Progressive Resistance Exercise Training Attenuated Renal Damages, but did not improve Muscle Force in STZ-Induced Diabetic Rats

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Abstract

A type of exercise (Resistance Exercise Training – RET) that is markedly applied to increase muscle mass has been prescribed to prevent muscle atrophy from catabolic conditions such as, diabetes and chronic kidney disease. However, how this type of exercise modulates renal system remains unclear. It prompted us to apply progressive RET to STZ-induced diabetes rats in order to investigate renal environment. Progressive RET was applied to Wistar rats under following exercise program: 6 to 12 climbs/day, 5 days/week, 8 weeks, at 50 to 80% of maximal loading test. After streptozotocin injection, animals were divided into four groups: two control groups (non-exercised and exercised) and two diabetic groups (non-exercised and exercised) with 4 to 6 rats/group. Kidney weight (KW), muscle weight, proteinuria and protein level (assessed by Western blot and multiplex technology) were determined.

It was found that RET did not protect exercised diabetic animals from loss of muscle mass. RET did not influence the performance in maximal loading test in diabetic groups (p>0.05); additionally, exercised diabetic animals did not recover body weight (p>0.05). KW was preserved in exercised diabetic animals (15.83 ± 2.6 mg/24h) when compared to diabetic group (37.66 ± 3.2 mg/24h, p<0.05). We assessed upstream and downstream of mTOR and found significant decreases in PI3K, p-Akt and p-4EBP1 in kidneys of exercised diabetic animals (p<0.05). Moreover, we observed that kidney protein level of TGFβ-1 was markedly decreased in exercised diabetic animals. Under progressive RET program, kidneys may be functionally protected and mTOR-signaling pathway plays important role on renal environment. However, skeletal muscle showed no improvement under this program, suggesting different mTOR modulation among renal and muscle functions.

Keywords: PI3K/Akt/mTOR-signaling pathway; Renal hypertrophy; Podocin; Fibrosis; Exercise prescription; Ladder climb; Muscle force

Introduction

Diabetic Nephropathy (DN) affects worldwide population and is the leading cause for end-stage renal disease. This complication is characterized by morphologic changes, including renal hypertrophy, augmented fibrosis and inflammation, increased proteinuria and also by a markedly impairment of renal function [1-3]. Increased kidney mass (i.e. renal hypertrophy) in DN is, in part, due to activation of mTOR (mammalian target of rapamycin, currently known as mechanistic target of rapamycin).

mTOR regulation includes cell growth, cell cycle, autophagy, and metabolism in several tissues such as, kidneys, skeletal muscle, vessels, heart and tumors. mTOR is an evolutionary conserved serine/threonine kinase and there are two complexes of mTOR that are functional and structurally different; complex 1 (mTORC1) is rapamycin (mTORC1 inhibitor) sensitive and is responsible for cell growth and proliferation. The complex 2 (mTORC2) does not respond to acute treatment of rapamycin and affects cell polarity and actin cytoskeleton; although both complexes can be activated by growth factors such as insulin and IGF-1, only mTORC1 responds to amino acids [4-6]. It is known that mTORC1 upstream regulation includes important proteins such as, lipid kinase phosphatidylinositol 3-kinase (PI3K), phosphatase and tensin homolog on chromosome 10 (PTEN), Akt (also known as protein kinase B) and tuberous sclerosis complex 2 (TSC1/2). The process begins with activation of PI3K that phosphorylates PI2 to PIP2; PTEN dephosphorylates PIP2 back to PIP2 and prevents Akt activation [7]. However, failure to increase PTEN expression culminates in increased Akt phosphorylation, and TSC1/2 (downstream of Akt) is also phosphorylated releasing mTORC1 from its inhibiting action. Downstream of mTORC1 is characterized by activation of ribosomal protein and S6 kinase and phosphorylation of eukaryotic initiation factor 4E biding protein 1 (4E-BP1), which increase cell growth [8-10].

RET is an important and essential applied method to gain muscle mass and it has been markedly accepted as treatment for skeletal muscle atrophy under catabolic conditions [4,5,11]. In skeletal muscle, mTOR-signaling pathway have been shown to play crucial role in hypertrophy [12,13]. Although, the joint position statement by American College of Sports Medicine (ACSM) and American Diabetes Association provided important information regarding prescription of RET and skeletal muscle adaptation in diabetes [6], little is known regarding how progressive RET modulates renal environment.

