EXPERIMENTAL STUDY

Crocin and voluntary exercise promote heart angiogenesis through Akt and ERK1/2 signalling in type 2 diabetic rats

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ABSTRACT

OBJECTIVES: The aim of this study was to evaluate the effects of crocin and voluntary exercise on heart angiogenesis through Akt and ERK1/2 signalling.

METHODS: Animals were divided into 4 groups as follows: diabetes (Dia), diabetic-crocin (Dia-Cro), diabetic-voluntary exercise (Dia-Exe), diabetic-crocin-voluntary exercise (Dia-Cro-Exe). Type 2 diabetes was induced by high-fat diet (4 weeks) and injection of streptozotocin (STZ) (i.p., 35 mg/kg). Animals received crocin orally (50 mg/kg), voluntary exercise was performed alone or together with crocin for 8 weeks. Akt and ERK1/2 levels were measured by ELISA and CD31 was detected by immunohistochemistry.

RESULTS: Akt and ERK1/2 levels in crocin and exercise groups were significantly higher than in diabetic group. Levels of Akt and ERK1/2 proteins and CD31 immunostaining were also significantly higher in crocin-voluntary exercise group in comparison to the other groups.

CONCLUSION: Our results revealed that treatments with crocin and voluntary exercise had synergistic effect on angiogenesis. Protective effects of these interventions are probably through the activation of Akt and ERK1/2 signalling pathways in the heart of diabetic rats (Fig. 3, Ref. 40). Text in PDF www.elis.sk.

KEY WORDS: type 2 diabetes, crocin, voluntary exercise, Akt; ERK1/2.

Introduction

Type 2 diabetes mellitus is a growing public health problem affecting millions of people worldwide. Its symptoms include progressive hyperglycemia, insulin resistance, and reduction in insulin secretion by pancreatic β-cells in response to increased blood glucose (1, 2). Patients with diabetes have a 4-fold greater risk of developing coronary arterial disease (3). Diabetes causes endothelial dysfunction and thereby contributes to the development of coronary artery disease (4). The molecular mechanisms underlying the impairment of angiogenesis in type 2 diabetes have remained largely unidentified. Recent studies revealed that diabetic abnormal angiogenesis was closely associated with an impaired Akt and ERK1/2 signalling (5). Hyperglycemia alters Akt and ERK1/2 signalling and leads to endothelial cell proliferative dysfunction (6). In diabetic patients, the expression of the vascular endothelial growth factor (VEGF) receptors decreases through reduced Akt phosphorylation, which in the long term worsens ischemic state. VEGF plays an important role in angiogenesis that activates cell proliferation via ERK1/2 (7, 8). On the other hand, expression of ERK1/2 protein, an activator of VEGF induced cell proliferation, was decreased in diabetic animals. Type 2 diabetes is associated with reduced Akt and ERK1/2 signalling along with impaired angiogenesis and collateral blood vessel formation.

Crocin, an active component of Crocus sativus (saffron), has been an important topic for research recently. Crocin exhibits a variety of effects including a decrease in the fasting blood glucose, HbA1c, lipid profile, and it also increases serum insulin levels (9). Furthermore, crocin has a protective effect on hypoxic damage of myocardial cells by elevation of VEGF expression (10). Active component of saffron enhances angiogenesis by a response of endodermic cells to VEGF (11). However, angiogenesis signalling pathways have not been extensively studied for understanding crocin-induced cardioprotection, particularly in diabetic animal models.

Exercise is an applicable treatment modality for diabetic patients. Training could also improve coronary vascular function by suppressing oxidative stress and inflammation in a type 2 diabetes (12). In response to training, two cellular pathways have been implicated in cellular growth: ERK and Akt (13). Aerobic exercise improves aging-induced downregulation of VEGF angiogenic signalling via phosphorylation of Akt protein in the heart, thereby contributing to the training-induced angiogenesis (14). Exercise reduced the heart disease risk and improved diabetic-mediated cardiovascular complications (15). Although many beneficial effects were reported for exercise, exhaustive exercise induces oxidative stress through a production of reactive oxygen species and can...
caused damage to organs (16). Voluntary exercise is ranged in mild/moderate exercise and is a useful therapeutic approach in cardiovascular diseases. In the animal model of voluntary exercise, the animal has free entrance to a running wheel and utilizes the wheel according to its physiological threshold for physical activity (17).

