Short Communication

Luteolin and fisetin suppress oxidative stress by modulating sirtuins and forkhead box O3a expression under in vitro diabetic conditions

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BACKGROUND/OBJECTIVE: Chronic hyperglycemia induces oxidative stress via accumulation of reactive oxygen species (ROS) and contributes to diabetic complications. Hyperglycemia induces mitochondrial superoxide anion production through the increased activity of nicotinamide adenine dinucleotide phosphate (NADPH) oxidase. This study aimed to determine whether fisetin and luteolin treatments suppress the oxidative stress by modulating the expression of sirtuins (SIRTs) and forkhead box O3a (FOXO3a) under hyperglycemic conditions in human monocytes.

MATERIALS/METHODS: Human monocytic cells (THP-1) were cultured under osmotic control (14.5 mmol/L mannitol), normoglycemic (NG, 5.5 mmol/L glucose), or hyperglycemic (HG, 20 mmol/L glucose) conditions, in the absence or presence of fisetin and luteolin for 48 h. To determine the effect of fisetin and luteolin treatments on high glucose-induced oxidative stress, western blotting and intracellular staining were performed.

RESULTS: Hyperglycemic conditions increased the ROS production, as compared to normoglycemic condition. However, fisetin and luteolin treatments inhibited ROS production under hyperglycemia. To obtain further insight into ROS production in hyperglycemic conditions, evaluation of p47phox expression revealed that fisetin and luteolin treatments inhibited p47phox expression under hyperglycemic conditions. Conversely, the expression levels of SIRT1, SIRT3, SIRT6, and FOXO3a were decreased under high glucose conditions compared to normal glucose conditions, but exposure to fisetin and luteolin induced the expression of SIRT1, SIRT3, SIRT6, and FOXO3a. The above findings suggest that fisetin and luteolin inhibited high glucose-induced ROS production in monocytes through the activation of SIRTs and FOXO3a.

CONCLUSIONS: The results of our study supports current researches that state fisetin and luteolin as potential agents for the development of novel strategies for diabetes.


Keywords: Diabetes mellitus, oxidative stress, monocytes

INTRODUCTION

Hyperglycemia contributes to diabetes and several diabetic complications, such as retinopathy, neuropathy, and nephropathy [1,2]. High glucose (HG) has been shown to induce inflammatory cytokines, chemokines, p38 mitogen-activator protein kinase, reactive oxygen species (ROS), protein kinase C (PKC), and nuclear factor-κB (NF-κB) activity in both clinical and experimental systems [3,4]. Hyperglycemia increases the intracellular ROS formation, which is the prevailing mechanism leading to endothelial cell dysfunction, cellular damage and diabetic complications [5]. The phagocyte nicotinamide adenine dinucleotide phosphate (NADPH) oxidase plays an important role in ROS production and consists of several proteins in resting cells, such as p47phox (NADPH oxidase cytosolic proteins), p67phox, and p40phox [6]. In our previous studies, we observed that hyperglycemia increased the p47phox expression and ROS generation [7,8].

Sirtuins are a class of proteins known as NAD-dependent deacetylases. The sirtuin family is divided into seven homologs (i.e., SIRT1-SIRT7) in mammals. Sirtuin affects a wide range of cellular processes, such as aging, apoptosis, inflammatory response, calorie restriction, energy efficiency, and stress resistance [9]. Previously, we reported that SIRT1 expression and superoxide anion production are altered in type 1 diabetes patients compared to normal healthy controls [8]. SIRT1 is a key mediator of various biological functions in mammals, including glucose-lipid metabolism, mitochondrial biogenesis, and inflammation [10].

The forkhead box O (FOXO) proteins are a family of transcription factors characterized by a conserved DNA-binding domain [11]. The members of the FOXO family, FOXO1 (FKHR),
FOXO3 (FKHRL1), FOXO4 (AFX), and FOXO6, play important roles in apoptosis modulation, energy metabolism, DNA damage repair, cell differentiation, oxidative stress, and other cellular functions [12]. In addition, FOXO is the known substrate of SIRT1, which regulates the FOXO transcription factors through direct binding or deacetylation or both [13,14]. However, there is little evidence on the exact interaction between SIRTs and FOXO under hyperglycemic conditions.

