Astragalus polysaccharides affect insulin resistance by regulating the hepatic SIRT1-PGC-1α/PPARα-FGF21 signaling pathway in male Sprague Dawley rats undergoing catch-up growth

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Abstract. The present study investigated the effects of Astragalus polysaccharides (APS) on insulin resistance by modulation of hepatic sirtuin 1 (SIRT1)-peroxisome proliferator-activated receptor (PPAR)-γ coactivator (PGC)-1α, peroxisome proliferator-activated receptor (PPAR)α-fibroblast growth factor (FGF)21, and glucose and lipid metabolism. Thirty male Sprague Dawley rats were divided into three groups: A normal control group, a catch-up growth group and an APS-treated (APS-G) group. The latter two groups underwent food restriction for 4 weeks, prior to being provided with a high fat diet, which was available ad libitum. The APS-G group was orally treated with APS for 8 weeks, whereas the other groups were administered saline. Body weight was measured and an oral glucose tolerance test (OGTT) was conducted after 8 weeks. The plasma glucose and insulin levels obtained from the OGTT were assayed, and hepatic morphology was observed by light and transmission electron microscopy. In addition, the mRNA expression levels of PGC-1α/PPARα, and the protein expression levels of SIRT1, FGF21 and nuclear factor-κB were quantified in the liver and serum. APS treatment suppressed abnormal glycolipid metabolism and insulin resistance following 8 weeks of catch-up growth by improving hepatic SIRT1-PPARα-FGF21 intracellular signaling and reducing steatosis. The suppressive effects of APS on liver acetylation and glycolipid metabolism-associated molecules contributed to the observed suppression of insulin resistance. However, the mechanism underlying the effects of APS on insulin resistance requires further research in order to be elucidated. Rapid and long-term treatment with APS may provide a novel, safe and effective therapeutic strategy for type 2 diabetes.

Introduction

There are two primary risk factors for type 2 diabetes mellitus (T2DM): Genetic predisposition and environmental factors, in which the latter has a more important role. After a period of growth inhibition, the linear growth rate usually exceeds the normal range; this phenomenon, known as catch-up growth, was first described >40 years ago by Prader et al (1). Catch-up growth has become the predominant cause for the accelerated spread of T2DM in developing countries (2), such as China, in which rapid economical growth lead to an increase in the prevalence of T2DM in populations who were subjected to nutritional deficiency at a young age. A previous study demonstrated that low birth weight is an independent risk factor for type 2 diabetes in China (3). Further studies demonstrated that catch-up growth following intra-uterine growth retardation in rats (IUGR rats) resulted in increased liver peroxisome proliferator-activated receptor (PPAR)-γ coactivator (PGC)-1 promoter histone 3/lysine 9 acetylation levels (4,5), and increased mRNA and protein expression levels of PGC-1α and PGC-1, resulting in increased hepatic glucose output (6). In addition to causing the epigenetic histone deacetylation changes in IUGR rats, sirtuin 1 (SIRT1) deacetylation also results in the de-acetylation of PGC-1α/PPARα and nuclear factor (NF)-κB (7), as well as other non-histone molecules, thereby improving liver glucose levels, lipid metabolism and inflammation, and regulating insulin secretion. Epidemiological studies regarding catch-up growth are difficult to perform due to the absence of ideal catch-up growth research platforms and data. The high fat diet weight-matching method following a long period or calorie restriction may better reflect the characteristics of a catch-up growth population (8,9). Therefore, the...
study of the association between PGC-1α/PPARα acetylation levels, inflammation, steatosis and hepatic insulin resistance in a liver model may further elucidate the high incidence of T2DM in developing countries. An increase in liver SIRT1 deacetylation levels results in abnormal epigenetic changes, improved liver lipid metabolism and reduced inflammation (7), as well as facilitating early prevention of T2DM. Astragalus exerts antioxidative effects (10) and increases neural activation in two important central glucose-sensing regions of the brain (the paraventricular hypothalamus and the nucleus tractus solitarius) thereby augmenting the counterregulatory response to hypoglycemia (11). The present study aimed to investigate the effects of astragalus on the suppression of hypoglycemia via the liver, as well as the underlying mechanism of these effects. APS is an active component of astragalus, which prevents the development of diabetic cardiomyopathy in diabetic rats via the PPARα-mediated regulatory signaling pathway (12). The association between APS and hepatocyte SIRT1-PGC-1α/PPARα-mediated regulation of hepatic glucose and lipid metabolism, as well as the therapeutic potential of APS in T2DM merits further research.

