Fibroblast Growth Factor 21 Attenuates Vascular Calcification by Alleviating Endoplasmic Reticulum Stress Mediated Apoptosis in Rats

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Abstract

Fibroblast growth factor 21 (FGF21), a hormone with multiple metabolic properties, has proven to be pleiotropic biological effects and may play pivotal role in numerous cardiovascular and metabolic diseases in the future. Vascular calcification (VC) is a concomitant pathological process of various cardiovascular and metabolic diseases. However, the effects of FGF21 on VC remain unclear. Therefore, in this research, we aimed to explore the roles and mechanisms of FGF21 in VC induced by vitamin D₃ plus nicotine (VDN) treatment rats. After 28 days VDN treatment, the calcium overload was confirmed by blood pressure, ultrasound imaging, calcium content, ALP activity and aortic pathological characteristics. In terms of FGF21, exogenous FGF21 can ameliorate the elevation of blood pressure, aortic calcification and related injury in VC rats. To investigate the mechanisms of FGF21 on VC, the endoplasmic reticulum stress (ERS) mediated apoptosis pathways were tested. As a method to detect apoptosis, the increased positive TUNEL staining cells were alleviated by FGF21 treatment. Furthermore, exogenous FGF21 can suppress the increased ERS chaperone, GRP78, in the calcified aortas. In the three pathways of ERS mediated apoptosis, we found CHOP pathway and caspase-12 pathway were involved in the treatment of FGF21, but not p-JNK/JNK pathway. Our study proved for the first time that FGF21 can inhibit the progress of VC by alleviating ERS mediated apoptosis in rats. FGF21 might be a new target for preventing and treating VC.

Key words: fibroblast growth factor 21, vascular calcification, endoplasmic reticulum stress, apoptosis

Introduction

Vascular calcification (VC) is a concomitant pathological process of various cardiovascular diseases[1]. The deposition of calcium-phosphorus contributed to the structural damage, fibrosis proliferation and the formation of calcium nodules in the middle layer of the artery, which can result in the thickening of arterial walls and the reducing of vascular compliance[2]. In contrary to the traditional view, VC is now regarded as an automatic, regulatable and revisable procedure controlled by multiple factors[3]. Particularly, previous researchers have proven that apoptosis of vascular smooth muscle cells (VSMCs) exist extensively in calcified aorta[4]. In addition, several pathological studies have found that apoptotic bodies released from VSMCs can initiate VC[5]. Thus, inhibiting apoptosis of VSMCs may alleviate VC.

Endoplasmic reticulum stress (ERS), a key reason to mediate apoptosis, has a significant effect on various metabolic disorders related diseases such as diabetes[6], atherosclerosis[7] and chronic kidney disease[8], all of which are related to VC. Furthermore, in the procedure of VC, calcium overloaded in VSMCs may disrupt the endoplasmic reticulum homeostasis and result in the unfolded protein response, which can induce the ERS and apoptosis[9].
Stated thus, to restrain ERS mediated apoptosis may attenuate the development of VC.

Recently, many studies have revealed that various biological factors such as cardiovascular peptides[10], cytokines and hormone[11] have key regulatory roles in the progression of VC. FGF21 is an endocrine hormone with multiple biological properties[12, 13]. Unlike other paracrine/autocrine FGFs, FGF21 is released from metabolic organs/tissues into blood and mediates its pleiotropic effects as an extracellular protein by activating its receptor FGFR1 and its cofactor β-Klotho[14, 15]. Increasing evidences point out that FGF21 plays a protective role on many cardiovascular events or metabolism associated diseases such as coronary atherosclerotic heart disease[16, 17], obesity[18, 19] and diabetes[20, 21]. Furthermore, our previous study has proven that FGF21 could ameliorate atherosclerosis via inhibiting ERS mediated apoptosis[22]. However, the biological functions of FGF21 in VC by modulating ERS mediated apoptosis are unknown yet. The thorough investigation into the mechanisms of the ameliorative effect of FGF21 on VC may contribute to the transformation from bench to bedside and be useful to discover a new strategy for treating VC.

Vitamin D₃ plus nicotine (VDN) induced rat model, first established by Niederhoffer et al, is a classical rat model of VC[23]. Therefore, we used the VDN rat model and administrated FGF21 to investigate the mechanisms of FGF21 on VC. Our results first reveal the evidences that FGF21 can protect VDN induced VC by inhibiting the ERS mediated apoptosis in vivo.