Flavonoids are reported to exert various biological activities such as anti-cancer, anti-inflammatory, and cardiovascular preventive effects [15,16]. Several flavonoids have demonstrated anti-diabetic effects, including reduction of aldose reductase activity, regeneration of pancreatic cells, and enhancement of insulin release [17].

Fisetin (3,3’,4’,7-tetrahydroxyflavone), a bioactive flavonol found abundantly in fruits and vegetables, has chemopreventive, neuroprotective, anti-oxidant, anti-inflammatory, and other beneficial health effects [18,19]. It also inhibits the expression of cell adhesion molecules, formation of ROS, and activation of NF-κB in vascular inflammatory responses [20]. Luteolin (3’,4’,5,7-tetrahydroxyflavone), a natural flavonoid found in fruits, vegetables, and medicinal herbs, also exerts anti-inflammatory, anti-allergic, anti-cancer, anti-oxidant, and other beneficial health effects [21,22]. Luteolin is also known to improve blood glucose, HbA1c, insulin levels, and fatty acid metabolism-related gene expression [23]. Our previous studies revealed that under hyperglycemic conditions, both fisetin and luteolin (3-10 μM) inhibit pro-inflammatory cytokine production in monocytes via epigenetic changes involving NF-κB [24,25]. However, the molecular mechanism of SIRTs and FOXO in regulating mononuclear superoxide under hyperglycemic conditions in the presence and absence of dietary agents remains unknown. In this study, we hypothesize that under hyperglycemic conditions, fisetin and luteolin suppresses the ROS production through a mechanism involving modulation of p47phox and SIRTs/FOXO in human monocytes. This is the first report on the interactions between SIRTs and FOXO under diabetic conditions and exposure to fisetin and luteolin.

MATERIALS AND METHODS

Reagents

Fisetin and luteolin were purchased from Sigma Aldrich (St. Louis, MO, USA). Fisetin and luteolin stock solutions were prepared and stored in dimethyl sulfoxide (DMSO), and diluted with culture medium as required. The vehicle control in all experiments was 0.1% (v/v) DMSO. Anti-FoxO family antibodies were procured from Cell Signaling Technology (Beverly, MA, USA) and anti-SIRT1, SIRT3 and SIRT6 were procured from Abcam (Cambridge, MA, USA). The BCM™ protein assay kit was purchased from Pierce (Rockford, IL, USA). Novex pre-cast Tris-Glycine gels were obtained from Invitrogen (Carlsbad, CA, USA). All other chemicals, unless otherwise stated, were obtained from Sigma (St. Louis, MO, USA).

Cell culture and treatment

The human monocytic THP-1 cell line was obtained from the American Type Culture Collection (Manassas, VA, USA). THP-1 cell were cultured in RPMI medium containing 10% fetal bovine serum and 1% antibiotics, and incubated at 37°C in a 5% CO2 atmosphere. Fisetin and luteolin dissolved in DMSO were used for treatment of the cells. The final concentration of DMSO was 0.1% (v/v) for each concentration. THP-1 cells (1 x 105 cells/mL) were cultured in the presence of osmol control (14.5 mM/L mannitol), normal glycemic (NG, 5.5 mM/L glucose) or hyper-glycemic (HG, 20 mM/L glucose) conditions, in the absence or presence of fisetin (3, 5 and 10 μM) and luteolin (3, 5 and 10 μM) for 48 h. After 48h incubation, the medium was collected for measuring the cytokine release; the cell pellet was washed with phosphate-buffered saline (PBS) and cells were harvested for further experiments.

