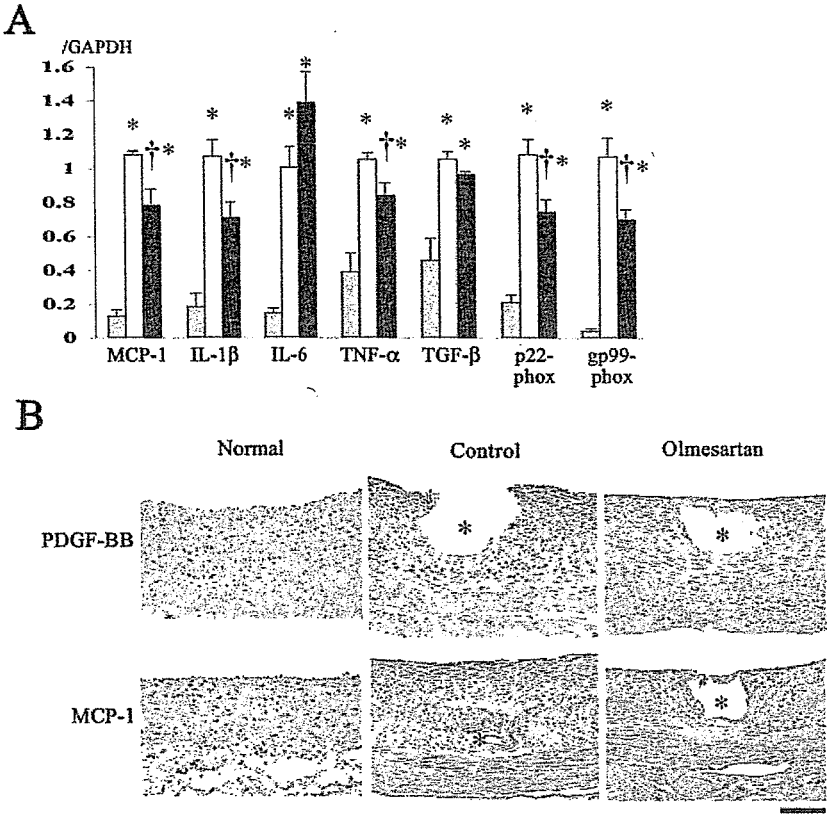

Figure 3. Effects of ARB on gene expression of proinflammatory factors, and immunohistochemical expression of PDGF and MCP-1. a, Effect of olmesartan on relative mRNA levels of various proinflammatory factors and NADPH oxidases 7 days after stenting in normal controls (▨, n=8), the no-treatment group (□, n=8), and the ARB-treated group (■, n=8). *$P<0.01$ vs uninjured normal artery; †$P<0.05$, vs the control group. b, Iliac artery sections from the uninjured normal animals and those from the control and olmesartan groups 10 days after stenting stained immunohistochemically with PDGF-BB and MCP-1. * indicates stent strut. Bar=100 μm. These immunohistochemical experiments were repeated 5 times, all with representative results.

ment with olmesartan partly attenuated the increased DHE fluorescence after stent implantation.

Inhibitory Effects of ARB on Inflammatory Changes and Apoptotic Cell Death in Rabbits

As we reported previously,[15] inflammatory changes and apoptotic cell death became evident 7 to 10 days after stent implantation in rabbits (Figure 2a and 2b). Treatment with olmesartan reduced such inflammatory changes and enhanced cell death in the intima after stenting.

Inhibitory Effects of ARB on Expression of Proinflammatory Factors and NADPH Oxidase Subunits

Treatment with olmesartan reduced the increased mRNA levels of monocyte chemoattractant protein (MCP)-1, interleukin (IL)-1β, tumor necrosis factor-α, p22phox, and gp91phox in rabbits (Figure 3a). Olmesartan did not affect the increased levels of IL-6 and transforming growth factor-β. Immunohistochemical staining performed 10 days after stenting revealed increased immunoreactive platelet-derived growth factor (PDGF)-β and MCP-1 in cells in the neointima and in smooth muscle cells in the media. This was attenuated by olmesartan treatment (Figure 3b). Treatment with olmesartan did not affect neovascularization in the neointima and adventitia or re-endothelialization 28 days after stenting (data not shown).

Effects of ARB on Transdifferentiation of Mononuclear Cells to Vascular Progenitor Cells

To investigate the potential contribution of vascular progenitor cells, peripheral blood mononuclear cells (MNCs) were isolated and cultured to stimulate the differentiation into SMPCs or endothelial progenitor cells (EPCs), as described previously.[16,17] The cells cultured in the PDGF-BB–enriched and basic fibroblast growth factor–enriched medium exhibited a hill and valley morphology that is characteristic of smooth muscle cells within 2 weeks. The smooth muscle cell phenotype was confirmed by immunostaining with antibodies specific for smooth muscle cell markers: SMPCs expressed α-smooth muscle actin (SMA), myosin, and calponin, which were all detected in human coronary artery smooth muscle cells and were not detected in MNCs and Cos-7 cells (data not shown). Expression of α-SMA gene in SMPCs was also confirmed by PCR analysis (data not shown). As reported,[16,17] the expression of inflammatory markers (MCP-1, IL-1β, etc) was greater in SMPCs than in cultured rabbit aortic smooth muscle cells (data not shown). The cells cultured in the vascular endothelial growth factor–enriched medium exhibited the typical cobblestone morphology of EPCs. The EPCs stained positively for von Willebrand factor and VE-cadherin and incorporated acetylated low-density lipoprotein (data not shown).

Analysis of colony-forming areas showed that the degree of transformation to SMPC was greater in MNCs from animals fed a high-cholesterol diet than in those from untreated, normal animals (Figure 4A). The transformation to SMPCs further enhanced in MNCs from animals that underwent stenting. Treatment of rabbits with olmesartan for 5 to 7 days suppressed the increased transformation to SMPCs induced by stenting. In contrast, there were no differences in the degree of transformation to EPCs among the groups. Immunohisto-

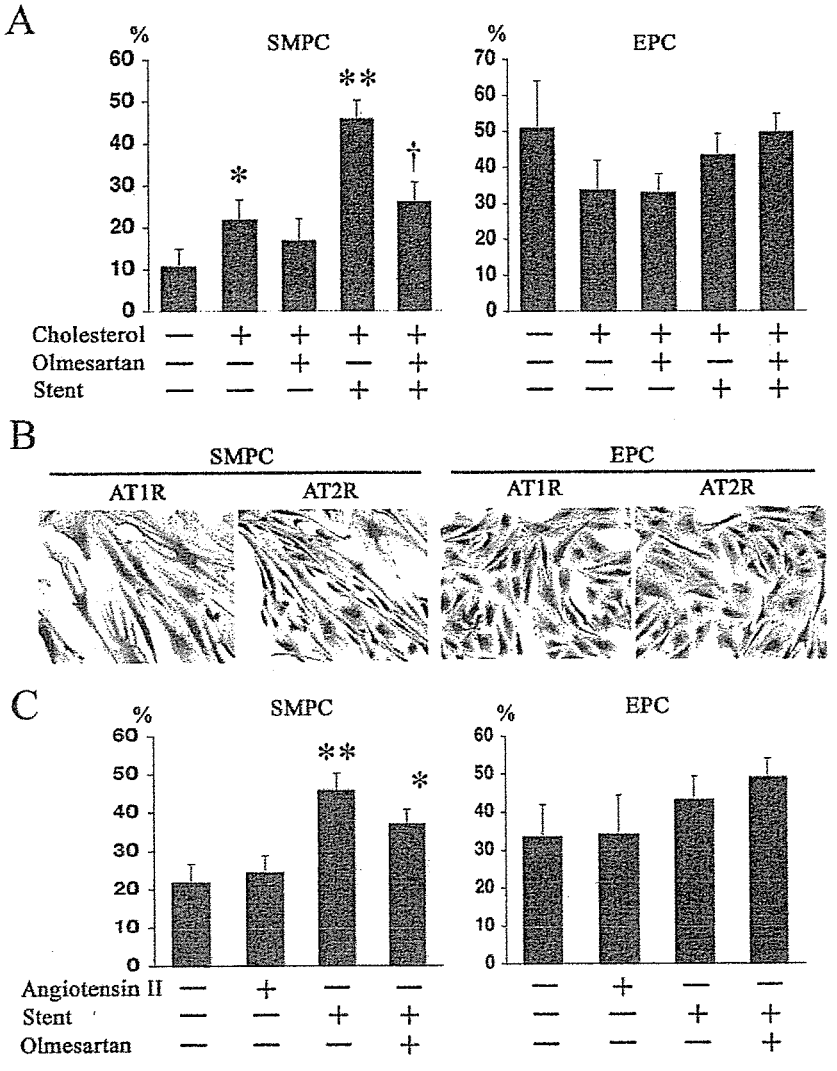

Figure 4. Effects of ARB on transdifferentiation of MNCs to SMPCs or EPCs. a, The degrees of transformation of MNCs to SMPCs (the percentages of α-SMA–positive area per well) or EPCs (the percentages of von Willebrand factor–positive area per well) in normal rabbits and those that received the high-cholesterol diet, high-cholesterol diet plus in vivo olmesartan treatment, high-cholesterol diet and stenting, or high-cholesterol diet, in vivo olmesartan treatment, and stenting (n=8 to 9 each). *$P<0.05$, **$P<0.01$ vs control (cholesterol [−] olmesartan [−] stent [−]). †$P<0.05$ vs high-cholesterol diet plus stenting (cholesterol [+] olmesartan [−] stent [+]). b, Immunohistochemistry for AT1 receptor and AT2 receptor in SMPCs and EPCs. c, The degree of transformation of MNCs from rabbits fed a high-cholesterol diet to SMPCs or EPCs. The effects of in vitro addition of angiotensin II were examined. In addition, the in vitro effects of olmesartan on the stent-induced increase in transformation were examined (n=8 to 9 each). *$P<0.05$, **$P<0.01$ vs control (angiotensin II [−] olmesartan [−] stent [−]).

chemical staining was then performed to examine the presence of AT_1 receptor and AT_2 receptor. Both receptors were found in SMPCs and EPCs (Figure 4B). We considered the possibility that AT_1 signals might be involved in increasing the transformation capacity of MNCs and, therefore, examined the effects of in vitro administration of angiotensin II or olmesartan on the transformation of MNCs. Angiotensin II did not enhance transdifferentiation, and olmesartan did not suppress transdiffrentiation in vitro (Figure 4C).