Materials and methods

Ethical approval. All experimental procedures were approved by the Ethics of Animal Experiments Committee of the Tongji University School of Medicine (Shanghai, China; approval no. TJMED-012-006). The present study was conducted according to internationally recognized guidelines on animal welfare, as well as the regulations regarding animal welfare in Shanghai, China, and was conducted in accordance with the guidelines of the Chinese Council on Animal Care.

Animals. A total of 28 six-week-old male Sprague-Dawley rats (Center of Experimental Animals, Tongji University School of Medicine, Shanghai, China), weighing 140-160 g, were housed in wire-bottomed cages in 22±1˚C with a 12-h light/dark cycle. The rats were raised on a commercial pellet diet (Center of Experimental Animals) consisting of 22% protein, 66% carbohydrates and 12% fat, and were provided with access to tap water ad libitum. The rats were randomly divided into three groups: A normal diet control group (NC; n=10), a catch-up growth APS-treated group (APS-G; n=9) and a catch-up growth model group (CUGFR; n=9). The rats in the NC group were raised on an ordinary diet for 4 weeks (60% of the diet intake of the NC group) following which they were fed with pellet diet for 8 weeks, and the rats in the CUGFR group were subjected to a dietary restriction for 4 weeks (60% of the diet intake of the NC group) following which they were fed with a high fat diet (42% calories from fat), which was provided ad libitum. The rats in the APS-G group were orally treated with 700 mg/kg APS (content of effective components, 69%), purchased from the Beijing Centre Biology Co. Ltd., (Beijing, China), for 8 weeks. The other groups were administrated with saline irrigation at the same dosage. The body weight of each rat was measured every 7 days.

Animal treatment. The rats were subjected to fasting 15 h prior to intraperitoneal injection with 20% urethane anesthetic (5 ml/kg; Shanghai Hengyuan Biological Technology Co. Ltd, Shanghai, China). The abdominal cavities of the rats were surgically opened following local disinfection. Blood samples (8-10 ml) were obtained from the abdominal aorta, maintained at room temperature for 10 min prior to centrifugation in an Eppendorf (EP) tube containing EDTA (1 mg/ml) and then centrifuged for 20 min at 2000 xg at 4˚C. The supernatant was placed into a sterile EP tube and preserved at -20˚C. The blood samples were used to determine the serum FGF21 levels using an ELISA kit (BioVendor, Asheville, NC, USA). Liver function, blood lipid levels and glycated hemoglobin (HBA1C) levels were also analyzed. Plasma fatty acid, total cholesterol (TC) and triglyceride levels were determined using a NEFAC kit (Wako Pure Chemical Industries, Ltd., Osaka, Japan), which measures oxidase and peroxidase activity (Trinder reaction). Low-density lipoprotein (LDL) and high-density lipoprotein (HDL) were measured using an immunoassay kit (Daichii Pure Chemicals Co., Ltd., Tokyo, Japan), alanine aminotransferase (ALT) was determined using an IFCC kit (Roche Diagnostics, Basel, Switzerland). HBA1C levels were determined by immunoturbidimetry (cat. no. D-68305; Roche Diagnostics). The rats were fasted for 15 h prior to intraperitoneal injection of 20% urethane anesthetic (5 ml/kg). The abdominal cavities of the rats were surgically opened by pattern clamps and ophthalmology scissors (Shanghai Kang Chau Medical Instrument Co., Ltd., Shanghai, China) followed by local ethanol disinfection (Sinopharm Co., Ltd., Shanghai, China). Subsequently, blood was drawn from the abdominal aorta and anti-coagulated with EDTA (1 mg/ml). Plasma was obtained by centrifuging the blood at 2000 xg at 4˚C for 20 min and frozen at 20˚C for further analyses. The liver, epididymal adipose and perirenal fat tissues were separated and weighed under sterile conditions after the rats were sacrificed by decapitation. The liver tissue samples were immediately dissected, fixed in 2% glutaraldehyde (Sinopharm Co., Ltd) and 10% formalin or frozen in liquid nitrogen until further analysis. The liver index (the ratio of liver weight to total body weight of the rats) and internal body fat ratio (epidymal fat weight plus perirenal fat weight) were calculated.