Hornberger et al. [14] demonstrated that a ladder climb apparatus is a well tool to improve skeletal muscle hypertrophy in normal rodents. We used the same apparatus to study the mechanism by which progressive RET might influence renal environment in diabetic rats. Additionally, because it is not fully understood how RET modulates renal system in diabetes mellitus, we believe that progressive RET might
modulate kidney environment through PI3K/Akt/mTOR-signaling pathway.

Material and Methods

Animals

Two months old male Wistar rats (n=4 to 6/group) obtained from Center of Biology and Medicine for Experimental Models of Federal University of Sao Paulo (CEDEME), were used throughout this study, and randomly assigned into 4 groups: two control groups (non-exercised and exercised) and two diabetic groups (non-exercised and exercised). Animals were fed standard laboratory chow and given tap water ad libitum while housed (4-5 per cage) in a temperature and humidity-controlled room (22°C and 60 ± 5%), with a 12:12 h light-dark cycle (lights on at 7 a.m.). All experimental procedures followed Institutional Guidelines for Care and Use of Laboratory Animals, and the Ethic’s Committee of Federal University of Sao Paulo, Brazil approved the protocols (Process number: 0185/11).

Diabetes was induced after 12h fasting by a single tail vein injection of streptozotocin (STZ, 50mg/kg-1 body weight; Sigma, Chemical, St. Louis, MO; in freshly prepared 0.01M citrate buffer, pH 4.5) [15,16]. Control animals were injected with citrate buffer only. Following this period, animals were kept for 3 days with free access to food and water. STZ-injected animals exhibited massive glycosuria and hyperglycemia within few days, and diabetes was confirmed by measuring fasting blood glucose concentration 5 days after drug injection. Fasting blood glucose was determined in blood samples obtained by tail prick, by using a strip operated glucometer (Accu-Check Sensor, Roche, Swiss) and rats were considered diabetic if they had fasting plasma glucose >250 mg/dL. In those animals we measured basic parameters such as body weight and renal function; we also measured protein level by western blot and multiplex technology. For that purpose, animals were anesthetized (ketamine 80 mg/kg + xylazine 12 mg/kg, IP) and a peritoneal incision was determined in blood samples obtained by tail prick, by using a strip operated glucometer (Accu-Check Sensor, Roche, Swiss) and rats were considered diabetic if they had fasting plasma glucose >250 mg/dL. In those animals we measured basic parameters such as body weight and renal function; we also measured protein level by western blot and multiplex technology. For that purpose, animals were anesthetized (ketamine 80 mg/kg + xylazine 12 mg/kg, IP) and a peritoneal incision was made to carefully remove kidneys. Subsequently, kidneys were weighted and immediately placed in liquid nitrogen. Next, kidneys were storage in -80°C.

Western blotting and reagents

Total kidney (30 to 50 mg) was mixed with 1mL of RIPA buffer. After protein concentration determination by BSA method, proteins ran in a SDS-page electrophoresis gel and were next electrotransferred to a 0.22μm PVDF membrane. Membranes were blocked (5% BSA) and incubated in primary antibodies (1:1000) at 4°C with gentle mixing overnight. Next, membranes were incubated in secondary antibodies (1:10000) at room temperature. Following the incubation, ECL detection was applied. Antibodies anti-PI3K p85 subunit, anti-GAPDH, and anti-total 4E-BP1 were purchased from Millipore (Temecula, CA, USA). Antibodies anti-total Akt, and anti-β-actin were purchased from Cell Signaling Technology (Beverly, MA, USA). Antibody anti-phospho 4E-BP1 was purchased from Calbiochem (Darmstadt, Germany). Antibody anti-podocin and anti-phospho Akt were purchased from Santa Cruz Biotechnology (Santa Cruz, CA).

Multiplex evaluation

We used kidney sample to assess transformer growth factor beta 1 (TGFβ-1) by multiplex assay (Milliplex xMAP multiplex technology – Millipore Billerica, MA, USA). Anti-TGFβ-1-labeled beads were re-suspended and aliquoted into a pre-washed 96 well filter plate. Twenty-five microliters of kidney sample or standards/controls were mixed with the beads in the assay buffer and incubated overnight at 4°C. Subsequently, the plate was incubated with biotinylated detection antibody and streptavidin-phyceroerythrin solution and run on the Luminex 200™ instrument (Luminex, Austin, TX).

