Reduced Connexin-43 Expression in the Aorta of Prehypertensive Rats

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Summary
Genetic component represents an important factor in the development of hypertension, which is known to be associated with changes in expression of vascular gap junction protein connexin 43 (Cx43). The aim of the study was to examine the distribution and expression of Cx43 in the aortic endothelium of adult normotensive Wistar rats (W), borderline hypertensive rats (BHR) and spontaneously hypertensive rats (SHR). Rings of the thoracic aorta were processed for immunofluorescence and Western blot analysis of endothelial Cx43 and for electron microscopy. Both, BHR and SHR exhibited significantly increased blood pressure vs. W (132 ± 2 mm Hg and 185 ± 3 mm Hg vs. 110 ± 2 mm Hg). Reduced Cx43 immunofluorescence was observed in the endothelium of BHR and these alterations were more pronounced in SHR. Western blot analysis showed significant suppression of Cx43 expression in the aorta of both BHR (p<0.05) and SHR (p<0.001) vs. W. Electron microscopy revealed local subcellular alterations of interendothelial connections in BHR including extended tight junctions. These alterations were more frequent and marked in SHR. The results indicate that connexin 43 expression is reduced in the aortic endothelium already in prehypertensive period, which may affect cell-to-cell communication and thus participate in acceleration of hypertensive disease.

Key words
Connexin 43 • Interendothelial connections • Aorta • BHR • SHR

Introduction
The vascular endothelium is a continuous monolayer formed by cells connected by various types of intercellular junctions that are involved in the control of endothelial permeability (Dejana et al. 1995). The structural and functional integrity of endothelium, which is promoted via the complex interaction of many substances, is essential for maintenance of tissue homeostasis as well as for regulation of vascular function. Such a versatile role of the endothelium requires coordination of the activity among individual cells, including gap junctional intercellular communication. Gap junctions are cell membrane protein channels clustered at cell-cell connections allowing exchange of ions and small molecules (up to 1 kDa) and thus direct intercellular communication. The properties of gap junctional channels are determined by the connexins, proteins that belong to a multigene family (Yeh et al. 2000). Disturbances in the endothelial integrity may affect cell-to-cell communication and thus signal transmission along the vessel wall (Payne et al. 2004).

Gap junctions have shown to play a significant role in many physiological mechanisms. Coupling of vascular endothelial cells as well as smooth muscle cells is thought to facilitate conduction of vasomotor responses along arteries by allowing for cell-to-cell transfer of electrical signals (Rummery and Hill 2004). Additionally, gap junctions are implicated in regulation of vascular tone (Christ et al. 1996, Chaytor et al. 1998) as well as in

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the control of vascular cell proliferation and migration (Simon and McWhorter 2003). In this regard, gap junctions may have important role in the development of the vasculature and in responses to blood vessel damage. Despite significant knowledge of connexin expression patterns in blood vessels, the contribution of specific connexins to endothelial communication is still not well understood (Krüger et al. 2002).

In the aorta, a large conduit vessel unlikely to be responsible for changes in peripheral resistance, the expression of three connexin isotypes Cx37, Cx40 and Cx43 was demonstrated (Van Kempen and Jongsma 1999, Haefliger et al. 2004, Rummery and Hill 2004). However, both the decrease and increase in Cx43 expression were reported in the aorta, depending on the experimental model of hypertension used (Watts and Webb 1996, Haefliger et al. 2004). Because the distensibility and compliance of the large artery differ among the individual models of hypertension, it was concluded that alterations in Cx43 expression represent an adaptive response to the changes occurring in the vascular wall during hypertension, possibly to maintain vascular elasticity (Haefliger et al. 2004).

Genetic predisposition to high blood pressure is a known risk factor for the development of hypertension and other cardiovascular diseases. Thus, normotensive animal models due to their genetic consistency need not be always appropriate for investigation of cardiovascular diseases. For such studies, a more suitable model is that of rats with a family history of hypertension, produced by the matting of spontaneously hypertensive dams with normotensive sires (Lawler et al. 1981, Bernátová et al. 2007). All procedures used were approved by the State Veterinary and Food Administration of the Slovak Republic.

For morphological analysis, rats were anesthetized by thiopental, the thoracic aorta was excised, cleaned of adherent tissue and processed for immunofluorescence, Western blot and electron microscopy.

**Basic parameters**

Blood pressure (BP), heart rate (HR) (measured non-invasively by tail-cuff plethysmography using the Statham Pressure Transducer P23XL, Hugo Sachs, Germany) and body weight (BW) were determined at the end of experiment.