Plasma ARB levels and Arterial Blood Pressure

The maximum drug concentration (C_{max}) levels of olmesartan at 15 mg/kg per day and valsartan at 50 mg/kg per day were 107±17 and 300±24 ng/mL, respectively. The C_{max} level of olmesartan at 3 mg/kg per day was 537±24 ng/mL in rabbits. Therefore, the dose of olmesartan used in rabbits is within a clinically relevant dose range. The C_{max} levels after oral administration of olmesartan at 5, 10, and 20 mg/body in hypertensive subjects are reported to be 149±21, 273±17, and 470±23 ng/mL (n=6, each), respectively. The C_{max} values after oral administration of valsartan at 80 and 160 mg/body in hypertensive subjects are reported to be 2830±920 and 5260±2300 ng/mL, respectively, according to the manufacturer's interview form. Thus, the doses of olmesartan and valsartan used in the present study are within or below the clinically relevant dose range.

Treatment with olmesartan showed no effect on systolic and diastolic arterial pressure. Systolic and diastolic pressure were 94±2 and 59±5 mm Hg in the control group and 92±2 and 52±6 mm Hg in the olmesartan group.

Discussion

We have demonstrated for the first time that oral treatment with 2 types of ARBs (valsartan and olmesartan) attenuated in-stent neointimal formation in nonhuman primates (cynomolgus monkeys), supporting the conclusions of the Val-PREST and VALVACE trials,[12,13] which involved a relatively small number of patients. Although it is uncertain which animal model is most appropriate for the evaluation of in-stent neointima formation (restenotic changes), a nonhuman primate model may have an advantage over nonprimate animal models, because vascular inflammatory and proliferative responses to injury in nonhuman primates are more similar to those in humans than are other, nonprimate models.

Hence, the use of nonhuman primates may work for evaluation of the efficacy of ARB on in-stent neointima formation in clinically relevant conditions.

To obtain mechanistic insight into the beneficial effects of ARB, we first examined the time course of local expression of RAS components in rabbits (Figure 2). We found that expression of all of the components (ACE, angiotensin II, AT_1 receptor, and AT_2 receptor) increased, mainly in cells composed of neointima (monocytes and smooth muscle cells), at early stages (7 to 10 days after stenting), and persisted until 28 days after stenting. This RAS activation colocalized with increased NADPH oxidase–dependent DHE fluorescence (generation of superoxide anion) and was associated with increased levels of NADPH oxidase subunit mRNAs, consistent with previous reports showing that increased reactive oxygen species can be detected in activated smooth muscle cells after balloon injury.[21,22] These in vivo observations are also consistent with previously published in vitro data suggesting that proliferation and migration of smooth muscle cells are critically mediated by oxidative stress via AT_1-mediated activation of NADPH oxidases.[23–29] Interestingly, treatment with ARB not only attenuated the levels of oxidative stress markers but also reduced the level of immunoreactive AT_1. These data suggest the presence of a positive feedback loop in which activation of AT_1 further enhances expression and activity of the AT_1 receptor in vivo, as seen in the present study.

It is known that oxidative stress–induced inflammatory and proliferative processes are central to neointima formation after vascular injury.[24,25] We and others have demonstrated that[1] increased monocyte-mediated inflammation or MCP-1 expression is associated with greater neointima formation after stenting,[26,27] and[2] anti–MCP-1 gene therapy[15,28–30] or administration of blocking antibody against the MCP-1 receptor[31] markedly reduces neointima formation after vascular injury. However, no previous study examined whether or not those inflammatory and proliferative changes after stenting depends on the AT_1 receptor. In the present study, we, therefore, examined the effects of ARB on monocyte recruitment and MCP-1 expression after stenting and found that ARB reduced monocyte/macrophage recruitment, as well as MCP-1 immunoreactivity and gene expression. Furthermore, ARB inhibition increased the expression of growth-promoting factors, such as PDGF and IL-1β. These data suggest that the beneficial effects of ARB may be attributed to the inhibition of oxidative stress–induced inflammatory and proliferative changes.

Recent studies have shown that peripheral blood contains bone marrow-derived progenitor cells, which contribute to neointima formation after injury.[16,18,32] However, the role of RAS in the recruitment/differentiation of progenitor cells into the neointimal cells after stenting has not been addressed. Here we found that differentiation to SMPCs increased in MNCs from rabbits fed a high-cholesterol diet and was further enhanced in those rabbits that had also undergone stenting. Differentiation into EPCs was not affected by either the diet or stenting. In vivo treatment with ARB suppressed the increased differentiation into SMPCs induced by diet or stenting. In contrast, in vitro treatment with angiotensin II or ARB did not affect the capacity to differentiate into SMPCs or EPCs. Therefore, the capacity to recruit or form SMPCs from MNCs after stenting might be determined by an AT1 receptor–mediated pathway in vivo and, thus, contribute to in-stent neointima formation.

It must be mentioned that ARB did not significantly reduced arterial blood pressure in rabbits. Although arterial pressure was not measured in monkeys, the dose of ARB used in the present study is reported to show no effects on arterial blood pressure in monkeys.[33] Plasma ARB level was within or below the clinical range. Furthermore, ARB did not affect serum lipid levels. Therefore, the beneficial effect of ARB on in-stent neointimal formation is likely to be independent of its effects on arterial blood pressure or serum lipid.

Perspectives

This study provides experimental evidence suggesting that oral treatment with ARB at a clinical dose range attenuates in-stent neointima formation in rabbits and nonhuman primates. The beneficial effects were associated with reduced local oxidative stress, reduced expression of MCP-1 and other inflammation-promoting factors, and reduced recruitment/differentiation of SMPCs, suggesting that ARB is of potential clinical benefit in patients who have undergone vascular interventions.

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Disclosures

None.

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Ovariectomy Augments Hypertension Through Rho-Kinase Activation in the Brain Stem in Female Spontaneously Hypertensive Rats

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Abstract—Estrogen protects against increases in arterial pressure (AP) by acting on blood vessels and on cardiovascular centers in the brain. The mechanisms underlying the effects of estrogen in the brain stem, however, are not clear. The aim of the present study was to determine whether ovariectomy affects AP via the Rho/Rho-kinase pathway in the brain stem. We performed bilateral ovariectomy in 12-week-old female spontaneously hypertensive rats. AP and heart rate (HR), measured using radiotelemetry in awake rats, were increased in ovariectomized rats compared with control rats (mean AP: 163±3 versus 144±4 mm Hg; HR: 455±4 versus 380±6 bpm). Continuous intracisternal infusion of Y-27632 significantly attenuated the ovariectomy-induced increase in AP and HR (mean AP: 137±6 versus 163±3 mm Hg; HR: 379±10 versus 455±4 bpm). In addition, we confirmed the increase of Rho-kinase activity in the brain stem in ovariectomized rats, and the increase was attenuated by intracisternal infusion of Y-27632 via the phosphorylated ezrin, radixin, and moesin (ERM) family, which are Rho-kinase target proteins. Furthermore, angiotensin II type 1 receptor expression in the brain stem was significantly greater in ovariectomized rats than in control rats, and the increase was partially reduced by intracisternal infusion of Y-27632. In a separate group of animals, we confirmed that the serum and cerebrospinal fluid 17β-estradiol concentrations decreased in ovariectomized rats. These results suggest that depletion of endogenous estrogen by ovariectomy, at least in part, induces hypertension in female spontaneously hypertensive rats via activation of the renin–angiotensin system and the Rho/Rho-kinase pathway in the brain stem. (***Hypertension.* 2006;48:651-657.**)

Key Words: estrogen ■ brain ■ nervous system, sympathetic ■ receptors, angiotensin ■ blood pressure ■ heart rate

The incidence of cardiovascular diseases is lower in premenopausal women than in age-matched men[1–3] and postmenopausal women.[4] The decreased protective effect against cardiovascular diseases, such as hypertension, in postmenopausal women is thought to be because of endogenous ovarian estrogen depletion. Estrogen decreases arterial pressure through direct effects on blood vessels[5,6] and through effects on central cardiovascular regulatory centers by modulating autonomic function of the cardiovascular system.[7,8] Hormone replacement therapy in postmenopausal women favorably affects cardiovascular regulation by improving baroreflex function and heart rate (HR) variability (HRV)[8] and by decreasing sympathetic nerve activity.[9] In the central nervous system (CNS), endogenous estrogen has numerous effects through estrogen receptor–dependent and -independent pathways.[10,11] Estrogen and estrogen receptors are present in the brain stem where the vasomotor centers, such as the nucleus tractus solitarius (NTS) and the ventrolateral medulla, are located.[12] Medullary injections of exogenous estrogen decrease arterial pressure, HR, and renal sympathetic nerve activity and enhance reflex control of the HR in male rats, as well as in ovariectomized female rats,[13] suggesting that estrogen has beneficial effects on autonomic functions.[14]

Rho-Kinase is a serine–threonine protein kinase and is one of the effectors of the small GTP-binding protein Rho. This pathway is involved in various cellular functions including smooth muscle contraction, actin cytoskeleton organization, cell proliferation, and cell motility.[15–18] In vascular smooth muscle cells, activation of this pathway contributes to the pathophysiology of hypertension via smooth muscle contraction.[19,20] In the CNS, the Rho/Rho-kinase pathway contributes to the formation of dendritic spines.[21] Dendritic spines form the postsynaptic contact sites of excitatory synapses in the CNS[22] and are thought to be involved in synaptic transmission.[23] We reported previously that Rho-kinase in the brain stem modulates glutamate sensitivity[24] and maintains arterial pressure via the sympathetic nervous system and that activation of the Rho/Rho-kinase pathway might contribute to the central mechanisms of hypertension.[25,26] Estrogen also regulates the formation of excitatory synapses on dendritic spines.[27] Estrogen treatment increases spine number and

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synaptic density on apical and basal dendrites of CA1 pyramidal neurons in ovariectomized adult female rats.[28,29] These findings led to the hypothesis that the effects of endogenous estrogen on central cardiovascular regulation involve alterations in Rho-kinase activity in the central cardiovascular center. Therefore, the aim of the present study was to determine whether the depletion of endogenous estrogen affects arterial pressure via the Rho/Rho-kinase pathway in the brain stem.