Based on these observations, this experiment was designed to test the hypothesis that crocin and voluntary exercise can improve the heart angiogenesis via Akt and ERK1/2 signalling in diabetic rats.

Material and methods

Animals

Male Wistar rats (200–250 g) were obtained from Tabriz medical faculty (Iran-Tabriz). They were housed four per cage except the rats in the exercise groups that were alone. Animals were housed in a room with a constant temperature of 24 °C, a relative humidity of 50 %, and a 12 h dark / light cycle with access to food and water ad libitum. This study was approved by the Animal Ethics Committee (document number 92197) in accordance with the instruction for the care and use of laboratory animals prepared by Tabriz University of Medical Sciences (Iran-Tabriz).

Experimental design

All rats were made diabetic by a high fat diet and STZ injection, then they were randomly divided into four groups (n = 7). Group 1 received a high-fat diet (4 weeks) and low dose of STZ (35 mg/kg) as a diabetic group (Dia). Group 2 received a single dose of crocin (50 mg/kg) for eight weeks after confirmation of Diabetes (Dia-Cro). Group 3 performed voluntary exercise for eight weeks after confirmation of Diabetes (Dia-Exe). Group 4 received crocin and simultaneously performed voluntary exercise for eight weeks after confirmation of Diabetes (Dia-Cro-Exe). Crocin (Sigma) was administered (50 mg/kg) 6 days a week for 8 weeks. For the assessment of voluntary exercise, rats were housed individually in a cage containing a wheel (1.00 m circumference, Tajhiz Gostar). This stainless-steel running wheel was equipped with a digital magnetic counter that was activated by wheel rotation. Each exercising rat had a separate running wheel in its cage that allowed to run voluntarily during 8 weeks of the study. The rats, whose running distance was lower than ~2000 m per day were eliminated before statistical analysis (18).

Development of type 2 diabetes

After 1 week ad libitum exposure to drinking water and standard rat pellet diet, rats were fed with high fat diet comprising 22 % fat, carbohydrate 48 % and protein 20 % in blend with standard laboratory chow consisting of 5 % fat, 53 % carbohydrate and 23 % protein for 4 weeks. After the period of dietary manipulation, a single low dose of STZ (35 mg/kg, dissolved in 0.1 M sodium citrate buffer at pH 4.4) was injected into each rat intraperitoneally (Sigma) (19). After 72 hours of streptozotocin administration, fasting blood glucose level was checked by using a portable glucometer (Glucoplus Inc., Quebec, Canada) in the blood collected from the tail veins. Animals with fasting blood glucose value > 300 mg/dl were considered diabetic. Animals with fasting blood glucose concentration < 300 mg/dl were excluded from the study (20).

Heart tissue preparation for Akt and ERK1/2 and CD31

On the final day of the experiment, rats were sacrificed under anaesthesia with ketamine/xylazine (88/10 mg/kg, i.p.). Heart tissue was immediately removed and washed with saline 0.9 %. Hearts were excised, frozen in liquid nitrogen and stored at deep freeze (~70 °C) for later measurements.

Akt and ERK1/2 assays

Heart tissue samples were weighted (0.1 g), homogenized in PBS (pH 7.2–7.4) and centrifuged for 20 min at the speed of 3000 rpm and 4 °C. Then supernatants were collected in a new tube and p-Akt (Cat. No. CK-E91385; Eastbiopharm Co., Ltd., Hangzhou, China), p-ERK1/2 (PT202/Y204) (Abcam, Cambridge, U.K, product code: ab176640) levels were measured using sandwich rat ELISA kits. The ELISA assay was performed according to the manufacturer’s instructions and normalized to the total protein concentration for each sample, as determined by the Bradford assay.