Resistance exercise training

Ladder climb was the method we used to exercise animals. In the past, Hornberger et al. [14] developed a specific training method for rodents. We used that apparatus to perform resistance exercise training in exercised control and exercised diabetic animals. First, all animals were adapted (one week) to the ladder for apparatus recognition and also to improve climb (3 to 6 climbs, 1 minute of rest between climbs). Second, all animals were submitted to a test to determine the exercise training prescription, and every two weeks in order to adjust prescription and to evaluate exercise training protocol efficiency. Test consisted of climbs until the animals were not able to complete it, and the last successful climb was considered 100% of maximal produced force to prescribe exercise training protocol. The test was performed in all groups. After the test, we designed a prescription that consisted in 50 to 80% of 100% of maximal obtained produced force. Initial resistance exercise training program consisted of 6 climbs, 50% of maximal produced force and 1 minute of resting between climbs. Every week, we increased the number of climbs until 12 (adjustment were performed every two weeks based on a new test). After animals reached 12 climbs, we progressively increased the intensity of training (based on percentage). Final exercise program was 12 climbs, 80% of maximal produced force, and 1 minute of resting between climbs. One day after the last RET section, animals were submitted to the last test to evaluate the efficiency of resistance exercise training program (Figure 1).

Renal function and urinary protein content

Two days after the last RET session, rats were housed in individual metabolic cages, and 24 h urine was collected and centrifuged at 3,000 rpm. Creatinine and proteinuria were analyzed. The animals (all groups) were anesthetized at the end of the protocol period (ketamine 80 mg/kg + xylazine 12 mg/kg, IP) and a peritoneal incision was made to carefully remove kidneys. Subsequently, kidneys were weighted and immediately placed in liquid nitrogen. Next, kidneys were storage in -80°C.

Statistical analysis

Results are expressed as means ± SEM. Statistical differences were determined by two-way analysis of variance (Two-way ANOVA) and Bonferroni post-hoc test; It was used GraphPad Prism 6.0 Software to analyze the data. Differences were considered significant at p<0.05.
Results

Progressive RET in diabetic animals prevents increased renal hypertrophy

STZ-induced diabetes increases kidney weight gain. This characteristic increase in kidney weight (measured by hypertrophy index) was found in diabetic group, and surprisingly progressive RET prevented it in diabetic exercised animals (Figure 2A; p<0.05). Because the effects of RET on kidney are not clear, we assessed the effects of progressive RET in renal function, and found that proteinuria (Figure 2B) was markedly reduced in exercised diabetic group (15.83 ± 2.6 mg/24h) when compared to diabetic group (37.66 ± 3.2 mg/24h, p<0.001). Also, we evaluated renal 4EBP1 activation (Figure 3C; p<0.001) and found that progressive RET decreased its expression in exercised diabetic group. Additionally, we measured TGF-β1 (Figure 4) and found that progressive RET in diabetic animals (325.8 pg/mL) have decreased TGF-β1 concentration when compared to non-exercised diabetic animals (437 pg/mL, p<0.05).

Progressive RET did not preserve skeletal muscle function in diabetic trained animals

Markedly loss of body weight is a characteristic set from STZ-induced diabetic rats. As showed in Table 1, diabetic animals showed decreased body weight when compared to controls groups (exercised or not); additionally, exercised diabetic animals do not increased (p>0.05) body weight after 8 weeks of progressive RET. Although, RET is

PI3K/Akt-signaling pathway was decreased in diabetic exercised animals following progressive RET

Because kidney weight was decreased in exercised diabetic group (Figure 2A), we assessed PI3K/Akt-signaling pathway that is involved in cell growth, in order to explore the effects of progressive RET on diabetes. We found that in exercised diabetic group PI3K expression was decreased by progressive RET (Figure 3A; p<0.01) similar to what was observed for Akt phosphorylation (Figure 3B; p<0.01) when compared to non-exercised diabetic group. We also evaluated renal 4EBP1 activation (Figure 3C; p<0.001) and found that progressive RET decreased its expression in exercised diabetic group. Additionally, we measured TGF-β1 (Figure 4) and found that progressive RET in diabetic animals (325.8 pg/mL) have decreased TGF-β1 concentration when compared to non-exercised diabetic animals (437 pg/mL, p<0.05).

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traditionally performed to increase muscle mass, surprisingly, we found that progressive RET was not able to increase muscle weight of extensor digitorium longus (EDL) and plantaris in exercised diabetic animals (p<0.05). Also, characteristic gain of muscle force after RET was not statistic significantly between diabetic groups (p>0.05). Moreover, these outcomes were independently of blood glucose level (p>0.05).