Oral glucose tolerance test (OGTT) and lipid parameters. An OGTT test was conducted as follows: A total of 2 g glucose/100 g body weight was orally administered following overnight fasting for 15 h. Blood samples were collected from the rat tail venous plexus 0, 15, 30, 60 and 120 min after glucose treatment, in order to measure the blood glucose and plasma insulin concentration levels. Blood glucose was measured by a glucometer (Roche Accu-Chek Performa; Roche Diagnostics). The blood was then centrifuged and placed into an EP tube containing 1 mg/ml EDTA (Shanghai Zurui Biological Technology Co., Ltd., Shanghai, China), in order to measure the blood insulin levels using an ELISA kit (EMD Millipore, Billerica, MA, USA). The plasma was then either immediately used for experimentation or frozen at -20˚C for further analyses.

Western blot analysis. The frozen rat liver tissue samples were homogenized three times, for 15 sec, in ice-cold radioimmunoprecipitation assay homogenization buffer (Merck Chemical Technology Co. Ltd, Shanghai, China)
containing Protease Inhibitor Cocktail Set III (cat. no. 539134; Invitrogen Life Technologies, Inc., Carlsbad, CA, USA) and Phosphatase Inhibitor Cocktail Set V (cat. no. 524629; Invitrogen Life Technologies, Inc.) using a PRO 200 homogenizer (PRO Scientific, Inc., Oxford, CT, USA). The homogenate was centrifuged at 4°C for 30 min at 12,000 x g and the supernatant was used as the total protein concentration. The total protein concentration then underwent protein quantification according to the Bradford method (Protein Assay kit II; cat no. 5000002; Bio-Rad Laboratories, Inc., Hercules, CA, USA) according to the manufacturer's instructions. Protein assay dye reagent concentrate (Bio-Rad) and BSA (Thermo Fisher Scientific, Waltham, MA, USA) were used as the protein standard. Protein (40 µg) was separated by 4-12% polyacrylamide gels [depending on the protein of interest; NuPAGE 4-12% Bis-Tris Gel 1.0 mmx12 well; MOPS SDS running buffer (20X); both from Invitrogen Life Technologies, Inc.] and transferred onto nitrocellulose membranes (iBlot transfer stacks nitrocellulose, regular; Invitrogen Life Technologies) using a semi-dry transfer protocol. Precision Plus Protein Dual Color standards (Bio-Rad) and Novex® Sharp Pre-stained Protein standard (Invitrogen) were used as the protein ladder. Following membrane blocking with Odyssey Blocking Buffer (LI-COR Biotechnology, Lincoln, NE, USA), the membranes were incubated with the appropriate primary antibody: Rabbit anti-NF-κB p65 polyclonal immunoglobulin (IgG) (1:200; cat. no. sc-372; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), mouse anti-SIRT1 monoclonal antibody (1:2,000; cat. no. ab110304; Abcam, Cambridge, UK), and rabbit anti-human PPARγ polyclonal antibody (0.2-1 µg/ml; cat. no. LS-B5864; 1:500 dilution; LifeSpan BioSciences, Inc., Seattle, WA, USA) antibodies, diluted with fluorescently-labeled secondary antibody blocking buffer [IRDye® 800CW goat anti-mouse IgG (H+L); 1:10,000; cat. no. 925-32210; or IRDye® 800CW goat anti-rabbit IgG (H+L); 1:10,000; cat. no. 925-32211; both from LI-COR Biotechnology]. The membranes were then washed twice with phosphate-buffered saline (PBS; Decent Biotech, Göttingen, Germany) with 0.1% Tween 20 (Sigma, Santa Clara, CA, USA) to remove all residual antibodies, and scanned using an Odyssey Infrared Imaging system with IR fluorescence scan Odyssey 2.1 software (LI-COR Biotechnology). Quantitative values were obtained from the densitometric measurements of the western blotting bands using Image Quant software (Tool, 1D gel analysis; GE Healthcare Bio-Sciences, Pittsburgh, PA, USA).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). The liver mRNA expression levels of PGC-1α and PPARγ were assessed by RT-qPCR. The total RNA was extracted using TRizol reagent (Invitrogen Life Technologies, Inc.; cat. no. 47122). cDNA was reverse-transcribed from 2 µg DNase-treated total RNA using an iScript cDNA Synthesis kit (Bio-Rad Laboratories, Inc.; cat no. 170-8891) according to the manufacturer's instructions. The following primer sequences, obtained from Invitrogen Life Technologies, were used: Forward, 5'-CCGACAACCTCAGCAAGTCTCCTC-3' and reverse, 5'-CCTTGCCTGGCCCTCAAAGTCTC-3' for rat PGC-1α; and forward, 5'-GGCGCTGCTATAATTGTGCTGTG-3' and reverse, 5'-GAAGGTTGATCCTGATGTTGTG-3' for PPARγ. The cDNA (diluted to 1:20) served as a template for the RT-qPCR; 100 µM primer solution was diluted to 5 pmol/µl, and qPCR was performed using SYBR Premix ExTaq (Takara Biotechnology Co., Ltd., Dalian, China). The PCR mixture contained 10.5 µl SYBR Premix ExTaq, 0.5 µl of each primer, 1 µl cDNA and 7.5 µl deionized water. PCR was performed with thermocycling conditions of 95°C for 30 sec, 40 cycles at 95°C for 5 sec, and 60°C for 30 sec in a DNA cycler apparatus (ABI Prism 7900HT; Applied Biosystems, Thermo Fisher Scientific). Genes were quantified using the ΔΔCt method with GAPDH as a housekeeping gene. For each set of reactions, the samples were analyzed in triplicate.

