Aqueous extract of post-fermented tea reverts the hepatic steatosis of hyperlipidemia rat by regulating the lipogenic genes expression and hepatic fatty acid composition

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Abstract

Background: Post-fermented tea has been used for the prevention of metabolic syndrome in Western China. Present study reports the biochemical mechanism of lipid-lowering effects of Jing-wei fu tea (JWFT), a variety of post-fermented tea on high-fat diet-induced hyperglycemia and obesity in rats.

Methods: Aqueous extract of JWFT was prepared by putting them in boiling water, and then concentrated under reduced pressure. The major compounds of JWFT were determined by high performance liquid chromatography (HPLC). High-fat diet fed rats were orally administered different doses of JWFT aqueous extract (0.5, 1.0 and 2.0 g/kg) for four weeks. At the end of this experiment, hepatic lipids, serum leptin and lipids levels were determined using enzyme-linked immunosorbent assay (ELISA). The hepatic fatty acid composition was analyzed using gas chromatography mass (GC-MS). The relative expression of lipids metabolism genes was analyzed using reverse transcriptase-polymerase chain reaction (RT-PCR).

Results: The results showed that JWFT inhibited the increase in the body weight, abdominal adipose weight, serum lipids and hepatic lipids, and decreased serum leptin levels of high-fat diet fed rats. JWFT normalized hepatic fatty acid composition of hyperlipidemia rats by up-regulating hepatic acetyl-CoA carboxylase (ACC), stearoyl-CoA desaturase 1 (SCD1) and fatty acid synthase (FAS) gene expression, and down-regulating carnitine palmitoyl transferase-1 (CPT1). Furthermore, the results showed that JWFT inhibited the absorption of lipids.

Conclusion: JWFT could mitigate the obesity-induced hepatic steatosis by regulating hepatic lipogenesis and lipolysis.

Keywords: Post-fermented tea, Lipids, Absorption, Lipogenic genes, Leptin

Background

Obesity is a balance disruption of energy intake and expenditure [1]. As a typical symptom of metabolic syndrome, hepatic steatosis was usually induced by high-fat and high carbohydrate diet [2]. Liver is the main organ for lipid uptake, de novo synthesis and fatty acid lipolysis [3]. Some lipogensis and lipolysis enzymes have been known to be involved in the lipids synthesis and metabolism. For example, acetyl-CoA carboxylase (ACC) converts the acetyl-CoA to malonyl-CoA, which is the direct carbon donor for de novo synthesis of fatty acid. Fatty acid synthase (FAS) organizes the structural skeleton of fatty acid with malonyl-CoA. Stearoyl-CoA desaturase 1 (SCD1) is a critical rate-limiting enzyme of de novo synthesis of monounsaturated fatty acids (MUFAs) [4]. Furthermore, increase in the carnitine palmitoyltransferase 1 (CPT1) substantially reduced the hepatic triglycerides level because it increased the fatty acid oxidation [5]. While the deficiency of SCD1 increased the oxidation of fatty acid, insulin sensitivity and resistance to diet-induced obesity and liver steatosis [6].

Many traditional herbal materials have attracted much attention because their various health benefits [7,8]. There has been a very long history of tea consumption as...
beverage or herbal medicine. Tea contains high contents of polyphenols and caffeine. Its biological activities have been widely studied with respect to cancer prevention, anti-inflammatory, anti-hyperlipidemia and antimicrobial effects [9,10]. Recently, post-fermented teas including pu-erh tea and fuzhuan brick tea have been reported with beneficial efficacy for metabolic syndrome in clinical intervention trials and animal models [11]. For many years, catechins and their gallate esters were widely reported as the main active compounds of green tea for anti-cancer and anti-obesity. In addition, (−)-epigallocatechin gallate (EGCG) showed the effects of inhibiting lipids absorption and synthesis [12]. But after post-fermentation, these compounds were highly decreased compared with unfermented tea. Therefore, it was suggested the catechins’ metabolite formed in post-fermentation may contribute to the health benefits of post-fermented tea [13].

“Fu” tea is a kind of post-fermentated tea according to the classification of processing method. Its history could be dated back to the Ming Dynasty during 1500 A.D. [14]. Jing-wei fu tea (JWFT) is made of the mature leaves of Camellia sinensis by undergoing an intensified fermentation of Eurotium cristatum [15]. The detailed manufacture process of this tea could be described as panning, rapid pile fermentation, rolling, drying, softening with steam, piling, tea brick pressing, fungal fermentation, and drying in order. The phase of “fungal fermentation” is critical for typical aroma, taste and unique phytochemical profiles.