Discussion

Resistance exercise training adaptations include increased muscle mass and function (i.e. force); it may occur in physiological or pathophysiological conditions such as, diabetes and CKD [11-13]. Using STZ-induced diabetes model we observed that skeletal muscle mass and force were not improved by progressive RET program applied in this study. However, there was observed a significant beneficial influence of progressive RET on kidney weight and proteinuria of exercised diabetic animals. Additionally, podocin expression was assessed and it was found increased in exercised diabetic animals when compared to non-exercised diabetic animals (Figure 2), which could be responsible for normalized proteinuria in exercised diabetic animals after 8 weeks progressive RET. This is relevant because podocyte is an important structure of glomeruli barrier, and preservation of podocyte integrity is imperative to avoid protein loss through glomerular membrane [1,8,17,18].

PI3K/Akt/mTOR-signaling pathway might be related with these adaptations and is crucial for understanding the complex mechanisms related to increased kidney weight and diabetes complications. In this
study, kidney weight gain could be correlated to augmented expression of PI3K, p-Akt and p-4EBP1; also quantification of TGFβ-1 in kidney was found increased in non-exercised diabetic animals. Potential effects of progressive RET were noticed and we observed attenuated and significant levels of protein expression (PI3K, p-Akt and p-4EBP1, and TGFβ-1 concentration) in kidney from exercised diabetic animals after 8 weeks of exercise. These outcomes suggest that progressive RET under 50 to 80% of maximal loading test improves renal system, however does not avoids diabetes-induced damage in skeletal muscle.

Mechanisms related to renal hypertrophy have been described in the literature. In compensatory renal hypertrophy by unilateral nephrectomy, the remaining kidney increases size and activation of mTOR is the main mechanism responsible for hypertrophy. Additionally, using rapamycin it could be blocked [18]. In DN, it has been shown that rapamycin was able to delay the progression of renal hypertrophy [19].

Catabolic condition-induced skeletal muscle atrophy, such as diabetes and CKD, normally lead skeletal muscle to loss of muscle weight and function, and this is at least in part due to increased inflammation and insulin resistance [10,20-23]. As we known, activation of muscle fibers by resistance exercise training (RET) or injury may increase mTORC1 activation improving protein synthesis and regulating several cellular process including translation and transcription [2,10,24]. On the other hand, atrophy process is provided by increased atrophic factors and decreased hypertrophy factors [10]. Protein breakdown may be increased through the activation of the ubiquitin-proteasome system that is markedly involved in muscle atrophy increasing mRNA and proteins known as atrogens such as, MuRF1 and Atrogin-1 [25-27]. On the other hand, catabolic conditions may decrease skeletal muscle hypertrophy with no involvement of atrogens, by decreasing activation of mTOR and its downstream, such as p70 S6K and 4EBP1 [10,21,28-31]. PI3K/Akt/mTOR-signaling pathway has been positively and extensively implicated on skeletal muscle hypertrophy under pathologic conditions. Only two weeks of muscle overload was reported to be sufficient to induce muscle hypertrophy and recruit satellite cells, which are responsible for skeletal muscle regeneration in a murine model of CKD [11]. In this study, skeletal muscle mass was not improved by progressive RET and it suggests that PI3K/Akt/mTOR-signaling pathway was not modulated or at least was not fully activated by RET; additionally, muscle force showed no difference between diabetic groups (Table 1). The fact that rats trained 5 days per week may not be excluded to explain skeletal muscle impairment; moreover, the ACSM guidelines recommend that RET must be performed as complement to aerobic exercise training [6]. Altogether, these outcomes might explain the absence of traditional response of skeletal muscle to progressive RET protocol used in this study.

The results from our study are encouraging, because it demonstrates that RET could be an important tool to prevent the evolution of diabetic nephropathy. It is also possible that preventing urinary protein content by RET, increase the adhesion to a regular RET program. Surprisingly, our study also demonstrates that skeletal muscles were not improved by our RET regimen, it suggests that different intensity or at least other type of exercise should be incorporated into a regular exercise program. As we known, other types of exercise, such as endurance exercise, are widely applied as treatment to diabetes and CKD [3,32,33]. Our group has demonstrated the protective effects of endurance exercise in diabetic animals that performed 14 weeks of running training on renal function [3]. It suggest that both, RET and endurance exercise should be applied for people who have diabetes.

In summary, this study provides important information regarding renal damages in DN and beneficial effects on kidneys induced by progressive RET. Kidneys from exercised diabetic rats decreased size and proteinuria, and increased podocin expression, this may improve renal function; additionally, PI3K/Akt/mTOR-signaling pathway modulates renal hypertrophy promoted by progressive RET. Finally, skeletal muscle from exercise diabetic animals were not improved after RET, suggesting that the prescription of RET used in this study induces different modulation among organs such as, kidneys and skeletal muscle. Different intensity of RET must be studied to provide complete beneficial effects of RET.

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