiting monocyte-mediated inflammation by blocking the MCP-1/CCR2 pathway in the lung. Some subtypes of PAH, such as systemic sclerosis-associated PAH, are known to be strongly associated with monocyte-mediated inflammation.\textsuperscript{56} However, as dysregulated immunity is a key pathophysiological factor of all PAH subtypes,\textsuperscript{2} the anti-inflammatory effect of pitavastatin-NP deserves to be tested in any subtype of PAH.

Currently approved drugs for the treatment of PAH (PDE5 inhibitors, endothelin receptor antagonists, prostacyclins, etc.) are thought to deliver their therapeutic effects mainly through vasodilating activities, but not through direct anti-inflammatory and anti-proliferative effects.\textsuperscript{2} In contrast, statins are known to inhibit the proliferation of vascular SMCs down-regulating Rho, which leads to a reduction in MAPK activity, a strong regulator of SMC proliferation.\textsuperscript{57} Statins also induce vasodilatory effects in some settings by the secondary activation of eNOS and prostacyclin synthase systems by down-regulating Rho.\textsuperscript{8,58} Previously, we showed that pitavastatin-NP directly inhibited the FBS-induced proliferation of human endothelial cells and pulmonary artery
In our short-term treatment protocol shown in Figure 3, intravenous pitavastatin-NP had no acute effects on RVSP, but treatment with the PDE5 inhibitor sildenafil decreased the elevated RVSP in the MCT-induced PAH model. Although we did not examine whether treatment with pitavastatin-NP activates eNOS or prostacyclin synthase systems in the chronic phases, these data suggest that the mechanism of pitavastatin-NP (mainly anti-inflammatory and anti-proliferative effects) in ameliorating MCT-induced PAH is different from that of sildenafil (vasodilating actions).

Clinical trials of statins that failed to show clinical benefits in patients with PAH suggested that the oral administration of statins at clinically approved doses appears to be insufficient to exert therapeutic effects. On the other hand, the therapeutic dose of statins in animal studies that showed its beneficial effects in PAH was relatively high (simvastatin 2-20 mg/kg/day, fluvastatin 1 mg/kg/day, or rosuvastatin 2-10 mg/kg/day). Since the oral administration of high-dose statins in humans is known to be harmful due to serious side effects, such as rhabdomyolysis, selective DDS with intravenous PLGA-NP can reduce the total amount of statin required. Indeed, in the present study, 1 mg/kg of pitavastatin-NP was equivalent to 30 mg/kg of pitavastatin alone relative to the reduction of RVSP. We have confirmed the safety limit of pitavastatin-NP in rats as 2 mg/kg (unpublished data in our laboratory), suggesting that its intravenous administration at the dose of 1 mg/kg can be safe and feasible. However, as these data are derived only from rodent models, it is difficult to estimate the appropriate doses of pitavastatin-NP for patients with PAH. Thus, phase II clinical trials are needed to confirm the efficacy and safety of pitavastatin-NP for PAH in humans.

In summary, nanoparticle-mediated targeting of pitavastatin to the small pulmonary arteries and inflammatory cells by intravenous administration attenuated the progression of established PAH and improved the survival in the MCT-PAH model, which was associated with anti-inflammatory and anti-proliferative effects. We have completed the phase I clinical trial on intravenously administered pitavastatin-NP (UMIN 000014940, UMIN 000019189) and confirmed the safety and tolerability in healthy subjects; a phase II clinical trial for patients with PAH is now under contemplation. Pitavastatin-NP can be developed as a novel category of therapeutics for PAH.
**Figure 10.** Effects of pitavastatin and pitavastatin-NP on the infiltration of monocytes and NF-κB activation 3 weeks after the MCT injection. **A:** Immunostaining with CD68 of lungs from rats in the 8 experimental groups. The bar graph indicates the number of CD68-positive monocytes per 5 high-power fields (HPFs) in each group. Scale bar: 50 μm, n = 8-10 each. **B:** Immunostaining with NF-κB (α-p65) of lungs from rats in the 8 experimental groups. The bar graph indicates the number of NF-κB (α-p65) -positive cells per 10 HPFs in each group. Scale bar: 50 μm, n = 8-10 each. *P < 0.05, **P < 0.01, ***P < 0.001 versus pitavastatin-alone and pitavastatin-NP groups with the same dose. #P < 0.01 versus the FITC-NP group. †P < 0.05, ††P < 0.01 versus the vehicle group.

**Figure 11.** Cytokine/chemokine levels in the lung measured by multiplex immunoassay. **A:** Heat map showing the means of the z-scores of the expression of cytokines/chemokines in the lungs of normal rats and MCT-induced PAH rats treated with the vehicle of pitavastatin-NP. The Z-score was calculated as z = (x - mean) / SD. **B:** Table showing the levels of cytokines/chemokines in the lungs of normal rats and MCT-induced PAH rats treated with the vehicle of pitavastatin-NP. P values shown in the table are versus the normal rat group. *P < 0.05, **P < 0.01 versus the vehicle group.
**Figure 12.** Comparisons of the effects of pitavastatin-NP and sildenafil on MCT-induced PAH. A: RVSP in the 4 experimental groups 3 weeks after the MCT injection. n = 8-10 each. B: RVH assessed as the ratio of the weight of RV/(left ventricle (LV) + septum (S)) in the 4 experimental groups 3 weeks after the MCT injection. n = 8-10 each. C: Remodeling of the small pulmonary arteries assessed as the percentage of fully muscularized pulmonary arteries in the 4 experimental groups 3 weeks after the MCT injection. n = 8-10 each. D: Immunostaining with CD68 of lungs from rats in the 4 experimental groups. The bar graph indicates the number of CD68-positive monocytes per 5 high-power fields (HPFs) in each group. Scale bar: 50 μm. n = 8-10 each. E: Immunostaining with NF-κB (α-p65) of lungs from rats in the 4 experimental groups. The bar graph indicates the number of NF-κB (α-p65)-positive cells per 10 HPFs in each group. Scale bar: 50 μm. n = 8-10 each. F: Immunostaining with PCNA of lungs from rats in the 4 experimental groups. The bar graph indicates the ratio of the number of PCNA-positive cells/total numbers of vascular cells in each groups Scale bar: 50 μm. **P < 0.01 versus the vehicle group. *P < 0.05, **P < 0.01 versus the normal rat group. n = 8-10 each.

**Figure 13.** Acute hemodynamic effects of pitavastatin-NP and sildenafil in MCT-induced PAH rats. Results of continuous hemodynamic measurements. Changes in RV systolic pressure (RVSP) and arterial systolic pressure (sAP) from baseline were plotted for 1 hour. n = 3-4 each.
Disclosures

Conflicts of interest: Dr. Egashira holds a patent on the results reported in the present study. Dr. Tsutsui reports receiving grant supports from Astellas Pharma and Daiichi-Sankyo, lecture fees from Astellas Pharma, Otuka Pharmceuticals, Takeda Pharmaceuticals, Daiichi-Sankyo, Daiichi-Sankyo, lecture fees from Astellas Pharma, Otuka receiving grant supports from Astellas Pharma and Dr. Egashira holds a patent on the

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