The aim of this paper is to elucidate JWFT’s function of regulating lipid metabolism and provide a novel understanding on lipid-lowering and anti-obesity activities of tea.

Methods
Materials
The raw material of Jing-wei fu tea is the mature large leaves of Camellia sinensis L. O. Kuntze. Jing-wei fu tea (JWFT) was provided by Cang-shan Co, Ltd. (Xianyang, Shaanxi, China) and formally identified by Professor Xiaochun Wan. Tea leaves were collected from tea plants (Camellia sinensis) in the Hanzhong district of China. To process tea, the mature leaves were dampened and fermented with the Eurotium cristatum under high humidity and temperature. All of the chemical standards used in this study were with the purity of more than 98%.

Preparation and analysis of JWFT extract
Aqueous extract was prepared by boiling minced JWFT powder with distilled water for 40 min at a ratio of 1:5 (w/v) three times. The filtered solution was combined and concentrated at 60°C with rotary evaporator to the required concentration, and then stored at 4°C. The procedures of sample preparation and HPLC analysis referenced published methods [16].

Animals and treatment
Fifty male Sprague–Dawley (SD, 16 weeks old) rats were purchased from the Animal Breeding and research center of Anhui Medical University, China. This study was approved by the Institutional Animal Care and Use Committee of Anhui agricultural university and carried out according to the guidelines of Care and Use of Animals Laboratory. These rats were housed in an environmentally controlled room (22 ± 2°C, 60 ± 5% relative humidity, and 12 hours light/dark cycle) and received food and water ad libitum. At the beginning of study, the body weights of rats were ranged from 160 to 200 g. After one week’s acclimatization, rats were randomly divided into five groups of 10 animals each. Four groups were fed high-fat diet and the fifth group was fed normal control diet (NC). One group of high-fat diet fed rats served as control group (HFC) and was maintained on the high-fat diet alone until the end of the study, while other three groups received high-fat diet and daily oral administrations (20 ml/kg) of JWFT extract for four weeks, at the dose of 0.5, 1.0 and 2.0 g/kg body weight, named as low (LJT), medium (MJT) and high (HJT) dose of JWFT treatment group. After four weeks’ treatment, the animals were fasted for 12 hours, and then blood was taken from the femoral artery into tubes and immediately centrifuged at 6000 rpm at 4°C for 5 min. All the samples were stored at −80°C until analyzed.

Body weight, abdominal fat, serum lipids and leptin analysis
Each rat’s body weight was recorded every week during the whole experiment. At the last day, rats were fasted for 12 h but free access to water, and then were anaesthetized. Blood was drawn from the abdominal aorta and rats were killed by exsanguinations. The intra-abdominal fat (including epididymal, mesenteric, and retroperitoneal fat) were removed and weighed. The body weights and abdominal fat of treatment groups were statistically compared with the HFC group. The levels of cholesterol, triglycerides and high density lipoprotein-cholesterol (HDL-C) were measured using ELISA methods with an automatic analyzer (SPECTRA MAX 190, Molecular Devices, US). Low density lipoprotein-cholesterol (LDL-C) levels were determined using the formula: LDL-C = TC−(1/5TG + HDL-C) according to the published methods [11]. The levels of serum leptin were determined using automatic radioimmunoanalyzer according to the leptin radioimmunoassay kit manufacturer’s (Beijing Northern institute of biotechnology, Beijing, China) directions.
Extraction and measurement of hepatic lipids

Total liver lipid was extracted by homogenizing with chloroform-methanol mixture (2:1, v/v) to a final dilution 20-fold the volume of the tissue sample according to the published method [17]. The extraction was carried out in a tissue tearor for three times. Pooled extract was dried using a nitrogen evaporator. Hepatic total cholesterol (TC) and triglycerides (TG) were determined by ELSIA according to the manufacturer’s directions.

Preparation and analysis of fatty acid methyl ester

Fatty acid methyl ester (FAME) was prepared by a safe rapid mean for methyl esterification with potassium hydroxide-methanol as a reaction solution, and then was extracted with hexane. 200 mg of lipids dissolved in 2 mL hexane and mixed well, then added 0.2 mL KOH-methanol solution (2 M). These tubes were mixed by vortex agitation for 30 sec and centrifuged at 5000 rpm for 5 min. The supernatant containing FAME was decanted and diluted with hexane, and then added 2 g sodium sulfate anhydrous for overnight.