Materials and methods

Materials and reagents

FGF21 was purchased from Phoenix Pharmaceuticals (Belmont, CA, USA). Alzet Mini-Osmotic Pumps (model 2004) were purchased from Durect Corp. (Cupertino, CA, USA). Vitamin D₃, nicotine, Alizarin-red S and Hoechst 33342 were purchased from Sigma-Aldrich (St. Louis, MO, USA). Calcium assay kit was purchased from Zhongsheng Biosino Bio-technology and Science INC (Beijing, China). Alkaline phosphatase (ALP) kit was purchased from Jiancheng Biological Corp. (Nanjing, Jiangsu, China). Antibodies against GRP78, CHOP and caspase-12 were purchased from Abcam (Cambridge, UK). Antibodies against p-JNK and JNK were purchased from Cell Signaling Technology (Danvers, MA, USA). Antibodies against GAPDH and all secondary antibodies were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). The kit for terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL) staining was purchased from Roche Applied Science (Indianapolis, IN, USA). All other reagents were of analytical grade.

Rats model of VC and experimental groups

Male Sprague-Dawley rats (180-200g) were purchased from the Laboratory Animal Center of Peking University (Beijing, China). All animal care and experimental protocols were performed in accordance with the Animal Management Rule of the Ministry of Health (Document No. 55, 2001), People’s Republic of China. Rats were randomly divided into three groups (n=13): (1) control group (Con group), (2) VDN group and (3) FGF21 treatment VDN group (FGF21 group). The method of establishing VDN model was described as previously[23]. Briefly, the VDN and FGF21 group rats were administrated with nicotine (25 mg/kg in 5 ml peanut oil) intragastrically and injected intramuscularly with vitamin D₃ (3×10⁵ IU/kg) at 9 a.m. on day 1. And the nicotine was administrated again at 5 p.m. on the same day. The control group rats received the saline injection and peanut oil oral gavage only. For FGF21 treatment, all rats were anesthetized by 1.5% pentobarbital sodium (40 mg/kg), then an osmotic mini pump was implanted subcutaneously. The FGF21 group pumps were infused with FGF21 (70 μg/kg/d), together with the infusion of saline for control and VDN group [24].

Ultrasound imaging

Before the examination, the rats were anesthetized by 1.5% pentobarbital sodium (40 mg/kg). Then the thoracoabdominal hair was shaved carefully, and the ultrasound transmission gel was smeared on the naked skin. The ultrasonographic images were obtained by the imaging system (Vevo770TM Visual Sonics Inc., Toronto, Canada) in B mode. All rats survived in the imaging process.

Blood pressure measurement and specimens collection

At the end of experiment, rats were anesthetized with ethylcarbamate (10 mg/kg, i.p.). Intubate a polyethylene catheter (PE-50) filled with heparin through the right carotid artery for 30 min, then the rat aortic blood pressure was measured by a pressure transducer of PowerLab instrument (model BL-420F) (TaiMeng, Chengdu, Sichuan, China). Finally, the blood samples were drawn and mixed with 500 KIU/ml aprotinin and 1 mg/ml ethylene diamine tetraacetic acid-Na₂. Plasma was obtained by blood samples centrifuging at 3000 g for 15 min at 4°C and stored at -70°C. The intact aorta was removed to 4°C PBS and all samples of tissues were stored at -70°C until use[24].
Hematoxylin and eosin (HE) staining

After fixed in formalin overnight, the aortic tissue was embedded in paraffin and sectioned for HE staining. Briefly, the slides were placed in the hematoxylin staining solution first, and then were washed and treated with 1% acidic alcohol. Afterwards, the slides were stained by eosin. Finally, the slides were washed, dehydrated and treated by xylene before microscopy. The aortic slides were evaluated by use of Image Pro Plus 6.0 system.

Alizarin-red S staining

After dehydrated and rinsed rapidly in distilled water, the aortic slides were placed in the Alizarin-red S staining solution for 30 min at room temperature. When a red-orange color appeared, removed all unbound stain from the slides, then the slides were photographed by use of a microscope, Olympus BX 50 (Olympus Optical, Tokyo, Japan).

Calcium content in aorta

The calcium content in aortic tissue was extracted and measured by a calcium assay kit according to the manufacturer instructions. Briefly, 10 mg of aortas were separated and dried at 55 ℃, and the dry weights were recorded. Then the dried tissue was nitrified by nitric acid. After removing the nitric acid by heating, the calcium was dissolved in the solution containing 27 NM KCl and 27 μM LaCl₃. Finally, the calcium content of the aorta was measured and standardized by the aortic tissues dry weight.

