Insulin-Related Liver Pathways and the Therapeutic Effects of Aerobic Training, Green Coffee, and Chlorogenic Acid Supplementation in Prediabetic Mice

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Background. The liver controls blood glucose levels via regulation of anabolic (glycogen synthesis and gluconeogenesis) and catabolic (glycolysis and glycogenolysis) processes through activation of the PI3K-AKT signalling pathway. The aim of this study was to assess the effect of aerobic training, green coffee, and chlorogenic acid supplementation on glucose metabolism-regulating pathways in prediabetic mice.

Methods. C57BL/6 mice were exposed to a high-fat diet and physical activity limitation to induce a state of prediabetes. After 12 weeks, mice were fed a high-fat diet compared to the control mice. The prediabetic mice were further treated with either green coffee, chlorogenic acid, or training or combinations of the same for 10 weeks. At the end of the experimental period, metabolic data (FBG, GTT, HOMA for IR, plasma level of insulin from systematic, AST, and ALT assessed into blood), histopathologic, and analysis of gene and protein expressions were obtained for target tissues.

Results. Training along with green coffee and chlorogenic acid supplementation improved complications of prediabetes including weight gain and elevated fasting blood glucose and plasma insulin levels. These effects were associated with the changes in mRNA levels of genes important in hepatic glycogen synthesis (GYS2), glucogenesis (PCK and G6PC2), and glycolysis (GK, PK, and PFKL). Conclusion. The training in conjunction with green coffee or chlorogenic acid is effective in the prevention of prediabetes in mice. As these interventions are relatively inexpensive and safe application to individuals with prediabetes appears warranted.

1. Introduction

Prediabetes (PD) is a metabolic disorder characterized by higher than normal and less than diabetic levels of blood glucose [1, 2]. Impaired fasting glucose (IFG) and glucose tolerance (IGT) tests assist in making the diagnosis [3, 4]. In prediabetic patients, normal levels of fasting blood glucose and glucose tolerance test are 100-125 mg/dl and 145-199 mg/dl, respectively. Prediabetics have 3-10 times the risk of progressing to type 2 diabetes mellitus (T2DM) compared to nonprediabetics [5]. The primary cause of PD is insulin resistance in the muscle, liver, and adipose tissues. This pro-
Progressive resistance to insulin leads to degradation in pancreatic beta-cell function and decreased insulin secretion and T2DM. Obesity is considered a primary factor accounting for insulin resistance. This is important, as current lifestyle factors such as sedentary routines and consumption of high-calorie diets promote weight gain and obesity [6, 7].

Insulin is secreted from beta cells of the pancreas into the portal vein. Thus, unlike other organs that receive insulin from systemic blood circulation, the liver is exposed to 2-3 times more insulin to suppress glycogenolysis and gluconeogenesis. The importance of insulin on whole-body glucose homeostasis in the liver is well documented [8, 9]. In hepatocytes, insulin acts through the PI3K/AKT signalling pathway. In hepatocytes, insulin binds to its receptors resulting in IRS1/2 phosphorylation activating the downstream target PI3K/AKT. Glycogen synthase kinase-3 (GSK3), transcription factor forkhead box protein O1 (FOXO1), and glucokinase (GK) are AKT pathway substrates [10, 11]. On the one hand,
insulin resistance and obesity, hepatic IRS1/2 undergoes serine phosphorylation to limit AKT activation and disrupt glycogen synthesis, thus promoting glycogenolysis and gluconeogenesis [12]. On the other hand, glycogen is also disrupted by insulin resistance [12]. Glycogen synthase 2 (GYS2), as the main enzyme in glycogen synthesis, is regulated by insulin through control of transcription and cytoplasmic to nucleus transit [9] (Figure 1).

Figure 2: HPLC chromatogram of CGAs in green coffee bean extract (BSK). (a) the percentages of chlorogenic acid and caffeine in green coffee. (b) the biochemical characteristics of chlorogenic acid and caffeine in green coffee.