Immunohistochemistry and morphometry. Following sacrifice, the livers of the rats were excised, and the liver wet weights were measured. A section of hepatic tissue (~3x3x10 mm) was removed via a horizontal incision 1 cm away from the edge of the right lobe. The hepatic tissue samples were fixed in 10% formalin, dehydrated with a graded series of ethanol, ethanol uranyl acetate and acetone (all from Sinopharm Co., Ltd.), and stained with hematoxylin and eosin (H&E; n=3 rats/group). Observation and image capture were conducted using a CKX31 light microscope (Olympus Corporation, Tokyo, Japan).

Transmission electron microscopy (TEM). Three 1-mm³ liver sections were fixed with 2.5% glutaraldehyde and stored at 4°C. The samples were subsequently analyzed at the Electron Microscopy Laboratory at the Shanghai University of Traditional Chinese Medicine (Shanghai, China). Following fixation, samples were rinsed with 0.1 M phosphate buffer and treated with 1% OsO₄ solution (Alfa Aesar Chemical Co., Ltd., Shanghai, China) for 3 h, dehydrated with a graded series of ethanol (30, 50, 80 and 90%) and propylene oxide, and embedded in epoxy resin (Shanghai Resin Factory Co., Ltd, Shanghai, China). Semi-thin (thickness, 500 nm) hepatic sections were stained with 1% toluidine blue (Sinopharm Co., Ltd) for histopathology. Suitable areas of ultrathin (thickness, 80-90 nm) hepatic sections were selected for ultrastructural analysis. These were sectioned using a diamond knife (diamond knife;cat no. 706602; Leica Microsystems, Wetzlar, Germany), mounted on a copper grid (Shanghai Chemical Experiment Audio Supplies Company Ltd, Shanghai, China), and stained with uranyl acetate and lead citrate. The sections were analyzed using a Tecnai-12 Bio T min TEM (Philips, Amsterdam, Netherlands), and observation and image capture were performed at an acceleration voltage of 80 kV.