For this purpose, we performed a bilateral ovariectomy in 12-week-old female spontaneously hypertensive rats (SHRs). Y-27632, a specific Rho-kinase inhibitor,[30] was then infused intracisternally for 2 weeks with a miniosmotic pump. Arterial pressure and HR were measured in awake rats using a radiotelemetry system.[31] In a separate group of animals, we confirmed that serum and cerebrospinal fluid (CSF) 17β-estradiol concentrations were decreased in ovariectomized rats. We then compared the Rho-kinase activity in the brain stem between control rats and ovariectomized rats. Finally, we compared the expression level of angiotensin II type 1 receptors (AT1Rs) in the brain stem between control rats and ovariectomized rats, because estrogen functions upstream of the renin–angiotensin system,[32,33] and the Rho/Rho-kinase pathway is downstream of the renin–angiotensin system.[34,35]

Methods

This study was reviewed and approved by the Committee on the Ethics of Animal Experiments, Kyushu University Graduate School of Medical Sciences, and was conducted according to the Guidelines for Animal Experiments of Kyushu University. Female SHRs (11-week-old, SLC Japan, Hamamatsu, Japan) were used in the present study. A flexible catheter containing an arterial pressure transmitter was introduced into the abdominal aorta just below the renal arteries under sodium pentobarbital (50 mg/kg IP) anesthesia. After surgery, rats were housed singly in cages and allowed unrestricted movement. Rats were fed food devoid of phytoestrogens (AIN 76A, Kyudo Co, Ltd). For 21 days after ovariectomy, arterial pressure and HR were recorded continuously for 10 minutes every other day using a UA-10 telemetry system (Data Sciences International).[25,31]

Surgical Procedures

Seven days after catheter implantation, we performed bilateral ovariectomy (OVX) or sham operation (control) in the then 12-week-old rats under sodium pentobarbital (50 mg/kg IP) anesthesia. Ovariectomized rats were further randomly divided into 2 groups. The first group received continuous infusion of vehicle (a-CSF, 0.25 μL/h; OVX-VEH rats), and the second group received continuous infusion of Y-27632, a specific Rho-kinase inhibitor (5 mmol/L Y-27632, 0.25 μL/h; OVX-Y rats), intracisternally for 2 weeks via a miniosmotic pump (Alzet model 1002; Durect Corp). The miniosmotic pump, filled with vehicle or Y-27632, was implanted subcutaneously in the back and connected to a polyethylene tube (PE 10). A small hole was then made in the atlantooccipital membrane, which covers the dorsal surface of the medulla, and the tip of the tube was placed intracisternally and fixed in place with tissue adhesive. After surgery, the rats were free to move about their cages. The infusion was calculated to last 14 days.

Power Spectral Analysis

To evaluate sympathetic activity, we performed power spectral analysis on day 11 after OVX or sham operation. HR was recorded from 20-minute ECGs performed in awake rats using the radiotelemetry system. The ECG data were calculated using the Powerlab system and Chart 4 software (AD Instruments), and the power spectra of the R–R interval was calculated using the maximum entropy method with MemCalc software (Suwa Trust Co, Ltd). The frequency bands were adapted for analysis in rats: a very low frequency band (VLF) of 0 to 0.25 Hz, a low frequency band (LF) of 0.25 to 0.8 Hz, and a high frequency band (HF) of 0.8 to 2.4 Hz.[36–38]

Western Blot Analysis

The animals of each group (control rats, OVX-VEH rats, and OVX-Y rats) were killed with an overdose of sodium pentobarbital on day 11 after OVX or sham operation, and whole brain stem tissues were obtained. The animals used in this experiment were different from those in which arterial pressure, HR, and ECG were monitored. The tissues were obtained as the whole brain stem to ensure that the same areas from each animal were used and then homogenized in lysing buffer containing 40 mmol/L HEPES, 1% Triton X-100, 10% glycerol, 1 mmol/L Na_3VO_4, and 1 mmol/L phenylmethylsulfonyl fluoride. The tissue lysate was centrifuged, and the supernatant was collected. The protein concentration was determined using a BCA protein assay kit (Pierce Chemical). A protein aliquot (15 μg) from each sample was separated on a 10% sodium dodecyl sulfate–polyacrylamide gel and subsequently transferred onto polyvinylidene difluoride membranes (Immobilon-P membrane, Millipore). Membranes were incubated with rabbit anti-phosphorylated ERM proteins [ezrin (Thr567), radixin (Thr564), and moesin (Thr558)], which are the target proteins of Rho-kinase (this primary antibody was made by Kaibuchi and colleagues[39–42] and has been characterized and used in many previous reports). Membranes were then incubated with a horseradish peroxidase-conjugated horse anti-rabbit IgG antibody (1:10 000). Immunoreactivity was detected by enhanced chemiluminescence autoradiography (ECL Western blotting detection kit, Amersham Pharmacia Biotechnology), and the film was analyzed using NIH Image (developed at the National Institutes of Health and available on the Internet at http://rsb.info.nih.gov/nih-image/). Western blot analysis for AT1R was performed as described above using rabbit anti-AT1R antibody (1:1000, Santa Cruz Biotechnology).

Measurement of Estradiol Concentration in the Serum and CSF

A separate set of 12-week-old female SHRs was divided into 2 groups. In the first group, bilateral OVX alone was performed (OVX rats). In the second group, sham operation was performed (Sham rats). The animals in each group were anesthetized with an overdose of sodium pentobarbital on day 11 after the intervention, and blood samples from the femoral vein were obtained to measure the serum 17β-estradiol concentration by radioimmunoassay performed by SRL Inc). Furthermore, a small hole was made in the atlantooccipital membrane, the tip of a tube connected to a syringe was placed intracisternally, and CSF was collected. Because only a small amount of CSF could be collected from each animal, samples from 5 animals of each group were pooled, and the 17β-estradiol concentration was measured.

Statistical Analysis

All of the values are expressed as mean±SEM. Two-way ANOVA was used to compare differences in mean arterial pressure (MAP) and HR between the Y-27632 and vehicle infusion groups. Comparisons between any mean values were performed by application of Bonferroni's correction for multiple comparisons. An unpaired *t* test was used to compare the baseline values and the effects of each intervention between groups. Differences were considered to be statistically significant when $P<0.05$.

Results

Effects of OVX and Intracisternal Infusion of Y-27632 on Arterial Pressure and HR Measured by Radiotelemetry

The time course of MAP and HR after OVX and intracisternal infusion of Y-27632 is shown in Figure 1. The baseline

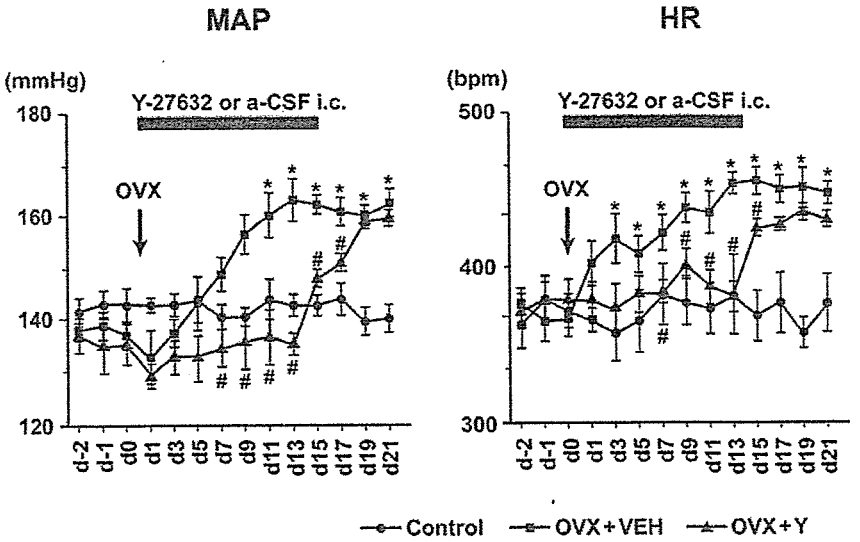

Figure 1. Chronic intracisternal infusion of Y-27632. Time course of MAP and HR after ovariectomy with continuous infusion of Y-27632 or vehicle intracisternally for 2 weeks with a miniosmotic pump. MAP and HR were increased in OVX-VEH rats. Y-27632 significantly attenuated the increase in MAP and HR in OVX-Y rats (n=4 for each). *$P<0.05$ vs control rats. #$P<0.05$ vs OVX+VEH rats.

values of arterial pressure in control rats, OVX-VEH rats, and OVX-Y rats were 192±7, 189±9, and 187±7 mm Hg (systolic arterial pressure); 115±3, 113±2, and 112±4 mm Hg (diastolic arterial pressure); and 142±3, 138±2, and 137±3 mm Hg (MAP), respectively. The baseline values of HR in control rats, OVX-VEH rats, and OVX-Y rats were 373±7, 372±6, and 362±7 bpm, respectively. Arterial pressure and HR were significantly increased in OVX-VEH rats. Y-27632 significantly attenuated the increase in arterial pressure and HR in OVX-Y rats. After discontinuing treatment with Y-27632, arterial pressure and HR increased to levels similar to those in OVX-VEH rats. In control rats, arterial pressure and HR did not change after the operation.

Effects of OVX and Intracisternal Infusion of Y-27632 on HR Power Spectra

The effects of OVX on HR power spectra are shown in Figure 2. The VLF and LF components were significantly increased in OVX-VEH rats. The VLF and LF components in OVX-Y rats were significantly reduced compared with those in OVX-VEH rats. The HF component, however, did not differ between control rats, OVX-VEH rats, and OVX-Y rats; therefore, the LF/HF ratio, which is considered to be a measure of sympathovagal balance, was also increased in OVX-VEH rats compared with the other groups.