**Immunofluorescence**

For Cx43 immunolabeling, series of cryostat cross sections of non-fixed frozen aortic tissue (10 μm) were preincubated with phosphate-buffered saline (PBS), fixed with 4 % paraformaldehyde for 15 min and membranes of aortic cells were permeabilized with 0.3 % Triton X-100 for 5 min. Afterwards, sections were blocked with PBS buffer containing 1 % goat serum for 30 min and incubated with primary monoclonal antibody mouse anti-Cx43 (1:100, Chemicon) for 2 h at room temperature. Sections were subsequently rinsed with PBS, followed by application of secondary antibody goat anti mouse IgG conjugated with FITC-fluorescein (1:50, Chemicon). Primary antibody was omitted in negative controls. After washing aortic sections were rinsed with PBS, mounted into Vectashield mounting medium (Baria, Germany) and viewed by Axiostar fluorescent microscope (Carl Zeiss). Pictures were digitized and transferred into PC.
Fig. 1. Immunodetection of Cx43 in endothelium and media of thoracic aorta of normotensive Wistar rats (A), borderline hypertensive rats (B), spontaneously hypertensive rats (C). E – endothelium, L – Lumen. Original magnification 80 x.

Western blot analysis

Frozen aortic tissue was homogenized in 10-fold amount of SB20 solution (20 % SDS, 10 mmol/l EDTA, 0.1 mol/l TRIS, pH 6.8) by sonificator UP 100H (Dr. Hielscher, Germany). Total protein content was determined by spectrophotometric method according to Bradford. Samples were fractionated by electrophoresis in 12.5 % polyacrylamide gel and blotted during night onto Trans Blot Transfer Medium (Bio Rad) at a constant current 50 mA. Membranes were preincubated for one hour at room temperature in 4 % solution of dry milk in TRIS-buffered saline – TBS (20 mmol/l Tris, 150 mmol/l NaCl, pH 7.4-7.6), rinsed in TBS and then incubated for 2 hours in mouse monoclonal antibody to Cx43 (Sigma) that was diluted 1:1000 in blocking buffer. After immunoblots were repeatedly rinsed in TBS, they were incubated for one hour with donkey anti-mouse IgG antibody coupled with alkaline phosphatase (Promega) that was diluted 1:2000. The bands were developed with the BCIP-NBT (5-bromo-4-chloro-3 indolyl phosphate/ nitro blue tetrazolium) method (Promega).

Densitometric analysis of Cx43 signals detected on Western blot was performed with morphometric software GelPro System (Media Cybernetics, USA).

Transmission electron microscopy

For TEM, aortae were cut into 3 mm rings and fixed with 2.5 % glutaraldehyde in 100 mmol/l cacodylate buffer (pH 7.4) for 3 h. After washing in cacodylate buffer, tissue was postfixed in 1 % OsO4, dehydrated via ethanol series, infiltrated by propylene oxide and embedded in Epon 812. Ultra-thin sections (65 nm) were cut on ultramicrotome (LKB Huxley, Stockholm, Sweden), counter-stained with uranyl acetate and lead citrate, then examined in the electron microscope Tesla 500 (Brno, Czech Republic).

Statistical analysis

All results are presented as mean ± SEM. Body weight, blood pressure and heart rate were analyzed using one-way ANOVA and followed by Duncan’s post-hoc test. All other data were analyzed using one-way ANOVA and Student t-test. Values were considered to differ significantly when p<0.05.

Results

Basic parameters

Body weight of Wistar, BHR and SHR was 309±3 g, 316±4 g and 287±4 g, respectively (p<0.01 SHR vs. W). Blood pressure of BHR and SHR was significantly increased (p<0.001) when compared to Wistar rats (132±2 mm Hg and 185±3 mm Hg vs. 110±2 mm Hg). Heart rate of Wistar rats and BHR did not differ significantly (414±11 bpm vs. 410±17 bpm), while elevated HR was found in SHR (452±10 bpm, p<0.05) vs. both Wistar and BHR.
Cx43 expression in the aorta

Immunofluorescent staining of Cx43 showed differences in distribution between normotensive Wistar and both groups of hypertensive rats (Fig. 1). In normotensive aortae, we observed intensive punctuated Cx43 immunolabeling lining in both the endothelium and the smooth muscle cell of media (Fig. 1A). The expression pattern of Cx43 in the aorta of BHR showed irregular and decreased intensity of punctuated staining in the endothelium (Fig. 1B) compared to Wistar rats. In addition, immunolabeled spots of Cx43 were sparsely scattered through the endothelium and media of the aorta in SHR (Fig. 1C).

Western blot analysis of total protein extracted from the aortae demonstrated two immunoreactive bands of Cx43 protein isoforms corresponding to phosphorylated (P1) and non-phosphorylated (P0) isoforms of Cx43 in aorta of Wistar rats (lane 1), BHR (line 2) and SHR (line 3). Values are mean ± SEM. * p<0.05 vs. control, ** p<0.001 vs. control. IOD – integrated density, BHR – borderline hypertensive rats, SHR – spontaneously hypertensive rats.