Statistical analysis. Statistical differences between the groups were evaluated using Student' unpaired t-test. The homeostasis model assessment of insulin resistance (HOMA-IR) was used, where: HOMA-IR = [fasting plasma glucose (mmol/l) x fasting plasma insulin (mU/l)]/22.5. The data were presented as the mean ± standard deviation. Data analyses were performed using SPSS 18.0 (SPSS, Inc., Chicago, IL, USA). P<0.05 was considered to indicate a statistically significant difference.
Results

Growth of the rats. The growth curve obtained from weighing the rats weekly demonstrated that their growth was arrested during the period of 60% food restriction, and rapidly increased following re-feeding on a high fat diet for 1 week; however, the rat body weights of the CUGFR group increased less compared with the NC group. The growth curve obtained from the rat weights demonstrated that the APS-G group exhibited a significant trend towards weight loss, as compared with the CUGFR group. Conversely, the rats in the CUGFR group significantly gained weight compared to the rats in the NC group (Fig. 1).

Effects of APS on insulin sensitivity, serum lipid, serum FGF21 and alanine aminotransferase (ALT) levels in high fat diet-fed rats following catch-up growth. Prior to treatment with APS, glucose tolerance curves demonstrated marginally lower (although not statistically significant) serum glucose levels in the CUGFR group rats at 0, 15 and 30 min compared with the NC group. At 60 min, following glucose gavage, blood glucose levels were significantly higher than the NC group; these glucose levels failed to return to pre-meal state (Fig. 2A). The insulin curve demonstrated significantly lower insulin levels at 15 min in the CUGFR group compared with the NC group (P<0.05) and that insulin increase was delayed (Fig. 3A). The insulin levels of the APS-G group significantly decreased at 15 min, and the area under the curve of glucose increased, although no statistically significant difference was observed (Fig. 2B, and Fig. 3A and B).

Following treatment with APS, no statistically significant difference was observed between the blood glucose levels of all three groups, and the blood glucose levels of the APS-G rats did not return to their pre-meal state (Fig. 2C and D). The insulin levels of the CUGFR group significantly increased, when compared with the NC group (P<0.01), and treatment with APS moderately reduced this increase, although the results were not statistically significant (Fig. 3D). After 15 min the levels of insulin were significantly lower in the APS-G group, when compared with the CUGFR group, and no significant decline in these insulin levels occurred throughout the experiment (Fig. 3C). The HOMA-IR and fasting insulin index were significantly higher in the CUGFR group, as compared with the NC group, and following treatment with APS, the two decreased significantly (Table I).

Effects of APS on liver index and visceral fat content ratio of high fat diet-fed rats following catch-up growth. The liver index and visceral fat content ratio of the rats were significantly higher (P<0.01 and P<0.05, respectively) in the CUGFR group, as compared with the NC group. Following 8 weeks of APS treatment, liver weight in the APS-G group reduced significantly compared with the CUGFR group, however the decrease in the liver index was not significant. Although the visceral fat content ratio of the CUGFR group was significantly higher than that of the NC group, treatment with APS did not significantly affect the visceral fat content (Table III).

Effects of APS on hepatic pathology and TEM of high fat diet-fed rats following catch-up growth. Liver H&E staining demonstrated large quantities of lipid droplet deposition in the liver tissue samples of the CUGFR group, as compared with the NC and APS-G groups. Following treatment with APS, the lipid droplet deposition of the APS-G group decreased, as compared with the CUGFR group, and the tissue samples exhibited lipid droplet fusion (Fig. 4).

TEM examination of the liver samples (Fig. 5) demonstrated that damaged mitochondria and lipid droplets were present in the hepatocytes of the CUGFR (Fig. 5C and D) and APS-G groups (Fig. 5E and F). The groups also exhibited clusters of damaged mitochondria, multi lamellar bodies, and large lipid droplets joined to form large lipid droplets and surrounded by autophagosomal membranes. In the CUGFR group, numerous lipid droplets were present, including fused lipid droplets, and the number of mitochondria and endoplasmic reticula was decreased in the cells. In addition, the cellular structure was severely impaired. Furthermore, the nuclear membranes were not continuous, exhibiting visible holes.

Following treatment with APS, the number of lipid droplets decreased in the APS-G group (Fig. 5E and F), however numerous impaired mitochondria and endoplasmic reticula...
remained visible, and the liver cells exhibited expanded endoplasmic reticula and mitochondrial coagulation-associated degeneration. The cell membranes were thick, but continuous (Fig. 5).