Gas chromatograph-mass spectrometer (GC-MS) analysis of fatty acids composition was performed using Shimadzu GC-MSQP 2010S system (Tokyo, Japan) equipped with a DB-5 capillary column (30 m × 0.32 mm ID) (Agilent, USA). The injection volume was 1 μL and split less. Helium was used as carrier gas and flow rate at 0.85 mL/min. The oven temperature program was from 150°C to 190°C at 18°C/min maintained for 1 min, 190°C to 200°C at 0.5°C/min held for 5 min, 200°C to 230°C at 2°C/min kept for 5 min. The injection and detector (FID) temperatures were 250°C and 280°C respectively. Mass detection was done with electric ionization at 1 keV by using selected-ion monitoring/scan mode for masses in the range of m/z at 50 to 600.

Analysis of the fecal lipids

In the 27th day, tea treatment and high-fat diet control groups’ animals were fasted for 12 h. After that, tea treatment groups were orally administered corresponding dose of JWFT. After 30 min, equal lipids (containing triglyceride and cholesterol) were given to each group treatment groups were orally administered corresponding dose of JWFT. After 30 min, equal lipids (containing triglyceride and cholesterol) were given to each group.

The feces were dried under reduced pressure at 40°C. One gram of dry feces was accurately weighed, and extracted with chloroform and methanol (2:1). The fecal TG and TC levels were determined using the kits from Nanjing Jiancheng Bioengineering Institute (Nanjing, China).

RNA extraction and reverse transcriptase-polymerase chain reaction (RT-PCR)

RNA was extracted from rat liver by homogenization in RNAPrep pure tissue kit (TianGen biotech CO. LTD, Beijing). Purity was determined using the 260/280 ratio on a Nanodrop 2000 spectrophotometer (Thermo Scientific, Wilmington, USA). Samples with a 260/280 ratio of 1.90 or greater were used for cDNA synthesis, using a cDNA reverse transcription kit (Takara, 5 × Primescript RT Master Mix) with Bio-Rad S1000™ Thermal cycler, programmed cycle of 37°C for 15 min, 85°C for 5 sec and 4°C until storage at −80°C.

The primers for SCD1, ACC, CPT1, FAS and 18S housekeeping gene were referenced the published references [18,19]. Primers were ordered from Sangon Biotech (Shanghai) Co., Ltd. RT-PCR was performed using SYBR Green qPCR mastermix (Takara, SYBR* Permix Ex Taq™II) on Bio-Rad CFX 96™ real-time PCR system. The reaction volume was 20 μL. The program had an initial incubation of 95°C for 30 sec, followed by 39 cycles of 95°C for 5 sec and 60°C for 30 sec following 39 cycles, a melt curve was produced by increasing the temperature from 60 to 95°C at a rate of 0.5 per minute. Analysis of data was performed by normalizing the threshold cycle number (Δct) of the gene of interest to the Δct for the housekeeping gene, 18S ribosomal RNA, using the 2ΔΔct method and expression HFC values relative to LJT, MJT, HJT values and expressing in arbitrary units.

Statistical analysis

Results were expressed as means ± SEM, with the number of determinations (n = 10) representing separate experiment. Data were evaluated at a 0.05 level of significance with one-way ANOVA with post-hoc testing by Fisher’s protected least significant differences procedure.

Results

Polyphenol and caffeine contents of JWFT water extracts

As shown in Table 1, the contents of catechins-gallates such as (−)-epigallocatechin gallate (EGCG), (−)-epicatechin gallate (ECG) and (−)-gallocatechin gallate (GCG) were very low in JWFT. Different from ripened pu-erh tea, the content of gallic acid was still very low [16]. These results indicated JWFT’s post-fermentation was not the same as other Chinese dark teas which highly increased the content of gallic acid.

Body weight, abdominal fat weight and serum lipids and leptin levels

The hyperlipidemia rats of HFC group gained more weight during the study than those fed a normal chow diet (p < 0.01). Medium and high dose (1.0, 2.0 g/kg) of JWFT extract exhibited a remarkable decrease in body weight and abdominal fat weight compared with the HFC (Figure 1, p < 0.05, 0.01). Serum TC and TG levels of HFC group were consistently much higher than the NC group after 28 days (p < 0.01). After four weeks’ treatment of high dose of JWFT extract (2.0 g/Kg), serum TC, TG
and LDL-C levels of HJT group were significantly decreased compared with HFC group (p < 0.05).