Western Blot Analysis of Rho-Kinase Activity

The extent of ERM phosphorylation, which represents Rho-kinase activity, was greater in OVX-VEH rats and in OVX-Y rats than in control rats (Figure 3A). The increase in ERM phosphorylation in OVX-Y rats, however, was significantly less than that in OVX-VEH rats. The expression level of the total ERM family did not differ between groups.

AT1R Expression Level

The AT1R expression level was significantly increased in OVX-VEH rats and OVX-Y rats compared with control rats (Figure 3B). The increase in AT1R expression in OVX-Y rats was significantly smaller than that in OVX-VEH rats (Figure 3B).

Effects of OVX on Serum and CSF Estradiol Concentrations

On day 11 after the intervention, the serum 17β-estradiol concentration was decreased in OVX rats compared with Sham rats (139±51 pg/mL versus 34±11 pg/mL; n=5 for each; $P<0.05$). In addition, the CSF 17β-estradiol concentration was also decreased in OVX rats (270.0 pg/mL versus 61.5 pg/mL).

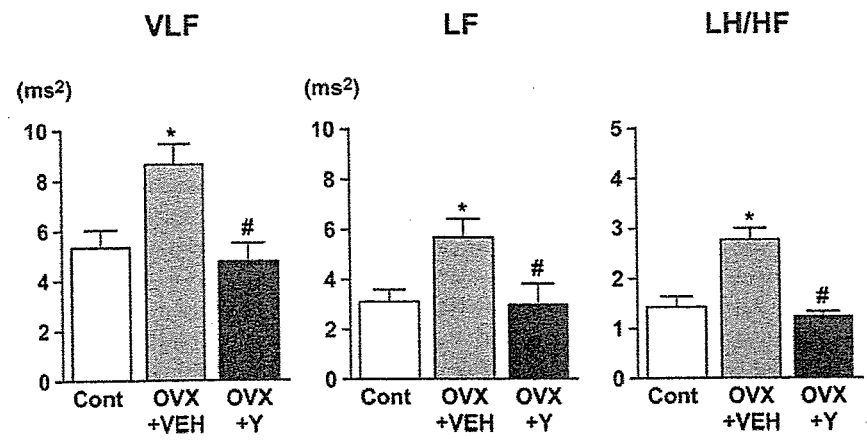

Figure 2. Evaluation of sympathetic nerve activity by power spectral analysis. Each graph shows the power density of each spectrum. The power density of VLF and LF reflects sympathetic nerve activity, and the ratio of LF/HF reflects the sympathovagal balance. Both parameters increased in OVX+VEH rats compared with control rats (n=5 for each). *$P<0.05$ vs control rats. #$P<0.05$ vs OVX+VEH rats.

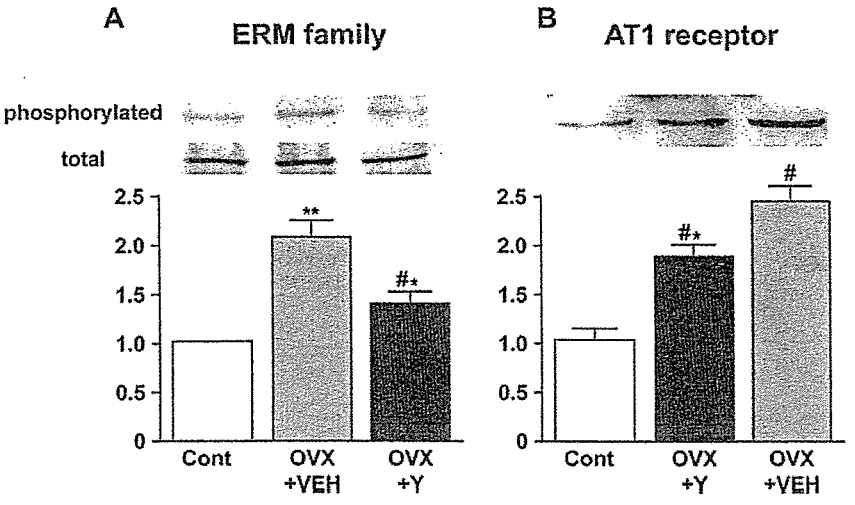

Figure 3. A, Western blot analysis demonstrating p-ERM expression in the brain stem. The p-ERM expression level was significantly higher in OVX-VEH rats than in control rats, and its increase was significantly attenuated by Y-27632 infusion. The data are expressed as the relative ratio to controls. The ratio of phosphorylated/total ERM family expression in control rats was assigned a value of 1 (n=5 for each). $^{**}P<0.01$, $^{*}P<0.05$ vs control rats. $\#P<0.05$ vs OVX+VEH rats. B, Western blot analysis demonstrating AT1R expression in the brain stem. The AT1R expression level was significantly higher in OVX-VEH rats than in control rats, and its increase was significantly attenuated by Y-27632 infusion. The data are expressed as relative ratio to control. The extent of the AT1R in control rats was assigned a value of 1 (n=4 for each). $^{**}P<0.01$, $^{*}P<0.05$ vs control rats. $\#P<0.05$ vs OVX+VEH rats.

Discussion

The present study demonstrated that in female SHRs, OVX induced an increase in arterial pressure via activation of the sympathetic nervous system, attributable, at least in part, to Rho-kinase activation in the brain stem. Furthermore, the results of the present study suggest that angiotensin II in the brain stem contributes to these mechanisms.

We used only female SHRs as a hypertensive model in this study. Previous studies focusing on the interaction between cardiovascular regulation and estrogen demonstrated that OVX has no effect on arterial pressure in normotensive animals.[43,44] Although the Rho-kinase activity and renin–angiotensin system in the brain stem might be affected by OVX in normotensive rats, the effects of OVX, like in postmenopausal women, are more obvious in hypertensive rats than in normotensive rats. Therefore, we used only SHRs in the present study. Previous studies also indicated that OVX alters arterial pressure only in salt-sensitive hypertensive models, such as SHRs and Dahl salt-sensitive rats.[45–47] There are reports that bilateral OVX in hypertensive rats enhances salt sensitivity but only increases arterial pressure in rats fed a high NaCl diet.[45,46] In the present study, arterial pressure and HR were significantly increased in OVX-VEH rats fed a standard NaCl diet compared with controls from day 7 after bilateral OVX. The discrepant results might be because of differences in age at the time of OVX or in the diet. In fact, rats ovariectomized at a young age and fed a normal diet only have a significant increase in arterial pressure when fed a high-NaCl diet,[45,46] whereas rats ovariectomized at an adult age and fed a phytoestrogen-devoid diet have increased arterial pressure under both a standard NaCl diet and a high-NaCl diet.[47] Furthermore, we measured arterial pressure by radiotelemetry. This system allowed us to measure arterial pressure in awake rats under relatively stress-free conditions compared with other methods, such as the tail-cuff method. In a preliminary study, when we measured arterial pressure by the tail-cuff method, OVX tended to increase arterial pressure, but the difference was not significant (data not shown). These methodologic differences might also cause discrepant results from previous studies. In fact, arterial pressure measured by radiotelemetry is significantly increased in ovariectomized Dahl salt-sensitive rats fed a phytoestrogen-free and low-salt diet.[48]

Intracisternal infusion of Y-27632 prevented the OVX-induced increase in arterial pressure, suggesting that endogenous Rho-kinase in the brain stem has an important role in OVX-induced hypertension in female SHRs. The concentration of Y-27632 (5 mmol/L) used in the present study has selective effects on the CNS.[26] Intracisternal drug infusion affects neurons in several regions of the autonomic cardiovascular area. We demonstrated previously, however, that the Rho/Rho-kinase pathway is activated in the NTS of SHRs but not normotensive rats, and Rho-kinase inhibition in the NTS induces a significant reduction in arterial pressure.[25] In addition, the NTS is positioned at the dorsal surface of the medulla. Therefore, Y-27632 might have greater effects on neurons of the NTS than on neurons of other nuclei, although we cannot exclude the possibility that it affects other autonomic areas in the brain stem. Together, these findings suggest that the effect of Y-27632 is mediated in part by the inhibition of Rho-kinase activity in the NTS.

In the present study, it is unlikely that the results are because of nonspecific effects caused by the surgical procedure, because continuous intracisternal infusion of vehicle using the same device did not suppress the increase in arterial pressure, and after discontinuing the Y-27632 infusion, arterial pressure increased to levels similar to those in OVX-VEH rats. In addition, ERM phosphorylation in the brain stem was significantly increased in OVX-VEH rats and reduced in OVX-Y rats, which strongly suggests that the Rho-kinase activity in the brain stem of OVX-VEH rats was increased and that Y-27632 suppressed Rho-kinase activity. We estimated the extent of ERM phosphorylation as a marker of Rho-kinase activity. The ERM family members are concentrated in the actin-rich cell surface, cross-link actin filaments with the plasma membrane, and contribute to cell–cell adhesion and maintenance of cell shape and cell motility.[49] Although the role of ERM family members in the central cardiovascular regulation is not clear, and ERM family members might be the substrates of other kinases, ERM phosphorylation is commonly used as an indicator of Rho-kinase activity.[25,26,34,40–42]

In the present study, intracisternal infusion of Y-27632 nearly abolished the OVX-induced increase in arterial pressure. We reported previously that Rho-kinase activity in the brain stem of hypertensive rats is increased compared with normotensive rats.[25,26] Therefore, intracisternal infusion of Y-27632 might suppress not only the OVX-induced increase in Rho-kinase activity but also basal Rho-kinase activity, thereby inducing a greater reduction in arterial pressure than that induced by OVX. In most previous studies, arterial pressure changes were evaluated several weeks after OVX. In the present study, ≈1 week after OVX, arterial pressure of ovariectomized rats was significantly increased compared with that of control rats. As mentioned above, we used more sensitive methods for measuring arterial pressure. In addition, we confirmed that the 17β-estradiol concentration, both in the serum and the CSF, decreased by ≈25% at 11 days after OVX. Although the serum and the CSF estradiol concentrations were markedly decreased by OVX, the concentrations were still high. We speculate that other organs, such as adrenal glands or adipocytes, produced estradiol after OVX. We did not address these issues, however, and do not have precise interpretations for this finding. In the present study, we primarily wanted to confirm that OVX reduced serum and CSF estradiol concentrations.