ALP activity in aorta

The ALP activity in aortic tissue was measured by an ALP assay kit followed its instructions. Briefly, the aortic tissue homogenate was mixed with a reaction mixture (stock substration solution: alkaline buffer =1:1) at a ratio of 1:1, respectively. Then the mixtures were incubated at 37 ℃ for 15 min. After incubation, 1.5 ml of the developer was added to each mixture, and the absorbance of each well at 520 nm was recorded. The activity of ALP in each specimen incubation, 1.5 ml of the developer was added to each well at 520 nm was measured. The ALP activity in aortic tissue was measured by the Bradford method and standardized by the aortic tissues dry weight.

Western blot analysis

The rat tissues were homogenized with a lysis buffer which contains 0.1 mol/L NaCl, 500 KIU/mL aprotinin, 0.01 mol/L Tris–HCl (pH 7.5) and 1 mmol/L ethylene diamine tetraacetic acid (EDTA). Equal amounts of protein sample were loaded at 10% or 12% SDS gels and then transferred to a nitrocellulose membrane. Afterwards, incubated the membrane with 5% non-fat dry milk for 1 h to block the non-specific proteins. Then the membranes were incubated with primary antibodies, anti-GAPDH (1:1000), anti-GRP78 (1:1000), anti-CHOP (1:200), anti-p-JNK (1:1000), anti-JNK (1:1000) and anti-Caspases-12 (1:500) overnight at 4 ℃, and with secondary antibody (HRP-conjugated anti-rabbit or anti-goat IgG) at room temperature for 1 h. The protein expression was analyzed by NIH image software and normalized to that of GAPDH. All experiments were repeated at least 3 times.

TUNEL staining

Apoptosis of aortas was evaluated by TUNEL staining according to the TUNEL TMR red kit. Briefly, the ascending aorta was separated and fixed in formalin overnight, then embedded in paraffin and cut at 40 μM intervals along the aortic tissue. Then the sections were permeabilized and stained with TMR Red. The nuclei were counterstained with Hoechst 33342. Finally, the fluorescence signal was monitored under a fluorescence inverted microscope (Leica Imaging Systems, Cambridge, UK).

Statistical analysis

The data were calculated as the mean ± SD form least three independent experiments and analyzed by a one-way ANOVA. The difference was accepted at P < 0.05. A P < 0.05 was considered statistically significant, and a P < 0.01 was considered very statistically significant.

Results

FGF21 improved the general characteristics of VDN rats

First, the effects of FGF21 on the blood pressure of VC rats were evaluated. Under VDN treatment, as compared to the normal group, the systolic blood pressure (SBP), diastolic blood pressure (DBP) and mean blood pressure (MBP) of model group began to increase at the day 7 by 21.0% (P < 0.01), 38.1% (P < 0.01) and 30.9% (P < 0.01), respectively. At the day 21, the blood pressures got to the top with 22.5% (P < 0.01), 43.9% (P < 0.01) and 33.9% (P < 0.01), respectively. At the end of our study, SBP, DBP and MBP of model group were dramatically increased by 22.5% (P < 0.01), 43.9% (P < 0.01) and 33.9% (P < 0.01), respectively. And compared with the model group, FGF21-treated group shows the mitigation of VDN-induced hemodynamic disorders at the beginning of day 14 by 7.9% (P < 0.01), 7.2% (P < 0.05)
and 7.4% \((P<0.05)\), respectively. And at the end of the study, the decreasing rate of SBP, DBP and MBP were 9.7% \((P<0.01)\), 19.9% \((P<0.01)\) and 15.5% \((P<0.01)\) compared to the VDN group, respectively (Figure 1A, B and C).

**FGF21 restored VDN-induced pathological injury of the aorta in VC rats**

Next, the influence of FGF21 on the structure of calcified aortas was identified. As shown in Figure 2A, the ultrasonography showed the formation of calcified plaques and aorta wall thicken distinctly, resulting in serious VC by VDN treatment. In terms of the microscopic pathological changes, the histopathologic examination also showed disordered elastic fibers and thickened vessel walls (Figure 2B) in the aortas. And compared with the VDN group, FGF21 treatment significantly restored the pathological disorder caused by VDN through relieving the irregular proliferation of fibrous tissue and formation of calcified nodules in the middle layer of the aorta (Figure 2A and B).