Following treatment with APS, the serum FGF21 levels markedly decreased in the CUGFR group, whereas the serum levels of FGF21 in the APS-G group significantly increased compared with the CUGFR group (Fig. 6; P<0.05).
Table I. Serum protein levels and HOMA-IR following treatment with APS.

| Item              | NC       | CUGFR    | APS-G    | P1    | P2    |
|-------------------|----------|----------|----------|-------|-------|
| TG, mmol/l        | 0.76±0.13| 0.81±0.27| 0.70±0.33| 0.504 | 0.521 |
| TC, mmol/l        | 1.46±0.41| 1.54±0.29| 1.75±0.46| 0.787 | 0.345 |
| LDL, mmol/l       | 0.19±0.12| 0.5±0.20a| 0.72±0.28| 0.0016| 0.185 |
| HDL, mmol/l       | 1.13±0.40| 0.92±0.10| 0.87±0.20| 0.245 | 0.59  |
| FFA, mmol/l       | 0.46±0.11| 0.74±0.08b| 0.44±0.14| 0.038 | 0.126 |
| ALT, U/l          | 37.75±23.2| 58±26.10b| 63.75±31.76| 0.041 | 0.419 |
| FBG, mmol/l       | 6.58±1.19| 6.0±0.37 | 6.0±0.51 | 0.279 | 1.0   |
| Fins, µIU/ml      | 5.82±2.24| 12.1±5.49b| 7.11±2.09| 0.024 | 0.089 |
| HOMA-IR           | 1.43±0.67| 3.02±0.87b| 1.83±0.42| 0.026 | 0.049 |
| HBA1C, %          | 3.85±0.11| 3.86±0.10| 3.87±0.14| 0.85  | 0.88  |

P1, CUGFR vs. NC; P2, APS-G vs. CUGFR. *P<0.01 and **P<0.05 vs. NC group; and ***P<0.05 vs. CUGFR. APS, Astragalus polysaccharide; HOMA-IR, homeostasis model assessment of insulin resistance; NC, normal control group; CUGFR, catch-up growth group; APS-G, APS-treated group; HBA1C, glycated hemoglobin; ALT, alanine aminotransferase; TG, thyroglobulin; TC, total cholesterol; LDL, low-density lipoprotein; HDL, high-density lipoprotein; FFA, free fatty acid; FBG, fibrinogen; Fins, fasting insulin.

Table II. HBA1C, serum lipid and ALT levels.

| Item   | NC       | CUGFR    | P-value |
|--------|----------|----------|---------|
| TG, mM/l | 1.0±0.22 | 1.49±0.45b | 0.013   |
| TC, mM/l | 2.17±0.48 | 2.77±0.71b | 0.05    |
| LDL, mM/l | 0.44±0.21 | 1.02±0.43a | 0.002   |
| HDL, mM/l | 0.77±0.28 | 0.48±0.18b | 0.012   |
| FFA, mM/l | 1.3±0.28  | 1.63±0.95  | 0.35    |
| ALT, µM/l | 47±7.13   | 48.46±13.6 | 0.85    |
| HBA1C, % | 3.99±0.09 | 4.18±0.13b | 0.0039  |

*P<0.01 and **P<0.05 vs. the NC group. HBA1C, glycated hemoglobin; ALT, alanine aminotransferase; NC, normal control group. CUGFR, catch-up growth group; TG, thyroglobulin; TC, total cholesterol; LDL, low-density lipoprotein; HDL, high-density lipoprotein; FFA, free fatty acid.

Table III. Liver index and visceral fat content ratio of the rats.

| Item       | NC       | CUGFR    | APS-G    | P1    | P2    |
|------------|----------|----------|----------|-------|-------|
| Weight, g  | 774.5±39.2| 786±48.33| 697.8±67.7| 0.101 | 0.019 |
| Liver weight, g | 17.14±1.78 | 31.67±2.47b| 27.49±4.05b| <0.001| 0.046 |
| Liver index | 2.31±0.25 | 4.03±0.17b| 3.94±0.41| 0.001 | 0.629 |
| E-P-fat, g  | 31.94±6.41| 42.55±10.62a| 39.36±13.26| 0.038 | 0.638 |
| V-fat, weight % | 4.52±0.53 | 5.75±0.99a | 5.19±1.35 | 0.018 | 0.446 |

P1, CUGFR vs. the NC group; P2, APS-G vs. CUGFR. *P<0.05 and **P<0.01 vs. the NC group; P*P<0.01 vs. CUGFR. NC, normal control group; CUGFR, catch-up growth group; APS, Astragalus polysaccharide; APS-G, APS-treated group; E-P, epididymal plus perirenal; V, visceral.