We estimated sympathetic activity using the HRV power spectral analysis. HRV is used as a noninvasive marker of autonomic outflow to the heart in a variety of disease states.[50] HRV has a very specific pattern in the frequency domains delineated by the HF, LF, and VLF components.[36] Spontaneous VLF power, which is 0 to 0.25 Hz, and LF power, which is 0.25 to 0.8 Hz, in the rat are particularly related to sympathetic nerve activity.[36–38] In fact, an increased LF component in R–R variability occurs in various conditions known to decrease baroreflex gain and increase sympathetic outflow, such as tilt, mental stress, and exercise.[51] On the other hand, the HF component is attributed to vagal and respiratory control, and the LF/HF ratio is used as an index of sympathovagal balance.[52] Therefore, we also used spectral analysis of the HR in the present study. The VLF and LF powers or the LF/HF ratio were greater in OVX-VEH rats than in control rats, and those in OVX-Y rats were significantly reduced compared with OVX-VEH rats. Furthermore, the HR increase could also be an indicator of activation of the sympathetic nervous system. In the present study, HR was greater in OVX-VEH rats than in control rats and that in OVX-Y rats was significantly reduced compared with OVX-VEH rats. These results indicate that sympathetic activity was increased by OVX, and intracisternal infusion of Y-27632 significantly attenuated the increase in sympathetic activity.

Estrogen decreases arterial pressure by acting on blood vessels or the kidney via the AT1R.[5,6,41] Furthermore, the renin–angiotensin system is also a major pathway of the central mechanisms of hypertension. Previous reports suggest that angiotensin II contributes to the neural mechanisms of hypertension.[53,54] In addition, inhibition of Rho-kinase activity suppresses angiotensin II–induced cardiovascular effects.[34] Therefore, we examined AT1R expression levels in each group to address the possibility of a partial interaction between the renin–angiotensin system and the Rho/Rho-kinase pathway in OVX-induced hypertension. The AT1R levels in the brain stem were significantly increased by OVX, and this increase was attenuated by intracisternal infusion of Y-27632. These results suggested that angiotensin II in the brain stem contributes to the mechanisms of OVX-induced hypertension in female SHRs. The Rho/Rho-kinase pathway is downstream of the renin–angiotensin system.[34,35] RhoA regulates the expression of AT1R,[55] however, and Y-27632 inhibits not only Rho-kinase activity, but also RhoA activity.[56] Therefore, Y-27632 might attenuate RhoA activity by direct effects or negative feedback mechanisms of the Rho/Rho-kinase pathway and, thus, lead to the inhibition of AT1R expression. The finding that Y-27632 had only a weak effect on AT1R expression, together with the results of previous studies,[34,35] suggest that the depletion of endogenous estrogen activates the Rho/Rho-kinase pathway in the brain stem and might also activate the renin–angiotensin system.

In conclusion, we demonstrated that the depletion of endogenous estrogen by OVX increases arterial pressure in female SHRs, at least in part, via activation of the renin–angiotensin system and Rho/Rho-kinase pathway in the brain stem.

Perspectives

It is not known how OVX induces increases in arterial pressure in female SHRs. In the CNS, both the Rho/Rho-kinase pathway and estrogen regulate the formation of excitatory synapses on dendritic spines.[27] Dendritic spines form the postsynaptic contact sites of excitatory synapses in the CNS[22] and are associated with glutamate sensitivity.[23] Therefore, estrogen depletion might induce morphological or functional changes in the dendritic spines via Rho-kinase activation. On the other hand, estrogen depletion increases arterial pressure and hypothalamic norepinephrine levels,[46] and hypothalamus neurons project to the dorsomedial medulla neurons, such as those in the NTS.[57] These findings suggest that estrogen depletion affects dorsomedial medulla neurons via changes in the hypothalamic norepinephrine levels. Although further studies are needed to clarify the mechanisms of these effects, the Rho/Rho-kinase pathway in the brain stem might be involved in the mechanisms underlying OVX-induced hypertension, because Rho-kinase in the NTS is involved in central mechanisms of cardiovascular regulation via modulation of the sensitivity of NTS neurons to glutamate.[24–26,58]

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Disclosures

None.

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Telmisartan downregulates angiotensin II type 1 receptor through activation of peroxisome proliferator-activated receptor γ

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Abstract

Objective: Telmisartan, an angiotensin II type 1 receptor (AT1R) antagonist, was found to have a unique property: it is a partial agonist of peroxisome proliferator-activated receptor gamma (PPARγ). Since previous studies have demonstrated that PPARγ activators suppressed AT1R expression, we examined whether telmisartan affects AT1R expression in vascular smooth muscle cells.

Methods: Vascular smooth muscle cells were derived from the thoracic aorta of Wistar–Kyoto rat. Northern and Western blotting analysis were used to examine AT1R mRNA and protein expression, respectively. The DEAE-dextran method was used for transfection, and the promoter activity of AT1R was examined by luciferase assay.

Results: Telmisartan decreased the expression of AT1R at the mRNA and protein levels in a dose- and time-dependent manner. Decreased AT1R promoter activity with unchanged mRNA stability suggested that telmisartan suppressed AT1R gene expression at the transcriptional level. However, the expression of AT1R was not suppressed by other AT1R antagonists such as candesartan or olmesartan. Since the suppression of AT1R expression was prevented by pretreatment with GW9662, a PPARγ antagonist, PPARγ should have participated in the process. The deletion and mutation analysis of the AT1R gene promoter indicated that a GC box located in the proximal promoter region is responsible for the telmisartan-induced downregulation.

Conclusion: Our data provides a novel insight into an effect of telmisartan: telmisartan inhibits AT1R gene expression through PPARγ activation. The dual inhibition of angiotensin II function by telmisartan – AT1R blockade and downregulation – would contribute to more complete inhibition of the renin–angiotensin system.

Keywords: Antihypertensive; Diuretic drugs; Atherosclerosis; Gene expression; Receptors; Renin–angiotensin system

1. Introduction

Angiontensin (Ang) II is a main final effector molecule of the renin–angiotensin system. Physiologically, Ang II plays an important role in controlling the blood pressure and the fluid volume [1]. However, Ang II is also involved in pathological conditions such as renal insufficiency [2], cardiovascular diseases [3] and metabolic disorders [4].

The effect of Ang II are mediated by Ang II receptors and so far two isoforms, type 1 receptor (AT1R) and type 2 receptor (AT2R), have been identified [5]. AT1R mediates most of the traditional effects of Ang II such as vasoconstriction, sodium retention, aldosterone secretion, and cell proliferation [1]. In contrast, AT2R mediates vasodilation and growth inhibition that opposes to the effects of AT1R [6]. However, it was reported that AT2R was hardly detected in blood vessel of adult animal [6].

Ang I converting enzyme inhibitors and AT1R antagonists are clinically used. Many clinical trials have demonstrated that these drugs are beneficial in the treatment of heart failure, renal failure and myocardial infarction. These drugs are also useful in preventing new-onset diabetes mellitus [7] and atrial fibrillation [8]. Telmisartan (Tel), one of the clinically

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available AT1R antagonists, was recently reported to have a partial agonistic effect on peroxisome proliferator activated receptor gamma (PPARγ) [9,10]. PPARγ is a nuclear receptor that regulates specific gene transcription [11]. The target genes of PPARγ are involved in the regulation of lipid and glucose metabolism [12], and inflammatory responses. Moreover, several studies have demonstrated that PPARγ activators are effective in preventing atherogenesis [13,14]. Therefore, Tel is focused for its additional therapeutic values in patients with metabolic disorders.

Previously, we and another group reported that activators of PPARγ such as 15-deoxy-$\Delta^{12,14}$-prostaglandin J_2 and pioglitazone (Pio) decreased the expression of AT1R in vascular smooth muscle cells (VSMCs) [15,16]. We, therefore, examined whether Tel, a partial agonist of PPARγ, affects the expression of AT1R in a similar way to PPARγ agonists in VSMCs.

2. Materials and methods

2.1. Materials

Tel and olmesartan (Olm) were generous gifts from Boehringer Ingelheim Co. and Sankyo Co., respectively. Candesartan (Can) and Pio were provided by Takeda Pharmaceutical Company. Dulbecco's modified Eagle's medium (DMEM) and fetal bovine serum (FBS) were purchased from GIBCO BRL. Bovine serum albumin (BSA) and Actinomycin D (ActD) were purchased from Sigma Chemical Co. Rabbit polyclonal antibody against AT1R [17,18] and α-tubulin were from Santa Cruz Biotechnology. Mouse polyclonal antibody against pERK and rabbit polyclonal antibody against ERK were from Cell Signaling Technology, Inc. Horseradish peroxidase-conjugated secondary antibodies (anti-rabbit IgG and anti-mouse IgG) were purchased from VECTOR Laboratories Inc. [α-^{32}P] dCTP was purchased from Perkin-Elmer Life Sciences.

2.2. Cell culture

All procedures and care of the animals were approved by the Committee on Ethics of Animal Experiments, Kyushu University and this study conforms with the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (NIH Publication No. 85-23, revised 1996). VSMCs were isolated from the thoracic aorta of Wistar–Kyoto rat by an explant method and maintained in DMEM supplemented with 10% FBS at 37 °C in a humidified atmosphere of 95% air −5% CO_2. VSMCs were cultured until grown to confluence, cultured in DMEM with 0.1% BSA for additional 2 days and used in the experiment. Cells between passages 4 and 14 were used.