**FGF21 ameliorated VC in VDN induced rats**

The roles of FGF21 in the mineralization of aorta were further investigated. Compared with the normal group, the model group showed stronger calcium salt deposition in aortas that the aortic calcium content was significantly increased by 7.03-fold \((P<0.01)\), Figure 3A). Meanwhile, the ALP activity, a key inducer of mineralization in tissue calcification, was also increased by 2.51-fold in aortas \((P<0.01)\), Figure 3B). In addition, the Alizarin-red S staining also showed widespread calcification in the middle layer of the aorta (Figure 3C). As compared with the VC alone group, FGF21 group showed lower calcium deposition by Alizarin-red S staining (Figure 3C) as well as markedly decreased the aortic calcium content, together with aortic ALP activity by 65.77% \((P<0.01)\) and 54.18% \((P<0.05)\) respectively (Figure 3A and B).

![Figure 1](http://www.ijbs.com) General hemodynamic tendency of rats in vitamin D3 plus nicotine (VDN) model. (A) Hemodynamic function of systolic blood pressure (SBP) tendency in vascular calcification (VC) rats. (B) Diastolic blood pressure (DBP) tendency in VDN induced VC rats. (C) Mean blood pressure (MBP) tendency in VDN induced VC rats \((n=11, \text{respectively})\). **P < 0.01 compared with the normal control group. #P < 0.05 compared with the VDN group. ##P < 0.01 compared with the VDN group.

![Figure 2](http://www.ijbs.com) Morphological study of calcified aortas by echocardiogram and Hematoxylin and eosin (HE) staining. (A) Images show short-axis views of ascending aorta, including the brachiocephalic artery branch, aortic arch and proximal descending aorta. Arrow: Calcified plaque. (B) HE staining of disordered elastic fibers in aortas of VC rats \((\text{magnification } 200\times)\).
Figure 3. FGF21 ameliorated VDN-induced VC in rats. (A) Effect of exogenous FGF21 on the aortic calcium content in VC rats (n=7). (B) Effect of exogenous FGF21 on aortic ALP activity in VC rats (n=7). (C) Alizarin-red S staining of vascular calcium deposition in aortas of VC rats (magnification 200×). **P < 0.01 compared with the control group. #P < 0.05 compared with the VDN group. ##P < 0.01 compared with the VDN group.

Figure 4. The apoptosis level of vascular smooth muscle cells (VSMCs) in rat aortas. Representative immunofluorescence staining of terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL, TMR Red) in calcified aortas. Blue indicates VSMCs nuclei by Hoechst staining (magnification 200×).

FGF21 inhibited apoptosis in the VSMCs

Since apoptosis of VSMCs has been proven to be a major cause of VC, the aortic tissues were drawn for TUNEL staining to test whether FGF21 shows effects on the apoptosis of VSMCs. Compared to the control group, the terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling-positive cells of aortic tissues could be observed dispersedly in the VDN group, which represented an extensive apoptosis in VSMCs. Whereas, compared to the model group, FGF21 treatment for 28 days significantly decreased the amount of terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling-positive cells in the aortic media (Figure 4). This result indicated that the effects of FGF21 on VC may be resulted from its inhibiting effect on apoptosis.

FGF21 down-regulated the ERS chaperones in calcified aortas

There have been studies proving that the apoptosis of VSMC in VC is caused by ERS. To assess whether FGF21 has function in the ERS pathways, we further detected the expression of GRP78, a main ERS chaperone in calcified aortas. Western blot analysis showed that the GRP78 level was significantly elevated by 108.8% (P < 0.01) in the model group as compared with controls (Figure 5A). And with the treatment of FGF21, the GRP78 protein level was down-regulated by 31.9% (P < 0.05, Figure 5A) which indicates that FGF21 may inhibit apoptosis by down-regulating the ERS pathways.