Relative mRNA expression levels of PGC-1α and PPARα, and protein expression levels of SIRT1, FGF21 and NF-κB in the liver. The expression levels of PGC-1α markedly increased in the CUGFR group when compared with the NC group, and were marginally increased by treatment with APS (Fig. 7A). The expression levels of PPARα in the CUGFR group were significantly lower as compared with the NC group, and these expression levels increased following treatment with APS (Fig. 7B; P<0.05). In the CUGFR group, the protein expression levels of FGF21 and SIRT1 were significantly lower than the NC group (Fig. 7C and D P<0.05), whereas the expression levels of NF-κB were significantly increased. In the APS-G group, the previously low expression levels of FGF21 and SIRT1 were increased (with the difference in SIRT1 being
significant compared with the CUGFR group; P<0.05). APS treatment appeared to exert no significant effect on the activity levels of NF-κB, where the expression levels of NF-κB had previously been significantly increased in the CUGFR group, as compared with the NC group (Fig. 7E and F).

Discussion

Catch-up growth may lead to non-alcoholic fatty liver disease (NAFLD), and accumulating evidence suggests that NAFLD is strongly associated with insulin resistance, which is an important factor in the development of T2DM (13). FGF21 regulates hepatic glycolipid metabolism and interacts with PPARα (14,15), and may serve as a biological target in NAFLD to improve hepatic insulin resistance. SIRT1 is a type of histone deacetylase that regulates NAD+ (16), and regulates certain transcription factors associated with glycolipid metabolism, including PGC-1α and PPARα (17,18). Therefore, the present study hypothesized that the SIRT1-PGC-1α/PPARα-FGF21 signaling pathway may be significant in the pathogenesis of hepatic insulin resistance under catch-up growth. Following diet restriction, the expression levels of SIRT1, PGC-1α and PPARα increased significantly, and the levels of hepatic glucose and fatty acid B oxidation increased significantly (19‑21). In a previous study, obese mice with genetically-engineered reduced levels of PGC-1α exhibited improved whole-body insulin sensitivity with increased levels of hepatic and circulating FGF21 (22). In a model of hepatic insulin resistance, glucose absorption was decreased due to the reduction of glucose transporter 1 and the phosphorylation of extracellular signal-regulated kinases 1/2, which were increased by FGF21 (23,24).
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The experimental rat model in the present study reflected the status of catch-up growth under a high fat diet following a period of calorie restriction. The results demonstrated that the expression levels of PGC-1α only marginally increased, whereas the expression levels of PPARα decreased significantly. Notably, the hepatic protein expression levels of SIRT1 and FGF21 were significantly decreased, with serum expression levels of FGF21 being marginally decreased. Simple variations in the diet, namely a restricted diet followed by a high-fat diet, appeared to result in a catch-up growth effect with significant decreases in SIRT1 and PPARα, while PGC-1alpha increased slightly. These results demonstrated that the effects on the expression of SIRT1 and its downstream factors (PGC-1α/PPARα) were more complex than a simple response to high-energy diet following calorie restriction, as the interactions between these factors became influenced by the increased energy levels and onset of catch-up growth. Furthermore, the results of the present study demonstrated that the insulin resistance index, liver index and visceral fat content ratio of the rats significantly increased in the catch-up growth group on the high-energy diet, with liver H&E staining and TEM indicating the presence of NAFLD pathological changes, which induce hepatic insulin resistance, as well as abnormal glucose and lipid metabolism. In addition, previous studies demonstrated that liver inflammation may cause oxidative stress, resulting in hepatic insulin resistance and NAFLD (25,26). The present study demonstrated that the protein expression levels of NF-κB in the CUGFR rat livers were significantly increased, when compared with the NC group, which indicated that inflammation contributed to the pathogenesis of NAFLD during the catch-up growth of rats. However, the underlying mechanism of the association.
between SIRT1-PGC-1α/PPARα-FGF21 and NF-κB remains unclear and requires further investigation.