Hepatic total cholesterol and triglycerides levels
High-fat diet caused the elevation of hepatic total TG and TC levels. JWFT treatment decreased the hepatic total TG and TC levels. As shown in Figure 2, hepatic TC levels were dose-dependently decreased by 15.93%, 17.24% and 29.56% at the doses of 0.5, 1.0 and 2.0 g/kg, respectively. JWFT also significantly inhibited the increase in hepatic total TG levels, and exhibited 42.34%, 55.49% reduction at medium and high doses respectively. Although the low dose of JWFT did not reduce TG level evidently, it had 27.57% reduction compared with HFC.

The above-mentioned results demonstrated that both medium and high doses of JWFT can lower hepatic total TG and TC levels.

Analysis of fatty acid and expression of CPT1, ACC, SCD1 and FAS of liver
Hepatic fatty acid composition was shown in the Table 2. When rat was given high-fat diet, C18:0, C18:1n-9 and C18:2 were significantly increased as previous report [20]. After the treatment of JWFT, the percentages of C14:0, C15:0, C16:1n-7, C17:0, C18:1n-9 were markedly decreased, only C18:0 significantly (p < 0.05) elevated in HJT group compared with HFC.

The difference of fatty acid composition between HFC and NC group (Figure 2) was profiled systematically by lipidomics. Principal components analysis (PCA) of GC-MS data sets showed that the HFC group was different from NC. JWFT treatment group presented a similar classification to NC group rather than HFC. It suggested that the characteristics of whole fatty acids of JWFT rats were similar to NC rats.

Hepatic stearoyl-CoA desaturase 1 (SCD1) activity could be estimated from the C16:1/C16:0 or C18:1n-9/C18:0 ratios. As shown in Figure 3, SCD1 activity was dose-dependently decreased in JWFT groups, but significantly increased (p < 0.05) in HJT group compared with NC rats.

In order to verify the reliability of estimation, we further analyzed the relative expression of SCD1, which was evidently (p < 0.05, p < 0.01) down-regulated in MJT and HJT groups with 74%, 96% reduction respectively (Figure 3). Furthermore, the expression of other lipogenic genes, ACC and FAS were also down-regulated in MJT and HJT.

Table 1 The contents of major polyphenols and caffeine of JWFT

| Compound | GA | GC | EGC | C | EC | CA | EGCG | GCG | ECG |
|----------|----|----|-----|---|----|----|------|-----|-----|
| Content (mg/g) | 0.794 | 0.804 | 3.969 | ND* | 0.763 | 22.521 | 1.966 | ND* | 1.904 |

*ND, not detected.

GA, gallic acid; GC, (−)-gallocatechin; EGC, (−)-epigallocatechin; C, (+)-catechin; EC, (−)-epicatechin; CA, caffeine; EGCG, (−)-epigallocatechin gallate; GCG, (−)-gallocatechin gallate; ECG, (−)-epicatechin gallate.

Figure 1 The effects of JWFT on the lipids levels. (A) The average body weight of rats each group from 0 to 4 weeks, (B) The weights of abdominal fat of rats after 4 weeks, (C) The levels of serum leptin of rats after 4 weeks, (D) The levels of serum total triglycerides of rats after 4 weeks, (E) The levels of serum total cholesterol of rats after 4 weeks, (F) The levels of serum LDL-C of rats after 4 weeks. *p < 0.05, **p < 0.01 compared with the HFC group.
The expression of CPT1 was up-regulated in HJT, which indicated the increasing of \( \beta \)-oxidation of fatty acids in liver (\( p < 0.05 \)).

Fecal total cholesterol and triglycerides levels
At the 27th day, equivalent lipids (cholesterol and triglycerides) were given to HFC, LJT, MJT and HJT groups. The feces excreted from 0 to 24th hour were analyzed. In all JWFT treatment groups (0.5, 1.0 and 2.0 g/kg), the total fecal TC levels were highly increased compared with the high fat group (\( p < 0.05 \), 0.01) as shown in Table 3.

Discussion
Dysregulations of hepatic lipid synthesis have been associated with several metabolic diseases. We need a
comprehensive understanding of lipogenesis and lipolysis regulation in disease process of metabolic syndrome [21]. For example, nonalcoholic fatty liver disease (NAFLD) is a common disease characterized by excessive accumulation of triglyceride in the liver. Excessive energy intake is an important contributor to the increasing of NAFLD in modern society [22]. In the present study, we intended to investigate the changes related to lipid metabolism of liver tissue induced by high-fat diet using a combined transcriptomic and metabolomic approach [23]. Furthermore, the variations of gene expression indicated the regulation pathways of molecular pathogenesis of hepatic steatosis. In view of the complexity of the hepatic steatosis, our study adopted a strategy of metabolomics profiling for investigating the hepatic fatty acids between normal and hyperlipidemia rats. With PCA analysis of fatty acids composition, it showed that the normal rats were highly different with the high fat induced rats.