FGF21 decreased the ERS mediated apoptosis pathways in VDN induced VC rats

As is well known, there are three major signaling pathways in the ERS, and CHOP, JNK and caspase-12 are three cardinal biomolecules in the branches of ERS...
mediated apoptosis. As the results showed above, ERS and apoptosis joined the VDN induced VC and FGF21 could restrain the ERS and apoptosis respectively. However, which signaling pathway can FGF21 play a role in is unclear. Therefore, we examined these three key biomarkers respectively by Western blot analysis furtherly. As shown in Figure 5B and 5D, in VDN group, the CHOP and cleaved caspase-12 significantly increased by 109.3% ($P < 0.01$) and 103.1% ($P < 0.01$), respectively, compared with the controls. Moreover, FGF21 significantly suppressed the CHOP and cleaved caspase-12 activation by 31.0% ($P < 0.05$) and 27.9% ($P < 0.05$), respectively. However, the level of JNK phosphorylation seemed not involved in the ERS caused by VDN (Figure 5C). These results demonstrated that FGF21 attenuated VC through suppressing ERS mediated apoptosis by inhibiting the activation of CHOP and caspase-12 pathways, but not by p-JNK/JNK.

**Discussion**

FGF21, a peculiar member of the FGFs superfamily, has multiple metabolic effects and anticipated therapeutic applications on various diseases such as atherosclerosis[26], diabetes[27], and chronic kidney disease[28]. In our study, FGF21 treatment can ameliorate the elevation of blood pressure induced by VC in clinical manifestation. Going deep into the pathology, exogenous FGF21 can alleviate the histological disorder, calcium content and ALP activity of the aortas. For the mechanism research, FGF21 can reduce the apoptosis mediated by ERS in the aortic tissue of VDN rats. These results suggest that FGF21 may mitigate VC by inhibiting ERS mediated apoptosis. Thus, FGF21 may be a new target to prevent and alleviate VC.

VC is a crucial risk factor of many cardiovascular diseases[29]. The increased vessel wall stiffness and vascular wall thicken resulted from VC leads to increased pulse pressure and pulse wave velocity, which subsequently brings on various cardio-cerebrovascular diseases[2]. Several mechanisms have been proven involved in the progress of VC, including apoptosis, loss of inhibition and osteogenic differentiation[30].

In our study, a rat model of VC was established using the VDN method, to research the effects of FGF21 on VC and reveal the mechanism behind it. VDN induced rat model, first established by Niederhoffer et al in 1997[23], is a classical rat model of VC which is stable and dependable, widely used to investigate VC. Hypervitaminosis D increases the deposition of calcium on the elastic fibers of arteries. And nicotine amplifies these effects of Vitamin D in arteries. VDN-induced calcium overload on the arteries has similarities with the calcification in human athero- and arteriosclerosis[31]. In our VDN model, the calcium content and ALP activity, key factors and markers for active osteochondrogenesis, were dramatically higher than that of the normal group. HE staining and Alizarin-red S staining observed that the aortic elastic plate of the VDN group was significantly thicker and more disordered, with breakage of the elastic fibers and widespread calcification compared with the control group. The increased SBP, MBP and DBP we found in VDN rats might result from the increased arterial stiffness, which consisted with the alterations in human hemodynamic characteristics of calcified arteries. All these data agree with those of prior researches[23] which indicated that our construction of the rat VC model was successful and stable.

As a special member of FGFs, FGF21 has a major role in maintaining cardiovascular homeostasis[12, 32]. In our research, compared with the model group, rats treated with FGF21 showed alleviated aortic calcification induced by VDN, as shown by decreased SBP, DBP and MBP, ameliorated aortic calcium content and ALP activity, and regular elastic fibers and less calcification in the vessel wall. These results suggested that FGF21 can inhibit VC in rats.

![Image](http://www.ijbs.com)
Apoptosis is considered as a forerunner to vascular barrier destruction, which is a critical mechanism serving to VC[33]. During the procedure of VC, both apoptotic bodies induced by exposing to calcium and phosphate deposition and matrix vesicles derived from VSMCs of vessels may lead to concentrate the local calcium-phosphate, which will provide an appropriate microenvironment for nucleation[34]. As well-known, the transformation of VSMCs phenotype into osteoblast-like phenotype is involved in the pathogenesis of VC. Apoptosis is involved in the process of VSMCs phenotypic transformation[35]. Apoptosis occurred prior to the onset of calcification and that VSMC “blebs” or apoptotic bodies could induce calcium concentration in a crystallized form. Apoptotic bodies derived from VSMCs have similarities with matrix vesicles. Therefore, apoptosis is a key initiating process for VC and that, once an original nucleation site is formed, calcium-phosphate crystal growth can occur[36]. Based on this theory, attenuating the osteoblast-like phenotypic transition of VSMCs through inhibition of apoptosis might be an effective treatment for VC. In addition, accumulating data suggests that apoptosis mediated by ERS is close related to VC because the calcium accumulation in the cytoplasm may damage endoplasmic reticulum homeostasis[37]. Endoplasmic reticulum is the primary site for Ca\(^{2+}\) storage. Any stimulus that interferes with calcium homeostasis may lead to endoplasmic reticulum function disturbance, accumulating unfolded or misfolded protein in the endoplasmic reticulum, namely ERS[38]. Therefore, inhibition of ERS induced VSMC apoptosis may alleviate the development of VC.