2. Material and Methods

2.1. In Vivo Experiments. This study was approved by the Ethics Committee of Royan Institute (ethics code: IR.A-CECR.ROYAN.REC.1399.075). All animals were treated according to the Animal Ethics Committee of Royan Institute’s recommendations.

Four-week-old male C57BL6 mice, weighing 12–14 g, were used in this study. Mice were housed in pathogen-free barrier facilities under controlled temperature (23°C ± 3°C) and humidity with a 12-h light/dark cycle. Access to food and water was ad libitum. After 1 week of adaptation, the mice were randomly divided into two groups: control (Ctrl; n = 5) and high-fat diet (HFD; n = 30) mice. Mice in the Ctrl group received a standard diet (containing carbohydrates 47.7%, fat 12.5%, and protein 20.5%) while mice in the HFD group received a diet containing 25% carbohydrates, 60% fat, and 15% protein for 12 weeks. Particularly, the fat ingredients in HF diets are saturated. After ensuring the emergence of PD, the HFD group was divided into six subgroups (n = 5): prediabetic mice (PD group), prediabetic mice treated with green coffee (200 mg/kg, PD/GC group), prediabetic mice treated with chlorogenic acid (100 mg/kg, PD/CGA), prediabetic mice which underwent training (PD/EX), prediabetic mice treated with green coffee + training (PD/EX. GC), and prediabetic mice treated with chlorogenic acid + training (PD/EX. CGA) [21–24].

Green coffee tablets were purchased from BSK (Zist Takhmir, Tehran-Iran). These tablets are a natural product made from 400 mg of standardized green coffee bean extract powder containing 2% caffeine and 50% chlorogenic acid, which these main substates analyzed by HPLC. Briefly, HPLC analysis of CGAs in green coffee bean extract (BSK) was performed using the HPLC-diode array detector gradient system (Agilent 1090 series) (Figure 2). Green coffee and CGA were administered as a gavage supplement, three times per week for 10 weeks.

During the experiment period, the mice were weighed every 7 days. At the end of the experimental period, animals were euthanized after a 12-h overnight fast under xylazine and ketamine anaesthesia. Serum and liver tissues were immediately stored at −80°C for further analysis. Figure 1 demonstrates the flowchart of the study.
2.2. Biochemical Analyses. Fasting blood glucose (FBG) and glucose tolerance tests (GTT) were performed at the end of the 12th and 22nd week of the intervention using a glucometer (Alpha TRAK glucometer, Zoetis, US) from a tail prick. For GTT, mice were fasted for 6 hours and then gavaged with 200 \( \mu l \) glucose solution. Blood glucose was measured at 0, 30, 60, 90, and 120 min later. The plasma level of insulin was determined with an Ultra-Sensitive Mouse Insulin ELISA Kit (ALPCO 80-INSMS-E01, Keewaydin Drive, USA) according to the manufacturer’s instructions. Plasma aspartate aminotransferase (AST) and alanine aminotransferase (ALT) levels were measured using Mouse AST and ALT ELISA Kits (ELISA Kit CSB-E12649m, UK and ELISA Kit- E01, Keewaydin Drive, USA) according to the manufacturer’s protocol.

2.3. Training Intervention. Aerobic training was carried out for 10 weeks (5 days/week). Briefly, all mice were acclimated to running on a treadmill (10 min, using different speeds) for 1 week. On day 1, an electric shock is applied to make the mice start running, after which they will run spontaneously. In other words, this method provides the sole external motivator through grid and mice quickly acquire that make maintain some distance from the grid when running [25]. Aerobic training was performed on a treadmill at low to moderate intensity (50%-60% MAV), 45 minutes per day for 5 days per week for 10 weeks, the exact time for training was between 10 p.m. and 11 p.m. The experiment was carried out in a quiet, well-ventilated room with low humidity at a temperature of 18 ± 2°C. The sedentary mice in the control group were treated similar to the training mice except they were not engaged in regular running. Each training session included 3 minutes of warm-up, 40 minutes of training, and 2 minutes of recovery. The initial training intensity was 15 m/min. The intensity was increased 2 m/minute every 2 weeks until it reached 23 m/min on the final week. Thus, the first and second week used an intensity of 15 m/min, the third and fourth week an intensity of 17 m/min, the fifth and sixth week an intensity of 19 m/min, the seventh and eighth week an intensity of 21 m/min, and the ninth and tenth week an intensity 23 m/min [26].