Astragalus is a commonly administered type of traditional Chinese medicine and APS is a type of water-soluble polysaccharide that is purified from astragalus. APS exhibits high biological activity (27), regulates inflammation (28,29) and protects the liver (30,31). APS is able to increase insulin sensitivity and lower blood glucose levels by decreasing endoplasmic reticulum stress in liver of patients with T2DM (32). APS treatment reduces the protein expression and activity levels of hepatic glycogen synthase kinase (GSK)3β, promotes hepatic insulin signal transduction, and decreases insulin resistance in the liver of diabetic KKAy mice (33). Furthermore, APS promotes the expression of myocardial PPARα, and increases the myocardial lipid metabolism of diabetic hamsters (12). However, to the best of our knowledge, no previous study has observed the effects of APS on the SIRT1-PGC-1α/PPARα-FGF21 signaling pathway in the liver of catch-up growth rats.

The results of the present study demonstrate that following treatment with APS, liver cells stained with H&E and observed by TEM exhibited markedly decreased lipid droplets, and the mitochondria, endoplasmic reticula and cell membrane ultrastructures were improved, as compared with those of the CUGFR group. In addition, the expression level of PPARα was markedly increased and the change in PGC-1α expression was not statistically significant. Furthermore, the protein expression levels of SIRT1 and FGF21 were significantly increased. The index of insulin resistance decreased, although not significantly. These results suggest that APS may counteract the pathology of NAFLD, by acting upon liver endoplasmic reticula, and reducing the protein expression and activity levels of hepatic GSK3β. Furthermore, APS may prevent NAFLD by increasing insulin sensitivity and promoting hepatic insulin signal transduction (32,33). The results of the present study also suggest that APS may improve hepatic glycolipid metabolism and suppress insulin resistance via modulation of SIRT1-PGC-1α/PPARα-FGF21. PGC-1α enhances the effects of PPARα through a conformational change, however, the present study demonstrated that APS did not increase the expression levels of PGC-1α and, therefore, there must be another mechanism that results in the increased protein expression levels of PPARα and FGF21. Previous studies have reported that SIRT1 increases the expression levels of PPARα (34,35) and the present study demonstrated that APS increases the protein expression levels of SIRT1. Therefore, the increased expression levels of PPARα and FGF21 in the liver may be due to an increase in PPARα deacetylation levels in liver cells; however, this hypothesis requires further investigation. Whether the effects of APS treatment act via the various polysaccharide groups, or via the β-D-(1→3) galactan or β-D-(1→6) activity groups of galactooligosaccharide side-chains also requires further investigation. In addition, a previous study on the effects of APS on blood glucose in rats demonstrated that APS treatment for 4 weeks exhibited no significant effect on the blood glucose levels of diabetic rats; however, following treatment with APS for 8 weeks, the decreasing effects on blood glucose levels became apparent. Furthermore, no effects were observed in the glucose levels of normal rats (36). Therefore, the results of the present study on the effects of APS on the morphological pathological changes of the liver, and on glucose and lipid metabolism further suggest that the active components of astragalus exerted significant effects on HOMA-IR and fasting insulin levels, although it did not affect hepatic insulin resistance. A study of longer duration would be required to observe the effects of APS on pathological and morphological changes in the liver, as well as on glucose and lipid metabolism. In addition various APS doses and purity levels require investigation to observe their effects on insulin resistance, glycolipid metabolism and NAFLD, to elucidate the ranges and determine guidelines for APS administration to provide early intervention strategies for the prevention of T2DM.

In conclusion, the suppressive effects of APS on liver acetylation levels and on associated glycolipid metabolism molecules contribute to decreasing insulin resistance. However, the mechanism underlying these effects remains to be elucidated, and requires further investigation. Early treatment with APS over longer time periods may provide a novel, safe and effective therapeutic strategy for the treatment of T2DM.

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