Consistent with previous research[39], in the aorta of VDN challenged rats, positive TUNEL staining confirmed the apoptotic VSMCs in the aortas. Furthermore, the Western blot analysis showed a significant increase of ERS markers, GRP78, in the VDN model group. GRP78, a defined biomolecular chaperone, is a critical regulator for the unfolded protein response network by binding to endoplasmic reticulum sensors and inhibiting their activation[40]. The up-regulation of the ERS landmark and the increased positive apoptotic cells suggested an ERS mediated apoptosis response during VC.

The initiation of ERS mediated apoptosis may trigger by three signaling pathways: CHOP, caspase-12 and JNK pathways[41]. Therefore, we researched the possible signaling pathways to ERS mediated apoptosis in VC furtherly. CHOP, one of the ERS response hallmarks, can directly regulate the expression of numerous pro-apoptotic proteins, which leads to enhanced oxidative damage and thus apoptosis[42]. In our VC rats, CHOP was up-regulated in the aortas, which demonstrated that CHOP pathway mediated apoptosis procedure was closely involved in VC. Caspases, a superfamily of cysteine proteases, are major mediators of programmed cell death, namely apoptosis. Among these superfamily, caspase-12 is activated only by ERS[43]. When activated to the cleaved form, caspase-12 can activate caspase-9 directly, which in turn activates caspase-3, and finally leads to apoptosis[43]. In our research, caspase-12 was cleaved, namely being activated, which suggested that caspase-12 pathway mediated apoptosis was also involved in VC. Besides, the JNK pathway mediates another apoptotic pathway[44]. However, in our VDN model, there were no changes in protein levels of p-JNK, which means that the p-JNK/JNK pathway may not be the major pathway of apoptosis in VC rats.

For FGF21 treatment, we found FGF21 administration can decrease TUNEL-positive cells in the aortas as well as alleviate the increased expression of ERS marker GRP78 in the calcified aortas. In the three pathways to ERS, the expression of CHOP in calcified aorta was significantly down-regulated under FGF21 treatment. Also, FGF21 can inhibit the caspase-12 activation but not the p-JNK/JNK. Therefore, FGF21 may reduce apoptosis via inhibiting ERS in VC. These results are in accordance with recent studies that FGF21 can inhibit ERS induced apoptosis of VSMC in atherosclerotic mice[22].

As the possible mechanism of the vascular protective and anti-ERS effect of FGF21, some reports demonstrate that FGF21 is a downstream factor of ERS which can exert beneficial feedback effects through counteracting ERS[45]. However, because FGF21 was originally found in the liver and highly secreted by

![Figure 6. Schematic of pathways for FGF21 signal transduction. FGF21 attenuated VC through suppressing ERS mediated apoptosis by inhibiting the activation of CHOP and caspase-12 pathways, but not by p-JNK/JNK.](http://www.ijbs.com)
the adipose tissue, most researches focused on the effects of FGF21 on the hepatocyte, adipocyte and even cardiomyocyte[46] that the effects of FGF21 on the artery has been neglected. Actually, the existence of FGF21, β-Klotho and FGFR1 in the cardiac microvascular endothelial cells and VSMCs suggests that FGF21 may be a key target in the physiological and pathophysiological aspects of artery[47].