2.4. RT-qPCR Analysis. The total RNA isolation from the liver tissue was performed using the TRIzol reagent (Thermo Fisher Scientific, Waltham, MA, USA). In order to remove contaminating genomic DNA, samples were treated with DNase I (TaKaRa, Japan). mRNA was reverse transcribed with 1 \( \mu g \) of total RNA using the cDNA synthesis kit (Biotechrabbit, Germany, Berlin) according to the manufacturer’s instruction. RT-qPCR was performed with SYBR Green (TaKaRa, Japan) using an Applied Biosystems real-time PCR thermal cycler (Thermo Fisher Scientific, Waltham, MA, USA). The evaluation of gene expression was carried out according to the 2\(^{-}\Delta\Delta CT\) method. Accordingly, the relative gene expression was calculated according to 18s rRNA as an internal control. Primer pairs were designed by the Beacon designer (Version 7.2, USA) and purchased from Micro-gene (Korea). The primer sequences are shown in Table 1.

2.5. Western Blot Analysis. Proteins were extracted from the tissue samples using the TRI reagent, according to the manufacturer’s protocol. Equal amounts of each protein sample (30 \( \mu g \)) were subjected to SDS PAGE and transferred to PVDF membranes. Membranes were blocked with a blocking buffer containing 10% skim milk (Millipore, USA) and 5% TBST. The membranes were probed with primary antibodies including anti-Akt antibody (1: 2000, ab30471, Elabscience, USA), anti-p-Akt (Thr 308) antibody (1: 1000, sc271966, Santa Cruz, USA), anti-\( \beta \) actin antibody (1: 1000, sc477878, Santa Cruz, USA), anti-glycogen synthase antibody (1: 1000, sc390391, Santa Cruz, USA), and anti-phospho-glycogen synthase antibody (Ser641) antibody (1: 1000, Cell Signaling, USA) for 1.5 hours at room temperature. Subsequently, the membranes were incubated for 1 hour at room temperature with an appropriate secondary antibody: horseradish peroxidase-(HRP-) conjugated goat antimouse IgG (1: 5000, Dako, P0447) or HRP-conjugated goat antirabbit IgG (1: 16000, Santa Cruz, SC2301). Bands were visualized by an Amer sham ECL Advance Western Blotting Detection Kit (GE healthcare). The activity of AKT protein was measured by comparing the level of phosphorylated AKT with the level of nonphosphorylated AKT. The ImageJ software (National Institutes of Health, Bethesda, MD, USA) was utilized for quantification of the intensity band.

2.6. Histological Studies. Immediately after the human euthanasia of the mice, the livers were removed, selected at random, sliced into small pieces, and were fixed in 10%
buffered formalin and embedded in paraffin. Fixed tissues were then cut into 5-μm thickness tissue sections. After deparaffinization and hydration, sections were stained with hematoxylin and eosin (H&E). The periodic acid-Schiff (PAS) was used to stain glycogen in the liver sections. Slides were observed using a light microscope.

2.7. Statistical Analysis. The statistical analyses were carried out using GraphPad Prism 8.4.3 software (GraphPad Software, San Diego, CA, USA). The paired sample t-test was performed to evaluate the prediabetic group compared to the control group. One-way analysis of variance (ANOVA) and Tukey’s post-hoc test were used to determine statistical differences.
significance between all treatment groups. *p-value < 0.05 depicts significant difference between samples. All experimental results are presented as mean ± SD.