After four weeks’ high fat diet feeding, the serum lipids levels, hepatic lipids levels, abdominal fat and body weight were significantly increased. In tea treatment groups, JWFT had the ability to lower the serum lipids (TC, TG and LDL-C). Intra-abdominal fat deposition is a key component of obesity. Some studies indicated that it was an indicator for type 2 diabetes and metabolic syndrome [24]. JWFT had a potent effect on decreasing intra-abdominal fat weight. The results also showed that body weight and abdominal fat weight were positively correlated with serum leptin levels. The serum leptin levels of HFC group was higher than NC group, while

| Fatty acid | NC       | HFC      | LJT       | MJT       | HJT       |
|-----------|----------|----------|-----------|-----------|-----------|
| C14:0     | 0.31 ± 0.06 | 0.215 ± 0.04 | 0.2 ± 0.02 | 0.18 ± 0.03 | 0.1 ± 0.03* |
| C15:0     | 0.577 ± 0.06 | 0.124 ± 0.02 | 0.094 ± 0.02 | 0.04 ± 0.02 | 0.033 ± 0.01* |
| C16:0     | 25.17 ± 0.85 | 24.53 ± 0.49 | 25.38 ± 0.4 | 25.59 ± 0.53 | 23.76 ± 0.65 |
| C16:1n-7  | 1.83 ± 0.28 | 0.665 ± 0.12 | 0.467 ± 0.06 | 0.429 ± 0.09 | 0.223 ± 0.05* |
| C17:0     | 1.4 ± 0.1 | 0.393 ± 0.05 | 0.349 ± 0.02 | 0.31 ± 0.02 | 0.241 ± 0.03* |
| C18:0     | 17.97 ± 0.67 | 21.62 ± 0.85 | 22.38 ± 0.67 | 23.46 ± 0.59 | 24.42 ± 0.88* |
| C18:1n-9  | 10.8 ± 0.83 | 15.08 ± 1.01 | 16.51 ± 1.13 | 16.14 ± 1.00 | 12.86 ± 0.73* |
| C18:1n-7  | 2.69 ± 0.11 | 1.75 ± 0.09 | 1.85 ± 0.34 | 2.09 ± 0.41 | 1.52 ± 0.06 |
| C18:2n-6  | 12.73 ± 0.64 | 13.02 ± 0.65 | 13.86 ± 0.24 | 13.51 ± 0.25 | 13.43 ± 0.37 |
| C18:3n-3  | 0.69 ± 0.08 | 0.76 ± 0.02 | 0.71 ± 0.05 | 0.72 ± 0.06 | 0.67 ± 0.09 |
| C20:3n-3  | 0.38 ± 0.03 | 0.27 ± 0.02 | 0.19 ± 0.01 | 0.18 ± 0.01 | 0.23 ± 0.02 |
| C20:3n-6  | 0.35 ± 0.04 | 0.18 ± 0.03 | 0.12 ± 0.02 | 0.11 ± 0.01 | 0.16 ± 0.03 |
| C20:4n-6  | 13.91 ± 0.61 | 13.30 ± 0.73 | 12.11 ± 0.45 | 12.26 ± 0.32 | 13.41 ± 0.55 |
| C20:5n-3  | 0.52 ± 0.06 | 0.32 ± 0.03 | 0.25 ± 0.02 | 0.21 ± 0.01 | 0.26 ± 0.02 |
| C22:6n-3  | 2.63 ± 0.17 | 2.34 ± 0.11 | 1.95 ± 0.15 | 1.69 ± 0.06 | 2.70 ± 0.22 |

NC, normal chow group; HFC, high dose of fat chow group; LJT, low dose of Jing-wei fu tea group; MJT, medium dose of Jing-wei fu tea group; HJT, high dose of Jing-wei fu tea group. All data are presented as mean ± SEM, *Compared with HFC, *P < 0.05.

Figure 3 The effects of JWFT on expression of lipids metabolism genes. (A) The ratios of C16:1/C16:0, (B) The ratios of C18:1n-9/C18:0, (C) The mRNA expression of hepatic lipogenic genes. *P < 0.05, **P < 0.01, ***P < 0.001 compared with the HFC group.
the leptin levels of were significantly lowered by JWFT supplementation.