Immunohistochemical assessments

For the investigation of angiogenesis in the heart tissue, samples from left ventricle were immersed into 10% formalin after excision, embedded in paraffin, and cut into 4 μm-thick slices. Sections were deparaffinized in xylene and dehydrated in a graded series of ethanol. Slides were incubated sequentially in proteinase K and treated by 0.3 % hydrogen peroxide for blocking endogenous peroxidase activity. Sections were overlaid by primary antibody CD31 (Santa Cruz Biotechnology (Santa Cruz, CA, USA), a marker of angiogenesis and incubated at +4 °C overnight. Sections were then washed and incubated with standard avidin–biotin complex (ABC; Santa Cruz Biotechnology) according to the manufacturer’s instructions. Finally, sections were assessed by light microscope (Olympus BX 40, Japan) with an independent pathologist (21). Capillaries were visualized in the myocardium as a brown precipitate. Vascular structures positive for CD31 were counted for 5 to 6 slides per animal and 10 fields per slide.

Statistical analysis

Statistical analysis was performed using SPSS v.18.0 (IBM Corp., Armonk, NY, USA). Between group differences were analysed using two-way analysis of variance (ANOVA). Post-hoc analysis for Akt and ERK data were Tukey’s test. The results are reported as the means ± SEM, and were considered significant at p < 0.05.

Results

Effect of crocin and voluntary exercise on Akt levels

After 8 weeks of treatment of diabetic rats with crocin and voluntary exercise, P-Akt levels in heart tissue in Dia-Cro and Dia-Exe were significantly (p < 0.05, p < 0.01 respectively) higher than in Dia group (Fig. 1). In the crocin combined with exercise group, the heart P-Akt levels were significantly increased compared to...
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Diabetic animals (p < 0.001). A significant difference was found between Dia-Cro-Exe group and Dia-Cro (p < 0.01) or Dia-Exe groups (p < 0.05). Figure 1 indicates that crocin combined with voluntary exercise had an additive effect in Akt protein levels in heart tissue.

**Effect of crocin and voluntary exercise on ERK1/2**

According to the Figure 2, two-way ANOVA indicated that treatment with crocin and exercise significantly enhanced the ERK1/2 levels in Dia-Cro (p < 0.01), Dia-Exe (p < 0.01), and Dia-Cro-Exe (p < 0.001) in comparison with the diabetic group. Also, ERK1/2 levels in Dia-Cro-Exe group were significantly higher than those of the other treatment groups (p < 0.001). These findings exhibit that p-ERK1/2 increased by crocin and voluntary exercise in the diabetic groups.

**Effect of crocin and voluntary exercise on angiogenesis**

Immunohistochemical staining for CD31 in the cardiac sections are shown in Figure 3. CD31 was present in the diabetic group treated with crocin and voluntary exercise. There was a significant increase in CD31 immunostaining in the cardiac tissue in the crocin-exercise-treated group (p < 0.001), when compared to the diabetic group (Fig. 3 A, D). Also, treatment of diabetic animals with crocin and voluntary exercise alone, significantly elevated angiogenesis compared to the diabetic group (all, p < 0.001) (Fig. 3 A, B, C).
Discussion