3. Results

3.1. HFD Promotes Prediabetes in Mice. The animals given the HFD showed a significant increase in body weight and the percentage of weight gain (Figures 3(b) and 3(c)). Additionally, the animals demonstrated increases in FBS, GTT (Figures 3(d) and 3(e)), and plasma insulin levels compared to the mice receiving the ND (Figure 3(f)). Consistent with these changes, the HOMA-IR index was significantly elevated in the HFD-fed mice (Figure 3(g)). The Western analysis of hepatic protein expression demonstrated less AKT protein phosphorylation in organs from the HFD versus the ND group (Figure 3(h)). Interestingly, the liver morphology and histopathology were comparable among groups and not grossly abnormal (data not shown). However, the ratio of liver weight to total bodyweight was greater for the HFD-fed mice (Figures 4(a) and 4(b)). As expected, PAS staining was less in liver samples from the HFD-fed mice (Figures 4(c) and 4(d)). Finally, the analysis of plasma levels of AST and ALT found that the enzymes were significantly higher in mice receiving the HFD compared to the ND-fed animals (Figure 4(e)).

3.2. HFD Is Associated with Changes in mRNA Levels of Genes Involved in Hepatic Glycogen Synthesis, Gluconeogenesis, and Glycolysis. In order to investigate the possible causes of hyperglycemia and reduced liver glycogen content, hepatic mRNA levels of genes associated with glycogen synthesis, glycolysis, and gluconeogenesis were examined. The livers from HFD-fed mice showed reduced expression of Gys2
mRNA (Figure 5(a)). The Western analysis of hepatic protein expression demonstrated less GYS2 protein phosphorylation in organs from the HFD versus the ND group (Figure 5(b)).

The mRNA levels of two gluconeogenesis enzymes, phosphoenolpyruvate carboxykinase (Pck1 = Peck) and glucose-6-phosphatase catalytic subunit 2 (G6pc2), were increased in mice on the HFD compared to the mice on the ND, although in the case of Pck1 was unchanged. (Figures 5(c) and 5(d)). Additionally, in the livers from the HFD mice, mRNA levels of glucokinase (Gk), pyruvate kinase (Pk), and (Pfk) phosphofructokinase were decreased compared to the controls (Figures 5(e), 5(f), and 5(g)).

3.3. Training, Green Coffee, and Chlorogenic Acid Mitigated PD-Associated Changes in HFD-Fed Mice. After induction of PD, the mice were treated with training (EX), green coffee (GC), chlorogenic acid (CGA), or a combination of these factors (EX+GC and EX+CGA). The GC used in this study contained 50% CGA and less than 2% caffeine, as revealed by HPLC (Figure 2). In the groups receiving GC and CGA, no weight loss was observed compared to the control PD group. In contrast, the mice that underwent EX regardless of other treatments showed significant weight loss compared to the control PD group (Figures 6(a) and 6(b)). All treatments improved blood glucose levels (Figure 6(c)) and glucose tolerance in the PD mice (Figure 6(d)). Moreover, all treatments resulted in decreased plasma insulin levels compared to the controls (Figure 6(e)). Importantly, EX was found to have the greatest effect on insulin levels. In keeping with these findings, the HOMA-IR index was lower in the treated PD mice compared to the untreated (Figure 6(f)). The accumulated protein level of hepatic AKT phosphorylation was higher in samples from all the intervention groups compared to the PD group (Figure 6(g)). This corresponded to the decreased liver to total body weight ratios in the treated compared to untreated PD mice. However, this trend was statically significant in the whole group-treated animals (Figure 7(a)). PAS staining in liver sections from