2.3. Northern blotting

Total RNA was prepared by acid guanidinium thiocyanate–phenol–chloroform extraction method [19]. Total RNA was electrophoresed on a 1.0% agarose, 1.0% formaldehyde gel, transferred to Hybond-N+ membrane (Amersham Biosciences) by a capillary transfer method in 10×SSC (1×SSC is 150 mmol/L of sodium and 15 mmol/L of sodium citrate) buffer overnight. The membrane was cross-linked by a UV cross-linker (Funakoshi Corporation). Prehybridization and hybridization were performed in a buffer containing 50% formamide, 5×SSC, 80 mmol/L sodium phosphate (pH 7.5), 2×Denhardt's Solution, 1% SDS, and 100 μg/L of heat-denatured salmon sperm DNA for 1 h and 16 h, respectively, at 42 °C. An ECORI fragment of the third exon of rat AT1A gene [20] and ribosomal RNA were labeled with ^{32}P by a Random Primer DNA Labeling Kit Ver.2 (Takara Bio Inc.) and used as a probe after heat denaturation. The hybridized membrane was washed twice with 2×SSC at room temperature, followed by two washes with 2×SSC/1% SDS for 30 min at 55 °C. The membrane was then exposed to a KODAK BioMax XAR Film at −80 °C. The hybridized membrane was stripped by boiling in 0.5% SDS solution and hybridized to a ^{32}P-labeled ribosomal RNA probe to obtain reference for the amount of applied RNA. Autoradiography was scanned and analyzed by a MacBAS Bioimage Analyzer (Fuji Photo Film Co). To analyze mRNA stability of AT1R, Actinomycin (Act) D (5 μg/mL) was added after 6 h of stimulation with Tel (10 μmol/L). In a control experiment, only ActD was added. Cells were harvested after 3, 6, 12, and 24 h of addition of ActD and expression level of AT1R mRNA was examined by Northern blot analysis.

2.4. Measurement of AT1R gene promoter activity

Five deletion mutants of AT_{1A} gene promoter were prepared by digestion with restriction endonucleases and

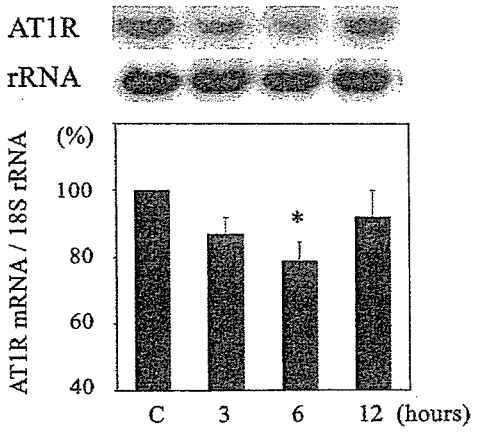

Fig. 1. Telmisartan (Tel) suppressed AT1R mRNA expression in VSMCs. VSMCs were incubated with Tel (10 μmol/L) for various periods as indicated in the figure. Total RNA was isolated and expression of AT1R mRNA and 18S rRNA (rRNA) was determined by Northern blot analysis. Radioactivity of AT1R mRNA was measured with an imaging analyzer and was normalized by radioactivity of rRNA. Values (mean±S.E.M.) are expressed as a percent of control culture in the bar graph (100%) (n=5). *P<0.05 vs control (c).

ligated to luciferase gene [21]. Confluent VSMCs were split by trypsin/EDTA solution and cells were prepared in a 6 cm tissue culture dish. At 80% confluence, 5 μg of AT1 promoter-luciferase fusion DNA and 2 μg of β-galactosidase gene were introduced to VSMC by the DEAE-dextran method according to the manufacturer's instruction (Promega Corporation). The cells were cultured in DMEM with 10% FBS for 18 h, washed twice with phosphate buffered saline, cultured in DMEM with 0.1% BSA for 24 h and stimulated with Tel (10 μmol/L) for 12 h. Then, the cells were lysed in 200 μL of Reporter lysis buffer (Promega Corporation). 100 μL of lysate was used for luciferase activity assay in a Lumat luminometer (LB 9501, Berthold, Germany). The β-galactosidase activity in the same sample was measured spectrophotometrically according to Sambrook et al. [22] and used to normalize the luciferase activity.

The AT1R promoter-luciferase construct with mutation in the GC-box-related sequence (wild type: TGCAGAGCAGC GACGCCCCCTAGGC mutant: TGCAGAGCAGCGA CGTTTCCTAGGC) was a generous gift from Dr. Sugawara (Tohoku University) [16].

2.5. Western blot analysis

VSMCs were lysed in a lysis buffer containing RIPA (100 mM sodium, 60 mM Na_2HPO_4, 100 mM NaF 10 mM EDTA, and 20 mM Tris), 1% aprotinin, 0.5% pepstatin A, 1 mmol/L PMSF, and 0.05% leupeptin. Protein concentrations were determined with the bicinchoninic acid protein assay kit (Pierce Chemical Co). Cell lysates were heated in a sample buffer (62.5 mmol/L Tris–HCl [pH 6.8], 10% glycerol, 2% SDS, 0.05% bromophenolblue, and

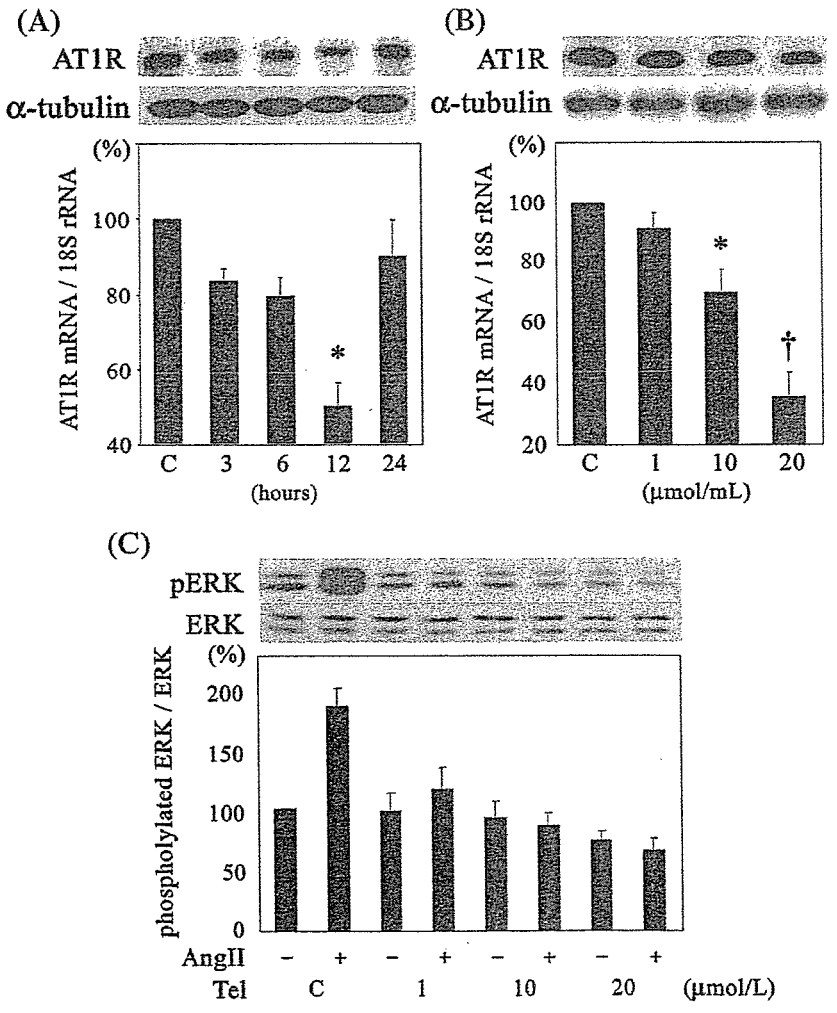

Fig. 2. Suppression of AT1R protein by telmisartan (Tel) in VSMCs. (A) VSMCs were incubated with Tel (10 μmol/L) for various periods as indicated in the figure. (B) VSMCs were incubated with Tel at concentrations varying from 1 to 20 μmol/L for 12 h. Expression of AT1R protein and α-tubulin was detected by Western blot analysis. The density of the specific band was scanned and quantified with an imaging analyzer. The ratio of AT1R to α-tubulin is shown in the bar graph. (C) VSMCs were incubated with Tel at various concentrations as indicated in the figure and stimulated by Ang II (100 μmol/L). Expressions of pERK and ERK protein were detected by Western blot analysis. The density of the specific band was scanned and quantified with an imaging analyzer. The ratio of pERK to ERK is shown in the bar graph. Values (mean±S.E.M.) are expressed as a percent of control (c) culture (100%) ($n=5$). *$P<0.05$ vs control. †$P<0.01$ vs control.

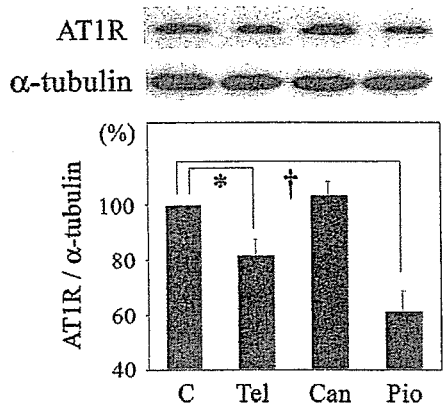

Fig. 3. Suppression of AT1R protein by telmisartan (Tel) but not by candesartan (Can) in VSMCs. VSMCs were incubated with Tel (10 μmol/L, 12 h), Can (10 μmol/L, 12 h), and Pio (10 μmol/L, 6 h). Expression of AT1R protein and α-tubulin was determined by Western blot analysis. Densitometric analysis was performed as described in the legend to Fig. 2. Values (mean±S.E.M.) are expressed as a percent of control (c) culture (100%) (n=5). *P<0.05 vs control. †P<0.01 vs control.

715 mmol/L 2-mercaptoethanol) at 95 °C for 3 min, electrophoresed on 12% SDS-polyacrylamide gel, and transferred to a polyvinylidene difluoride membrane (Immobilon-P, Millipore). The blots were blocked with TBS-T (20 mmol/L Tris–HCl [pH 7.6], 137 mmol/L NaCl, 0.1% Tween 20) containing 5% skim milk at room temperature for 30 min. The AT1R protein expressions were detected by ECL chemiluminescence (Amersham Pharmacia Biotech) according to the manufacturer's instructions. The membranes were exposed to X-ray film. The membranes were stripped by incubating them in a buffer containing 62.5 mmol/L Tris–HCl, 2% SDS, and 100 mmol/L 2-mercaptoethanol at 50 °C for 30 min and reprobed with an antibody against α-tubulin by the same procedure. Phospholylated ERK and ERK (which recognizes both phosphorylated and nonphosphorylated forms) were examined by the same method.