Transmission electron microscopy

The electron microscopy of the aortic endothelial cells of Wistar rats showed normal structure of the endothelial gap and tight junctions (Fig. 3). On the other hand, the focal significant subcellular alterations of inter-endothelial connections in aorta of SHR were found (Fig. 4) manifested by irregular cell overlapping and extended tight junctions containing degraded membrane structures. In the aortic endothelium of BHR (Fig. 5) local irregular extended tight junctions were observed. In addition also long gap junctions connecting two adjacent cells in different stages of subcellular injury could be seen. These changes were less marked and their occurrence was less frequent than those observed in SHR.

Discussion

In the present study we demonstrated alterations of Cx43 distribution in endothelial monolayer, decreased Cx43 expression and ultrastructural alterations of endothelial gap junction and tight junctions in the aortic wall of spontaneously hypertensive rats. However, the most important finding of this study is that above mentioned changes in vascular structure of the aorta were present also in prehypertensive period. Although the occurrence and frequency of structural changes was less pronounced in prehypertensive period than in fully established hypertension, data clearly suggest that already borderline elevation of BP may be associated with impairment of endothelial monolayer followed by remodeling of vascular wall.

Arterial hypertension is one of the most important risk factors for development of other cardiovascular diseases. Despite significant research progress on the field of hypertension, the role of vascular changes in the development of hypertension is still unclear. Several authors showed significant functional
abnormalities in the various types of blood vessels in SHR, including endothelial dysfunction (Konishi and Su 1983, Čačányiová et al. 2006, Török et al. 2006) while other authors, including our group, did not observe reduced endothelial function in SHR or BHR (Konishi and Su 1986, Wirth et al. 1996, Bernátová et al. 2007, Puzserová et al. 2007). On the other hand, vascular remodeling and decreased elasticity of arteries was observed in adult (Gabriels and Paul 1998, Haefliger et al. 2001, Kansui et al. 2004) as well as in young SHR (Koprdová et al. 2007). The vascular remodeling is also accompanied with increased endothelial permeability and disturbances of endothelial integrity – both determined by alterations of cell-to-cell connections including gap junctions. The aorta, which is a sparsely innervated and electrically quiescent vessel, is likely to be particularly dependent on gap junctional communications for coordinating the response of endothelial cells and smooth muscle cells to diverse signals (Christ et al. 1996). Cx43, together with Cx37 and Cx40, represent major gap junction proteins expressed in endothelium of large vessels including aorta (Bruzzone et al. 1993, Gabriels and Paul 1998). The Cx43 expression may vary depending on the species and various regions of vascular tree (Bruzzone et al. 1993, Little et al. 1995).

Several studies have reported that the high blood pressure alters connexin expression in vascular system. It is interesting that Cx43 expression was increased in aorta of rats made hypertensive by clipping one renal artery or in DOCA-salt model, while it was decreased in aorta of rats made hypertensive by the inhibition of NO synthase activity (Haefliger et al. 1999, Haefliger and Meda 2000). In SHR, elevated Cx43 was observed in the mesenteric artery (Kansui et al. 2004) while unchanged expression was observed in the caudal artery (Rummery et al. 2002). The above mentioned results indicate different regulation of Cx43 in various experimental models of hypertension and support the heterogeneity of Cx43 expression in the...
Various parts of the vascular tree. Our finding demonstrated decreased Cx43 expression in the aorta already in prehypertensive rats, which was more pronounced in the aorta of rats with fully established hypertension. Suppression of Cx43 expression, together with subcellular alterations of endothelial connections in the aorta of BHR indicate that even moderately increased blood pressure may represent a risk factor affecting the intercellular communication, endothelial permeability and integrity which may consequently influence aortic function in later period of life (Wirth et al. 1996) or in interaction with other environmental factors, such as stress (Bernátová et al. 2007).

Regarding the relation between Cx43 and aortic function, we cannot conclude whether reduced Cx43 expression and junctions’ morphology in the endothelium were associated with alterations in the aortic function because we did not monitor the dye transfer between endothelial cells. However, our recent study showed the ultrastructural injury of the endothelial cells in Wistar-mothered BHR (Okruhliková et al. 2007), suggesting possible impairment of endothelial function in the aorta. This impairment was supposedly related rather to EDHF-mediated responses (Kansui et al. 2004) than NO, since we observed elevated NO production in the aorta of both BHR and SHR (Bernátová et al. 2007).

In conclusion, the present study demonstrated decreased Cx43 expression and ultrastructural alterations of interendothelial junctions in the aorta of both BHR and SHR. The results indicate that already moderately increased blood pressure might represent a risk factor affecting interendothelial communication that may likely contribute to the further vascular wall remodeling and thus supposedly to participate in acceleration of hypertensive disease.

**Conflict of Interest**

There is no conflict of interest.

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