The expression of microRNA of major enzymes involved in de novo synthesis, such as SCD1, ACC and FAS were all significantly decreased after high-fat diet, while the expression of CPT1 was increased. After the treatment of JWFT, total hepatic cholesterol and triglycerides levels were also decreased markedly by JWFT treatment, which suggested a potentiality on reversing adiposis hepatica. Furthermore, high dose of JWFT can change the fatty acid composition of high-fat fed rats’ liver. Most of saturated and monounsaturated fatty acids percent-age were decreased significantly, such as C14:0, C15:0, C16:1n-7, C17:0 and C18:1n-9. The C16:1/C16:0 or C18:1n-9/C18:0 ratios may be adjusted by the hepatic SCD1 activity [20]. The ratio of C16:1/C16:0 and C18:1n-9/C18:0 had a notably reduction with high dose JWFT treatment, meanwhile, the level of relative expression of SCD1 was down-regulated considerably in MJT and HJT group. So, it was reasonable that the accumulation of SCD1 substrates (SFA, C16:0 or C18:0) and reduction of products (MUFA, C16:1n-7 or C18:1n-9).

The increased fat accumulation in the liver of HFD rats caused a down-regulation of lipogenic genes and the up-regulation of genes involved in catabolism of free fatty acids, possibly by a feedback mechanism. We found that several enzymes, which have been known to be involved in the de novo synthesis, were down-regulated after the high-fat diet. Parallel down-regulation of SCD1 in our study at mRNA levels is consistent with potentially reduced fatty acid composition in liver of NAFLD [25]. Furthermore, the CPT1 was known to be involved in β-oxidation of fatty acid was also up-regulated in the HFLD-fed group. In four weeks’ experiment, the up-regulation of β-oxidation genes and down-regulation of de novo synthesis could be effective on serum lipids if the experiment had been conducted for a longer duration.

**Conclusions**

In conclusion, JWFT exhibited a potential anti-obesity effects for high-fat diet induced mice, which may be mediated by firstly inhibiting the absorption of lipids, secondly strengthening the feedback regulation of expression of de novo lipogenic genes, and up-regulating CPT1 expression in the liver. Furthermore, the RT-PCR results also showed that rats resisted the over uptake of lipids by down-regulating the expression of lipogenic genes. It was supposed that in a short treatment, the absorption of lipids was firstly controlled, because the low dose of JWFT also promoted the excretion of lipid in feces. In a long term, the regulation of metabolism should be responsible for the lipid controls.

**Abbreviations**

CDT: Chinese dark tea; JWFT: Jingwei fu tea; GA: Gallic acid; C: (−)-Catechin; EC: (−)-Epicatechin; GCC: (−)-Epicatechin gallate; GCG: (−)-Epicatechin gallate; EGC: (−)-Epigallocatechin; GGC: (−)-Epigallocatechin gallate; EGC: (−)-Epigallocatechin gallate; TC: Total cholesterol; TG: Total triglycerides; NC: Normal chow group; HFC: High dose of fat chow group; LJT: Low dose of Jing-wei fu tea group; MJT: Medium dose of Jing-wei fu tea group; HJT: High dose of Jing-wei fu tea group.

**Competing interests**

Authors declare that they do not have any conflict of interests.

**Authors’ contributions**

XCW and JSZ participated in the research design, analysis of the data and reviewing of the manuscript. LZ and LZ conducted all the experiments and draft of the manuscript. All of the authors have read and approved the final manuscript.

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Table 3 The levels of total cholesterols and total triglycerides of feces

|          | HFC (mmol/100 g) | LJT (mmol/100 g) | MJT (mmol/100 g) | HJT (mmol/100 g) |
|----------|------------------|------------------|------------------|------------------|
| TC       | 11.14 ± 1.88     | 17.33 ± 1.32*    | 18.45 ± 2.56*    | 18.60 ± 1.29**   |
| TG       | 6.34 ± 1.00      | 6.74 ± 0.41      | 7.48 ± 0.78      | 8.63 ± 1.38      |

HFC, high dose of fat chow group; LJT, low dose of Jing-wei fu tea group; MJT, medium dose of Jing-wei fu tea group; HJT, high dose of Jing-wei fu tea group. All data are presented as mean ± SEM, *Compared with HFC, *P < 0.05, **P < 0.01.
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