Cell viability assay

The toxicity of fisetin and luteolin on the cultured THP-1 cells was evaluated using the Cell Counting Kit-8 (CCK-8, Dojindo Molecular Technologies, Rockville, MD, USA), according to the manufacturer’s protocol. Briefly, THP-1 cells were seeded at 4 x 103 cells/well in a 96-well plate, and subsequently treated with various concentrations of the phytochemicals (3, 5 and 10 μM) for 48 h. Absorbance was measured using a Wallac EnVision microplate reader (PerkinElmer, Turku, Finland). The inhibitory effect of phytochemicals on the growth of cells was assessed as the percentage of cell growth reduction compared with vehicle-treated cells, which were defined as 100% viable.

Preparation of nuclear and cytoplasmic lysates

After treatment with fisetin and luteolin, the medium was aspirated and cells were washed twice in PBS (10 mM, pH 7.4). Nuclear lysates were prepared using NE-PER Nuclear and Cytoplasmic Extraction Reagents (Pierce, Rockford, IL, USA). Lysates were collected and cleared by centrifugation, and the supernatant was aliquoted and stored at -80°C.

Western blot analysis

For western blot analysis, treated cells were homogenized at 4°C in buffer containing 10 mM Tris-HCl (pH 7.5), 150 mM NaCl, 0.5% (v/v) Tween 20, 1 mM PMSF, and one protease inhibitor cocktail tablet (Roche, Basel-Stadt; Basel, Switzerland), and then centrifuged at 10,000 x g for 15 min. Protein concentration of the lysate was determined using the BCM™ protein assay kit. Samples (40 μg of total protein) were mixed with sample buffer (100 mM Tris-HCl, 2% sodium dodecyl sulfate, 1% 2-mercaptoethanol, 2% glycerol, and 0.01% bromophenol blue (pH 7.6)), incubated at 95°C for 15 min, and loaded on 12% polyacryla-mide gels. Electrophoresis was performed using the Mini Protean 3 Cell system (Bio-Rad, CA, USA). The resolved proteins were transferred on a nitrocellulose membrane (Schleicher & Schuell BioScience, Saxony, Dassel, Germany). For immunoblotting, the membranes were washed and incubated in blocking buffer (10 mM Tris-HCl pH 7.5, 150 mM NaCl, 0.1% Tween 20, and 3% nonfat dry milk), and then incubated with diluted primary antibodies (1:1000) for 2 h at room temperature. Following incubation with the primary antibody, membranes were washed 3 times and then probed with diluted secondary antibodies (1:2000) for 1 h. The membranes were washed 3 times (15 min each) and detected by chemiluminescence ECL.
kit (Amersham Life Sciences Inc., Pittsburg, PA, USA).

**Intracellular H$_2$O$_2$ staining**

Cells were cultured in a 6-well plate and incubated at 37°C overnight to allow attachment, after which they were cultured overnight in serum-free medium. After treatment with fisetin and luteolin for 48h, 2',7'-dichlorodihydrofluorescein diacetate (DCFH-DA) (5 μM) was added and incubated with the cells for 15 min. The cells were then washed with 1 × PBS buffer, and fixed with 10% formaldehyde. Fluorescent images were captured with a fluorescent microscope. The signal quantification was assessed by ImageJ software.

**Statistical analysis**

Each experiment was performed at least 3 times. All data are expressed as the mean ± standard error (SE). The significance of differences between groups was determined by applying the student’s t-test. Statistical significance is expressed as *P* < 0.05, **P** < 0.01.

**RESULTS**

**Cytotoxicity of luteolin and fisetin on monocytes exposed to hyperglycemic conditions**

Cytotoxic effects of fisetin and luteolin on THP-1 cells exposed to hyperglycemic conditions were investigated using the CCK-8 assay (Fig. 1). No toxicity was observed for both compounds at concentrations 3, 5 and 10 μM, at 48 h treatment under hyperglycemic conditions. All further experiments were performed in the non-toxic concentration range of fisetin and luteolin (3, 5 and 10 μM).