The present study showed that crocin and voluntary exercise treatment increased the Akt and ERK1/2 activation in diabetic rat heart. In addition, they stimulated angiogenesis in heart tissue of diabetic rats. For the first time, our findings demonstrated that heart angiogenesis and its upstream molecules including Akt and ERK1/2 were upregulated in response to crocin combined with voluntary exercise in rats with high-fat diet-induced type 2 diabetes. Type 2 diabetes represented a formidable unmet medical health problem with an elevated risk of cardiovascular disease. In this pathologic case, there is an inadequate collateral vascular formation that ultimately results in enhanced cardiovascular morbidity and mortality rates (4). Based on previous studies it was found that in diabetes, angiogenesis is regulated dependently on organ-tissue, and cell type-specific manner (22). In the diabetic heart tissue, VEGF (pro-angiogenic molecules) signalling pathways are disturbed and collateral vessel formation is interrupted (23). VEGF exerts many of its effects via the AKT and ERK1/2 pathways (24). During developmental vasculogenesis, the AKT pathway regulates venous specification, whereas the ERK pathway regulates arterial specification (25). VEGF activates AKT via a phosphatidylinositol 3-kinase (PI 3-kinase)-dependent pathway, and this downstream effector may play a principal role in the “outstanding” biological processes of VEGF, such as: endothelial cell proliferation, migration, and survival (26). There are several lines of evidence indicating that under diabetic condition, the expression level of phosphorylated Akt and ERK1/2 is decreased in cardiomyocytes (27), in spite of high level of VEGF (28). Our results proved that crocin combined with voluntary exercise enhanced angiogenesis and its upstream components of angiogenic signalling such as: Akt and ERK1/2. Previous studies showed that training promotes cardiac angiogenesis by enhancing the angiogenic pathways such as: ERK1/2 and PI3K/Akt both in healthy and obese animals (29, 30). Research confirmed that the expression levels of VEGF-A and VEGF-R2 were increased in saffron therapy group and it was suggested that the protective effects of saffron might be related to its ability to promote angiogenesis (11, 31). Cardioprotective effects of crocin were reported in some studies that are related to modulation of antioxidant activities and cardiac biomarkers (32). Crocin prevented retinal ischemia/reperfusion injury in retinal ganglion cells through the PI3K/Akt signalling pathway (33). Saffron strongly enhanced the phosphorylation of ERK in muscle cells and increased insulin sensitivity. Also, when the cells were exposed to saffron and insulin simultaneously, PI3K/Akt pathway was more greatly stimulated compared to those of individual treatments (34). Several rationales may be considered for this crocin-exercise-induced upregulation of Akt and ERK1/2. One possible elucidation for the ameliorated phosphorylation levels of Akt and ERK1/2 in heart tissue of the diabetic rat might be a mechanism related to glycemia (35). It was reported that hyperglycemia decreased Akt and ERK1/2 activities and further inhibited the proliferation and migration of cells necessary for neovascularization (36). Since it was found that normalized glycemia brought the phosphorylation and activity of Akt and ERK1/2 to a normal level (37), so our results that crocin combined with exercise increased P-Akt and P-ERK 1/2 levels could support this hypothesis. An additional reason for upregulation of Akt and ERK1/2 in heart tissue of diabetic rats by crocin-exercise may act via activating mitogen-activated protein kinase (MAP kinase/ERK1/2) and PI3K/Akt (34, 38, 39). Furthermore, there is a cross-talk between MAPK, PI3K pathways and hypoglycemic effects. Activation of these pathways promote the translocation of GLUT4 from an intracellular pool to the cell surface, which in turn can lead to an improvement of insulin stimulated glucose uptake and subsequently reduce glucose levels in diabetic cases (40).

Accordingly, training and crocin administration might be beneficial to enhance the cardiac survival in diabetic heart by ERK1/2 and Akt signalling pathways. This is the first report regarding increased angiogenesis due to administration of crocin combined with voluntary exercise through activation of Akt and ERK1/2 in heart tissue of type 2 diabetic rats. This study represents an initial step toward unravelling the mechanisms that underlie angiogenesis induced by crocin combined with voluntary exercise in type 2 diabetic model. In conclusion, our findings suggest that crocin combined with voluntary exercise can reverse reduced angiogenesis by high glucose in heart tissue of type 2 diabetes. In addition, crocin and voluntary exercise induce coronary angiogenesis by activating the Akt and ERK signalling pathways. Indeed, combination of voluntary exercise with crocin administration in type 2 diabetic rats has beneficial effects compared to each treatment.

References