Figure 5: HFD is associated with changes in hepatic metabolic enzyme mRNA levels. (a) RT-qPCR analysis of Gys2 in the livers from the Ctrl and HFD mice. (b) Immunoblotting analyses of p-GYS2 and total GYS2 protein in the livers from the Ctrl and HFD groups, (c) RT-qPCR analysis of G6pc2, and (d) Peck mRNA in the livers from the Ctrl and HFD mice. (e–g) RT-qPCR analysis of Pk, Pfk, and Gk mRNA in the livers from the Ctrl and HFD mice. All values are presented as mean ± SD. * = p < 0.05, ** = p < 0.01, and *** = p < 0.001 indicate significant statistical differences in the HFD compared to Ctrl mice.
Figure 6: Continued.
treated PD mice was also judged less compared to the controls (Figures 7(b) and 7(c)). Similarly, serum levels of AST and ALT were decreased in all intervention groups (Figures 7(d) and 7(e)).

3.4. Training, Green Coffee, and Chlorogenic Acid Treatment Alter Hepatic mRNA Levels of Genes Associated with Glycogen Synthesis, Gluconeogenesis, and Glycolysis. In all treatment groups, hepatic mRNA levels of Gys2 and levels of phosphorylated Gys2 were increased (Figures 8(a) and 8(b)) compared to the untreated PD group. mRNA levels of Pck were increased in the EX and EX+GC-treated PD mice versus the controls (Figure 8(c)). As well, EX was associated with increased hepatic G6pc2 mRNA, whereas in animals treated with GC or EX+CGA groups mRNA levels were decreased (Figure 8(d)). Hepatic mRNA levels Pfk1, Gk, and Pk were found to trend up in PD mice regardless of the intervention compared to the controls. However, the difference was only significant in the EX-treated animals (Figures 8(e) and 8(g)). mRNA levels of Pfk1 were significantly increased in the CGA-, EX-, and EX+GC-treated groups (Figure 8(f)).

4. Discussion
In the current study, the effects of training in combination with green coffee and chlorogenic acid in the HFD-fed PD mice were evaluated. To the best of our knowledge, this is the first published study to address this. Consistent with the findings in people, the HFD-fed mice displayed
metabolic dysregulation including elevated FBG, insulin and AST/ALT levels, abnormal glucose tolerance, and increased HOMA index. Additionally, increased liver weight, decreased glycogen levels, and AKT phosphorylation were observed. Furthermore, the HFD-fed mice showed disruption in hepatic glycogen synthesis as evinced by reduced expression of Gys2, increased gluconeogenesis via increased Pck and G6pc2, and increased glycolysis via decreased expression of Gk and Pk.

Therapeutic intervention with training, green coffee, or chlorogenic acid found that training was most efficacious at resisting HFD-mediated metabolic dysregulation including glucose balance, glycogen accumulation, and body and liver weight changes. This is consistent with clinical data that identified constant training as having an impact on metabolic syndrome and diabetes [27].

In mice, HFD is an accepted means of inducing metabolic imbalance (Xu, D., Jiang et al. in 2019, Asare-Bediako et al. in 2020 and Mu, H. N) [28–30]. EX, GC, and CGA independently reduce FBS and improved insulin resistance. We found that while EX caused weight loss in the PD mice, GC and CGA alone had no effect, as we found, others reported that EX promoted weight loss and improvement in body composition. However, there effects of GC and
Figure 8: GC, CGA, EX, EX. GC, and EX.CGA treatment alters mRNA expression of multiple metabolic genes in livers from PD mice. (a) RT-qPCR analysis of Gys2 mRNA levels in the liver from PD and all treated animal groups. (b) Immunoblot of protein levels of p-GYS2 and total GYS2 in the liver from PD and all treated groups. (c) RT-qPCR analysis of G6pc2 and (d) Pck mRNA in the livers from PD and all treated animal groups. (e–g) RT-qPCR analysis for Pk, Pfkl, and Gk mRNA levels in the livers from PD and all treated groups. All values are presented as mean ± SD. * = p < 0.05, ** = p < 0.01, and *** = p < 0.001 indicate significant statistical differences in the treated compared to PD mice.
CGA on weight loss are controversial [17, 20, 31–33]. It is possible that variation in study design and dosages explain variation in reported outcomes.