**Evaluation of luteolin and fisetin effects on O$_2^-$ production in hyperglycemic conditions using immunofluorescence assay**

We examined the production of superoxide ions using immunofluorescence analysis (Fig. 2). Under hyperglycemic conditions, superoxide production was increased compared to normal glucose conditions. However, in the presence of fisetin and luteolin, the superoxide production decreased to levels similar to that under normal glucose conditions.

**Effect of luteolin and fisetin on p47phox expression in hyperglycemic conditions**

p47phox is a component of monocyte NADPH oxidase that produces ROS [26]. To obtain further insight into ROS production under hyperglycemic conditions, we examined whether fisetin and luteolin treatments suppressed the expression of the p47phox gene (Fig. 3). As expected, the expression of p47phox gene was greater in HG conditions than in NG conditions. The increased expression of p47phox gene was downregulated by luteolin (Fig. 3A) and fisetin (Fig. 3B) treatments.

**Luteolin and fisetin treatments increase the expression levels of SIRTs and FOXO3a in HG-induced THP-1 cells**

To identify the specific underlying mechanisms, we
excretion, and oxidative stress [8-11]. Many studies suggest that SIRT1 is an effective therapeutic target for diabetes, and various natural compounds that can serve as activators of SIRT1 have been researched [10]. Loss of SIRT1 activity may be associated with metabolic diseases, such as diabetes [27]. Of the seven members of the sirtuin family, SIRT1, SIRT3, and SIRT6 play important roles in diabetes, inflammation, and oxidative stress [27-29]. SIRT3 is localized to the mitochondria and regulates metabolism, adenosine triphosphate (ATP) generation, and multiple additional metabolic processes [28]. SIRT6 is a chromatin-associated deacetylase known to be involved in aging, inflammation, metabolism, and stress response [29]. Among the FOXO family, FOXO3a is closely related to oxidative stress, and upregulates the transcription of ROS scavenging enzymes superoxide dismutase 2 (SOD2, also known as MnSOD) and catalase [11-12]. SIRT1 is especially associated with FOXO transcription factors and regulates gene-specific transcription [13-14]. SIRT3 also enhances deacetylase FOXO3a in mitochondria to prevent oxidative stress through DNA-binding to the MnSOD promoter [30]. Previously, we have shown that the increased expression of p47phox under hyperglycemic conditions decreases by fisetin and luteolin treatments. In addition, we observed that the increased expression of p47phox under hyperglycemic conditions decreases by fisetin and luteolin treatments. Venogopal et al. [7] showed that p47phox, an essential component of monocyte NADPH oxidase, is required for ROS generation under HG conditions. In type 2 diabetes patients, there was significant decrease in the monocytic SIRT1 expression and increase in the p47phox expression, compared to the control [7]. Moreover, SIRT3 plays an important role in mitochondrial oxidative stress by reducing the ROS production. We have previously reported the role of SIRT1 and FOXO3a in regulating monocyte superoxide under HG conditions [8]. However, the roles of SIRT3 and SIRT6 were not clearly understood in our previous study. In this study, we observed that SIRT1, SIRT3, SIRT6, and FOXO3a expression was decreased in hyperglycemia-induced monocytes. Conversely, the downregulated expressions of SIRT1, SIRT3, SIRT6, and FOXO3a were restored by fisetin and luteolin treatments to levels similar to normoglycemic conditions. Based on these findings, we suggest that hyperglycemia downregulates the SIRT1-FOXO3a pathway, resulting in p47phox expression and ROS generation. Fisetin and luteolin treatments under HG conditions modulate the SIRT1, SIRT3, SIRT6, and FOXO3a expressions, and subsequently decrease the p47phox expression and ROS production.

In conclusion, fisetin and luteolin inhibit ROS production through the activation of SIRT and FOXO3a expression. Therefore, we propose that fisetin and luteolin are potential agents for the prevention and treatment of diabetes and its associated complications.

CONFLICT OF INTEREST

The authors declare no potential conflicts of interests.
REFERENCES