For the pathogenesis of liver disease study, ERS stimulated by triglycerides can directly increase FGF21 expression via a PERK-ATF4-CHOP dependent mechanism[48]. When the ERS occurs, one of ERS specific pathway, containing CHOP, is activated, and consequently expression of endogenous FGF21 is upregulated[49]. Furthermore, intraperitoneal injection of tunicamycin, an endoplasmic reticulum stressor, to normal mice can also induce hepatic FGF21 expression directly[50]. The same results are also confirmed in the adipocyte[45] and H9C2 cardiomyocytes[51]. Especially for the CHOP, some studies have found CHOP can induce the expression of FGF21 by activating transcription via promoter elements and enhancing FGF21 mRNA stability in ERS[49]. And in our previous study, we have confirmed that the expression of endogenous FGF21 were increased in the development of VC[52]. Taken together, the CHOP pathway in ERS may play a crucial role in ERS induced FGF21 elevation in the related diseases. These findings may further explain the correlation between ERS and increased FGF21 expression. As a result, the elevation of FGF21 can feedback control the ERS responses and attenuate the development of the ERS related diseases. The negative feedback mechanisms can prevent an excessive activation during the ERS and they can trigger cellular recovery. For instance, Zarei et al. revealed that FGF21 may protect against hepatic steatosis by alleviating ERS[53]. Other researchers reported that the administration of FGF21 could reverse the suppression of adiponectin expression, counteracts obesity and alleviates insulin resistance via inhibiting ERS in mice[45, 54]. In particular for the cardiovascular diseases, FGF21 could reduce the hypoxia-induced apoptosis of human pulmonary arterial endothelial cells and prevent endothelial dysfunction via the inhibition of ERS[55]. In addition, FGF21 overexpression can protect against ERS induced cardiomyocyte injury via increasing cell viability and reducing cell apoptosis[51]. FGF21 can also protect against ischemia cerebral injury via decreasing ERS[56]. In addition, our team first reported that FGF21 can protect against atherosclerosis at least in part by its inhibitory effects on ERS mediated apoptosis[22]. These researches clearly demonstrate that FGF21 administration or over-expression can protect tissues and organs against the adverse effects induced by ERS. In our study, administration of FGF21 in turn ameliorated the ERS responses of VC by suppressing CHOP and caspase-12 pathways. These imply that the ERS-FGF21 axis creates a negative feedback loop and our results show that FGF21 inhibits two of the three ERS induction downstream effectors in VC. Therefore, FGF21 appears to be a regulatory factor in the proteostasis network and not merely an endpoint regulator in the negative feedback loop. ERS is thought to be caused not only by the excess misfolded or unfolded proteins in the endoplasmic reticulum but also by pathogenic conditions, such as atherosclerosis[57] and diabetes[58]. In addition, treatment with chemical chaperones that reduce ERS can improve several metabolic diseases, including VC and obesity[59]. Based on our data, together with these previous findings, we propose that FGF21 may function as a critical regulator in the proteostasis network to maintain protein homeostasis by protecting against ERS not only in the VC but also in other metabolic disorders caused by ERS.

There were also several limitations in our research, which still need to be explored in the future. Firstly, we have been yet regretting that we did not confirm our in vivo study in vitro. Actually, it is worth to have a further discussion that in vitro study we plan next. In addition, our team had just published a paper recently which matched our in vivo study in vitro[60]. Involvement of RANK/RANKL/OPG/TRAIL system in VC has been researched widely, but it is still full of controversy. In our previous study, FGF21 may inhibit VSMCs calcification through OPG/RANKL system by P38 and PI3K/AKT pathways in vitro. Secondly, in this study, we used the echocardiography to evaluate the calcification condition, but in the clinical study, intravascular ultrasound (IVUS) is more common to investigate the vascular lumen morphology and calcification condition. However, the rat aorta was too tiny that the IVUS cannot pass. Subsequently, we plan to use the IVUS in a VC pig model to evaluate the VC continuously and dynamically.

In conclusion, FGF21 is likely an endogenous cardiovascular protective peptide. Using a rat model of VC, we propose that FGF21 can significantly alleviate the progression of VC in vivo. These data, to our knowledge, provide the first evidence that in the aorta of VDN challenged VC rat, the expression of apoptotic cells and pro-apoptosis protein levels of ERS pathway: CHOP and cleaved caspase-12, were decreased by FGF21. Thus, FGF21 is a potent cardiovascular protective peptide which can inhibit VC at least in part by alleviating ERS mediated
apoptosis in VSMCs. This study provides new avenues and targets for prevention and treatment of VC.

Abbreviations

FGF21: fibroblast growth factor 21; VDN: vitamin D3 plus nicotine; VC: vascular calcification; FGFR1: fibroblast growth factor receptor 1; ALP: alkaline phosphatase; VSMC: Vascular smooth muscle cell; ERS: endoplasmic reticulum stress; GRP 78: glucose-regulated protein 78; JNK: c-JUN NH2-terminal kinase; CHOP: C/EBP homologous protein.

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Competing Interests

The authors have declared that no competing interest exists.

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