2.6. Statistical analysis

Statistical analysis was performed with 1- or 2-way ANOVA and Fisher test, if appropriate. Statistical significance was designated as P<0.05. Values are expressed as mean±S.E.M.

3. Results

3.1. Tel reduced the expression of AT1R

VMSCs were incubated with Tel (10 μmol/L) for various periods. The expression level of AT1R mRNA was gradually decreased with a peak suppression at 6 h of incubation (Fig. 1). Though we did not remove Tel from the medium, the expression of AT1R demonstrated a transient suppression. The mechanism of its recovery at 12 h of incubation is unknown. However, there are some reports that demonstrated a transient or a biphasic gene expression induced by thiazolidinediones (TZDs) [23,24], which seems to be consistent with our results. Western blot analysis revealed that Tel reduced AT1R protein level with a peak reduction at 12 h of incubation (Fig. 2A), and that Tel suppressed AT1R expression in a dose-dependent manner (Fig. 2B). As shown in Fig. 2C, preincubation with Tel at the same concentration as used in Fig. 2B almost completely inhibited the Ang II-induced ERK phosphorylation.

Following experiment used 10 μmol/L of Tel, which is the minimal dose that suppressed AT1R expression. Pio, one

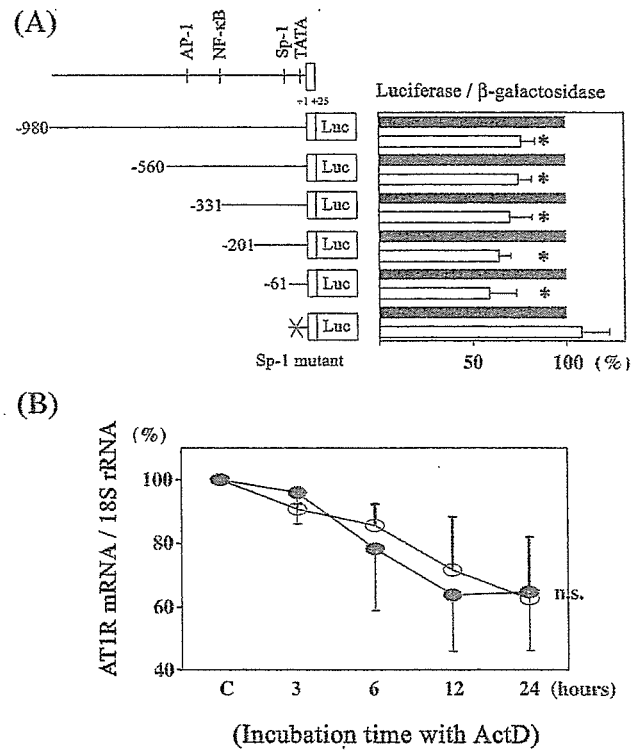

Fig. 4. Effect of telmisartan (Tel) on AT1R gene promoter activity and AT1R mRNA stability. (A) The scheme of deletion mutants of AT1R promoter/luciferase fusion DNA construct and SP-1 mutant construct is indicated. These luciferase constructs were introduced to VSMCs with LacZ expression plasmid by the DEAE-dextran method. Then VSMCs were stimulated with Tel (10 μmol/L) for 12 h. Relative luciferase activity of unstimulated VSMCs (control) was set as 100%. Solid and open bars indicate the relative luciferase activity of unstimulated and Tel-stimulated VSMCs transfected with the same construct indicated in the left panel, respectively. Values (mean±S.E.M.) are expressed as a percent of control culture (n=6). *P<0.05 vs unstimulated cells. n.s. not siginificant. (B) VSMCs were incubated with Tel (10 μmol/L) for 6 h and then ActD (5 μg/mL) was added. In a control experiment, only ActD was added to the medium. Total RNA was isolated at the indicated time after ActD supplementation and expression levels of AT1R mRNA and rRNA were determined with method described in the legend to Fig. 1. Expression level of AT1R mRNA was normalized with that of rRNA. The normalized AT1R mRNA expression before addition of ActD in each group was set as 100% (c), (n=3). n.s. not siginificant.

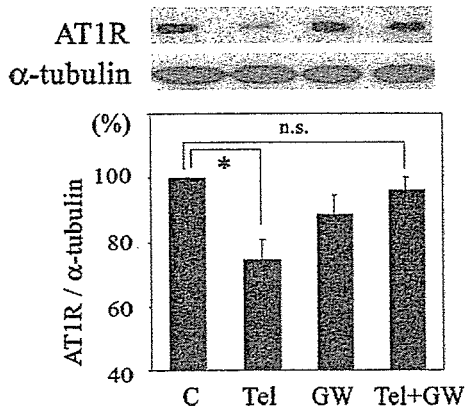

Fig. 5. The effect of GW9662, a PPAR γ antagonist, on telmisartan (Tel)-induced AT1R downregulation. VSMCs were preincubated with GW9662 (5 μg/mL) for 30 min and incubated with or without Tel (10 μmol/L) for 12 h. Expression of AT1R protein and α-tubulin was determined by Western blot analysis. Densitometric analysis was performed as described in the legend to Fig. 2. Values (mean±S.E.M.) are expressed as a percent of control culture (100%) ($n=5$). *$P<0.05$ vs control. n.s. not significant.

of TZDs and a full PPARγ agonist, inhibited AT1R expression as previously described [16]. We examined the effect of other AT1R antagonists on AT1R expression. Can (Fig. 3) and Olm (data not shown) had no effect on AT1R expression.

3.2. Tel inhibits AT1R expression at the transcriptional level

Deletion mutants of AT1 promoter/luciferase fusion DNA were used to locate the response element responsible for Tel-induced AT1R suppression (Fig. 4A). The suppression was observed in all constructs from −980/+25-luc to −61/+25-luc, so we supposed that the response element may exist in the DNA segment between −61 bp and +25 bp. Since Sugawara et al. [16] had previously reported the crucial role of a GC-box-related sequence within the −58/−34 region of the AT1R gene promoter in PPARγ-induced AT1R suppression, we hypothesized that the same site may also be important in Tel-induced suppression. The luciferase construct with mutation in GC box (Sp1 site) failed to respond to Tel (Fig. 4A), indicating the important role of Sp1 site in Tel-induced downregulation. In addition, Tel did not affect the degradation rate of AT1R mRNA (Fig. 4B). These data suggested that Tel inhibits AT1R gene transcription and does not affect AT1R mRNA stability.

3.3. Tel-induced AT1R downregulation is PPARγ dependent

To examine the role of PPARγ in Tel-induced AT1R suppression, we examined the effect of GW9662, a PPARγ antagonist. Although GW9662 itself did not affect AT1R expression, preincubation with GW9662 blocked AT1R suppression induced by Tel (Fig. 5).

4. Discussion

In the present study, we demonstrated that Tel, an AT1R antagonist, suppressed the AT1R expression through the PPARγ-mediated pathway. This is the first study that demonstrates the suppression of AT1R expression by AT1R antagonist in VSMCs.

The expression of AT1R was suppressed at both mRNA and protein levels. The results of the promoter assay and the mRNA stability assay suggested that the suppression occurred at the transcriptional level rather than the post-transcriptional level.

The involvement of PPARγ on AT1R suppression was confirmed by experiments using GW9662. GW9662 prevented the Tel-induced suppression of ATR expression, indicating the critical role of PPARγ in this pathway. The ability to activate PPARγ is reported to be unique to Tel and irbesartan (Irb) among several AT1R antagonists [10]. In our study, though we have not examined the effect of Irb, Can (Fig. 3) and Olm (data not shown) had no effect on AT1R expression. Our results seem to be consistent with the previous report [10].

Schupp et al. reported that a subset of AT1R blockers (ARB), Tel and Irb, induced PPARγ activity and promoted PPARγ-dependent differentiation in 3T3-L1 adipocytes [9]. These ARB activated PPARγ by direct interaction with the ligand binding domain (LBD) of PPARγ [9].

The docking studies of the molecular binding model explained the difference in the ability to activate PPARγ among several ARB and full agonists of PPARγ by comparing their interaction with residues of several helixes [10]. According to this model, Tel fits in the LBD of PPARγ surrounded by helixes H3, H6, and H7. However, Tel does not interact with activation function-2 helix that is responsible for receptor activation and stabilization by full PPARγ agonists. Irb, Can, and Olm, so-called tetrazole-containing ARBs, made contact with helix H3 but not H7. The difference in the interaction with these helixes of LBD of PPARγ might contribute to their potential to activate PPARγ.

On activation by a ligand, PPARγ regulates the expression of several genes involved in lipid and carbohydrate metabolism [25,26] and inflammatory responses [27]. From the molecular insight, these effects are basically due to two different transcriptional regulatory mechanisms, one is transactivation and the other is transrepression. Transactivation is dependent on PPARγ response element (PPRE). Upon activation, PPARγ forms a heterodimer with retinoid X receptor and binds to PPRE in the promoter region of the target genes such as CD36 and glucose transporter 4. In contrast, transrepression involves interference with other transcription factors such as NF-κB and AP-1. Because there is no consensus sequence of PPRE in the AT1R gene promoter up to −980 bp, the suppression of AT1R gene transcription should have occurred through the latter mechanism.

This assumption is well substantiated by a previous study which reported that suppression of AT1R gene expression by

PPARγ activation was independent of PPRE [16]. The authors demonstrated that the −58/−34 region of the AT1R gene promoter, including Sp1 binding site, was essential to the PPARγ activator-induced AT1R suppression. They concluded that activated PPARγ inhibited Sp1 function by direct protein–protein interaction. Our study with Tel also demonstrated the essential role of Sp1 binding site in the suppression of AT1R expression. Therefore, Tel may inhibit Sp1 function through the activation of PPARγ resulting in downregulation of the AT1R expression.

Our previous studies showed that downregulation of AT1R by PPARγ activator attenuated cellular response to Ang II [15]. However, Ang II induced ERK activation was almost completely blocked by Tel even at 1 μmol/L, which did not affect AT1R expression because of AT1R blocking effect (Fig. 2C). Therefore, AT1R binding effect is expected at the lower concentration of Tel and dual effect of AT1R binding and AT1R downregulation is expected at higher concentration.