The liver controls blood glucose levels via the regulation of anabolic (glycogen synthesis and gluconeogenesis) and catabolic (glycolysis and glycogenolysis) pathways. By activation of PI3K-AKT signalling pathway, insulin causes progression of glycogen synthesis and gluconeogenesis and inhibits glycolysis and glycogenolysis. In PD and T2D, liver glycogen storage is dysregulated. Conversion of glucose to glycogen is performed by GYS2. In a feedback manner, GYS2 is activated is dysregulated. Conversion of glucose to glycogen is performed by GYS2. In a feedback manner, GYS2 is activated.

In summary, the results of this study demonstrated that aerobic training in conjunction with green coffee lowered blood glucose and mitigated metabolic changes associated with PD in mice. These data support further study looking at long-term effects with a goal of possible translational applications.

**Abbreviations**

ADA: American Diabetes Association  
AKT: Protein kinase B  
ALT: Alanine aminotransferase  
AST: Aspartate aminotransferase  
BSK: Bonyan Salamat Kasra  
CHA: Chlorogenic acid  
EX: Training training  
FBG: Fasting blood glucose  
FOXO1: Forkhead box protein O1  
G6PC2: Glucose-6-phosphatase 2  
GC: Green coffee  
GK: Glucokinase  
GSK3: Glycogen synthase kinase 3  
GTT: Glucose tolerance test  
GYS2: Glycogen synthase 2  
H&E: Hematoxylin and eosin  
HOMA: Homeostatic model assessment of insulin resistance  
IGF: Impaired fasting glucose  
IGT: Impaired glucose tolerance  
IRS1/2: Insulin receptor substrate 1  
MAV: Micro aerial vehicle  
PAS: Periodic acid-Schiff  
PD: Prediabetes  
PCK: Phosphoenolpyruvate carboxykinase  
PFKL: Phosphofructokinase, liver type  
PDK: Pyruvate kinase  
T2DM: Diabetes mellitus type 2.

**Data Availability**

Data are available on request.

**Additional Points**

*New & Noteworthy.* (i) Training, green coffee, chlorogenic acid, and the combination of training with green coffee and chlorogenic acid can improve the symptoms and complications of prediabetes by regulating the glycogen synthesis mechanism by activating the signalling pathway of IRS1, PI3K, AKT, GSK3, and GYS2. The results were somewhat effective. The results were so effective that blood sugar, plasma insulin, and liver enzymes were in the normal range. (ii) Training, green coffee, and chlorogenic acid can improve prediabetic conditions by activating genes involved in glycolysis mechanisms, such as PK, PFKL, and GK, and prevent the disease from progressing to type 2 diabetes. (iii) Green coffee and a combination of training and chlorogenic acid, by controlling the genes involved in the mechanism of...
gluconeogenesis, interfere with the activity of this signalling pathway and thus can prevent the production of glucose and glucose accumulation in the blood.

**Ethical Approval**

The approval of mouse usage in this study was obtained by the Ethics Committee of Royan Institute.

**Conflicts of Interest**

The authors declare that they have no conflicts of interest to declare.

**Authors’ Contributions**

M.A. contributed to the study design, data collection, analyses, data mining, data interpretation, and manuscript writing. S.M.M. contributed to the study design, data collection, data analysis, data interpretation, and manuscript writing. K.G.H. contributed to the data interpretation. Z. S. contributed to the data interpretation, technical assistance, manuscript writing, and final approval of the manuscript. F.K. contributed to the study design and data collection. S.S.H. contributed to the data collection. M.H.S. is the liver pathologist. P. R. contributed to the manuscript writing. M.H.N.E. contributed to the data analysis and data interpretation. All authors read and approved the final manuscript.

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