TZDs, synthetic PPARγ activators, are reported to inhibit atherogenesis by regulating various gene expressions. Several studies have demonstrated the anti-atherogenic effects of PPARγ activators in both animal models and human. Rosiglitazone (Rosi), one of TZDs, was shown to have additive effects on plaque regression in the combination treatment with simvastatin in an atherosclerotic rabbit model [28]. Anti-atherogenic effect of Rosi was also reported in a diabetes–atherosclerosis mouse model [29]. AT1R antagonists are reported to suppress atherogenesis. Strawn et al. demonstrated that losartan attenuated atherogenesis in monkeys with hypercholesterolemia [30]. Based on these studies, Tel may be more efficient in suppressing atherosclerotic vascular diseases due to its properties of PPARγ activation and AT1R antagonism.

TZDs are also effective in improving insulin sensitivity and they are already utilized to treat patients with type 2 diabetes [31]. Although the precise mechanism for PPARγ-mediated insulin sensitization is not clear, TZDs reduced fasting and postprandial glucose concentration, and insulin level. Ang II also affects insulin sensitivity [32]. The inhibition of AT1R by losartan improved insulin sensitivity [33] and, in the LIFE study, losartan significantly reduced the incidence of new-onset diabetes in patients with hypertension compared with atenolol [34]. Therefore, the dual effects of Tel may synergistically improve insulin sensitivity.

The dual function of Tel, an AT1R antagonist and a partial agonist of PPARγ, may be quite useful for the treatment of patients with hypertension with complications such as diabetes and atherosclerosis. Considering the blockade of Ang II, the downregulation of AT1R through the activation of PPARγ adds further possibility of Tel. This may result in more complete inhibition of the Ang II. However, it remains to be determined whether Tel suppresses the expression of AT1R in vivo and is superior to other AT1R antagonists in terms of the inhibition of the progression of cardiovascular diseases. Further study is needed.

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A third-generation, long-acting, dihydropyridine calcium antagonist, azelnidipine, attenuates stent-associated neointimal formation in non-human primates

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Background Calcium antagonists have been shown to reduce atherogenesis and improve clinical outcomes in atherosclerotic vascular disease. No study has so far, however, addressed the effects of calcium antagonists on stent-associated neointimal formation. We therefore investigated whether a third-generation calcium antagonist, azelnidipine, attenuates in-stent neointimal formation in non-human primates.

Method Male cynomolgus monkeys were fed a high cholesterol diet for 4 weeks, and were randomly assigned to three groups: a vehicle group and two other groups treated with azelnidipine at 3 and 10 mg/kg per day for an additional 24 weeks ($n = 12$ each). Multi-link stents were then implanted in the iliac artery.

Results Azelnidipine at the high dose reduced neointimal thickness (0.25 ± 0.02 versus 0.19 ± 0.02 mm; $P < 0.05$). Azelnidipine also reduced local oxidative stress and monocyte chemoattractant protein 1 (MCP-1) expression. No difference was found between the three groups in the degrees of injury score, inflammation score, plaque neovascularization, or plasma lipid levels. Azelnidipine also reduced MCP-1-induced proliferation/migration of vascular smooth muscle cells *in vitro*.

Conclusions This study demonstrated for the first time that azelnidipine attenuates in-stent neointimal formation associated with the reduced expression of MCP-1 and smooth muscle proliferation/migration in the neointima. These data in non-human primates suggest potential clinical benefits of azelnidipine as a 'vasculoprotective calcium antagonist' in patients undergoing vascular interventions. *J Hypertens* 24:1881–1889 © 2006 Lippincott Williams & Wilkins.

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Introduction

Dihydropyridine calcium antagonists have been used worldwide for patients with hypertension and atherosclerotic vascular disease. Recent clinical trials with calcium antagonists provided evidence suggesting that reducing arterial blood pressure close to normal ranges is of great importance in reducing cardiovascular events [1–4]. Notably, in the Comparison of Amlodipine versus Enalapril to Limit Occurences of Thrombosis (CAMELOT) trial [5], compared with placebo and enalapril, the third-generation calcium antagonist amlodipine was shown to reduce hospitalization as a result of unstable angina and revascularization in patients with coronary artery disease already being treated with aspirin, beta-blockers, or statins. These latest clinical trials suggest that there might be pleiotropic actions of calcium antagonists beyond blood pressure lowering. The pleiotropic actions of calcium antagonists may be distinct from pharmacological actions related to L-type calcium channel blockade, but may be attributable to their lipophilic character leading to a high affinity with membrane phospholipid of arterial wall cells such as vascular smooth muscle cells (VSMC) [6]. The vasculoprotective actions of such calcium antagonists include an improvement in endothelial function, an anti-inflammation effect, anti-oxidant function, antiproliferation of VSMC and so forth.

In animal experiments, calcium antagonists have been shown to prevent or attenuate atherosclerosis [7,8], hypertension-induced vascular remodeling [9], heart failure [10] and neointimal formation after balloon injury [11]. However, no study has so far addressed the effects of

calcium antagonists on stent-associated neointimal formation. Investigating this subject must be of clinical importance, because stent implantation is now the major revascularization technique worldwide in patients with atherosclerotic vascular disease. Although a drug-eluting stent reduces the restenosis rate in selected arterial lesions, stent-associated restenosis remains an unsolved clinical issue in high-risk lesions. The cellular mechanism of restenosis after stenting is presumed to be neointimal formation as a result of VSMC proliferation/migration, recruitment of bone marrow-derived progenitor cells or inflammation in response to injury [12–14].

Azelnidipine is a newly developed third-generation calcium antagonist and has antihypertensive action that is comparable to amlodipine [15]. Azelnidipine has strong lipophilicity and affinity to membranes of VSMC [16,17]. Therefore, the aim of this study was to investigate whether azelnidipine at a clinical dose range attenuates in-stent neointimal formation in non-human primates (cynomolgus monkeys). To gain clinical significance for the results obtained, we used a non-human primate model of stent-associated neointimal formation [12].

Materials and methods

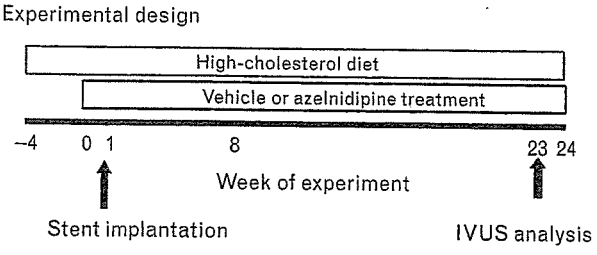
Experimental animals

Thirty-six 5-year-old male cynomolgus monkeys weighing 4.2–6.6 kg were purchased. The study protocol was reviewed and approved by the Committee on the Ethics of Animal Experiments, Kyushu University Graduate School of Medical Sciences. The animal care before and after the operation of the animal model was performed in Gaoyao Kangda Laboratory Animal Science and Technology in China. A part of this study was performed at the Station for Collaborative Research and the Morphology Core, Kyushu University Graduate School of Medical Sciences.

Animal model of in-stent restenosis

All animals were fed a high-cholesterol diet (0.5% cholesterol and 6% corn oil) for 4 weeks before stent implantation (Fig. 1). Ticlopidine 100 mg and aspirin 81 mg were started 7 days before the initial procedure. Aspirin was to be given and continued until animals were killed at 6 months, and ticlopidine was administered for 28 days. The animals were randomly assigned to three groups as follows: (i) no treatment vehicle control group (0.5% carboxymethyl cellulose sodium salt); (ii) low-dose azeenidipine (3 mg/kg per day; donated by Sankyo Pharmaceutical Co., Tokyo, Japan) group; (iii) high-dose azelnidipine (10 mg/kg per day) group ($n = 12$ each). Azelnidipine treatment was carried out once a day by gavage for 24 weeks. One week after grouping, all monkeys were anaesthetized with ketamine hydrochloride (10 mg/kg intramuscularly) and sodium pentobarbital (30 mg/kg intravenously), and underwent the placement of a 15 mm-long Multilink stent mounted over the 3-mm balloon implanted in the iliac artery (30 s inflation at 6 atm followed by 60 s inflation at 8 atm, resulting in a stent-to-artery ratio of 1.2 : 1.0) as described [12]. After the operation, all monkeys were fed the same high-cholesterol diet.

Fig. 1



Experimental design. IVUS, Intravascular ultrasound.

Histopathology and immunohistochemistry

Stented arterial sections were excised and fixed for 24 h with 95% ethanol and 1% acetic acid. Each segment was divided into two parts at the centre of the stent. The proximal part was embedded in methyl methacrylate mixed with *n*-butyl methacrylate to allow for sectioning through metal stent struts. Serial sections were stained with elastica van Gieson and haematoxylin eosin. To evaluate the stent area, lumen area, medial area, and neointimal area, the neointimal thickening (neointimal area/length of internal elastica lamina; IEL) were measured. An immunohistochemical study was performed with antibodies against human smooth muscle actin (1A4; Dako, Tokyo, Japan) and von Willebrand factor (vWF; Dako). To evaluate the degree of neovascularization the number of arterioles in the intima and adventitia were counted.

The distal part was used for immunohistochemical analysis. After stent struts were gently removed with micro forceps, the tissue was dehydrated, embedded in paraffin, and cut into 5-μm thick slices. They were subjected to immunostaining using antibodies against human monocyte chemoattractant protein 1 (MCP-1; Santa Cruz, California, USA) and C-C chemokine receptor 2 (CCR2, Santa Cruz) as described [12]. A single observer, who was blinded to the experiment protocol, performed the morphometry. All images were captured by an Olympus microscope equipped with a digital camera (HC-2500) and were analysed using Adobe Photoshop 6.0 (Adobe Systems, San Jose, California, USA) and Scion Image 1.62 (Scion, Frederick, Maryland, USA) software.

Local oxidative stress with fluorescent dihydroethidium micrographic analysis

Frozen, enzymatically intact, sections (10-μm thick) just at the proximal edge of stented iliac arteries were incubated with dihydroethidium (10 μmol/l; Sigma, Tokyo,