

The Effect of Matcha (*Camellia sinensis*) and Yerba Mate (*Ilex paraguariensis*) Aqueous Extract on Oxidative Stress Status in Obese Rats

Sawsan Adel Noor, Maysa M. El-Mallah and Hany G. EL-Masry

Master's degree

Nutrition and Food Science Department, Faculty of Home Economics, Helwan University, Cairo, Egypt.

Abstract

The present work aimed to investigate the effect of matcha and yerba mate tea aqueous extract on the oxidative stress status of obese rats. Thirty-five adult male Sprague-Dawley rats were divided randomly into 7 equal groups (n=5) as follow: Group 1: negative control, was fed on a basal diet. Groups 2-7 were given lead acetate (LA) (200mg/L) in drinking water for 4 weeks to induce oxidative stress. The rats were then rendered obese by feeding on HFD for 4 weeks. Group 2 was kept as a positive control group (HFD + LA). Groups 3 and 4 were administrated orally with 0.5 and 1 ml of match aqueous extract, whereas 0.5 and 1ml of yerba mate aqueous extract were given to groups 5 and 6, respectively. Group 7 was oral administered with 1 ml of a matcha and yerba mate aqueous extract mixture for 4 weeks. At the end of experiment, rats were scarified and serum was collected for biochemical analyses. Results showed that the administration of HFD-LA group (control positive) resulted in significant elevations in body weight gain, feed efficiency ratio, peritoneal fat pad, serum total cholesterol, triglyceride, low-density lipoprotein cholesterol (LDL-c), very low-density lipoprotein cholesterol (VLDL-c), leptin, nitric oxide (NO), malondialdehyde (MDA), alkaline phosphatase (ALP), aspartate aminotransferase (AST), alanine aminotransferase (ALT), uric acid, urea and creatinine levels as compared to the negative control, and levels of serum high-density lipoprotein cholesterol (HDL-c) and glutathione peroxidase activity (GPx) were significantly decreased. On the other hand, oral administration with mate and matcha extract and their combination attenuated these adverse effects and biochemical alterations caused by HFD-LA administration. In conclusion, yerba mate and matcha exhibit antioxidants and hypolipidemic activity and are effective in reducing body weight in obese rats. The study recommends that consumption of yerba mate and matcha may be beneficial for individuals who suffer from obesity and hyperlipidemia.

Keywords: Matcha, Yerba Mate, Lead acetate, Oxidative stress, Obesity, High fat diet, Rats.

INTRODUCTION

Obesity is an excessive accumulation of fats and a body mass index (BMI) above 30 kg/m². Obesity is a complex disease having an important public health impact worldwide, and its prevalence is increasing (**Pérez-Torre *et al.*, 2021**). The recent global epidemiological data on obesity prevalence show that approximately 2.1 billion subjects are categorized either as overweight or obese (**Naomi *et al.*, 2023**). Obesity is classified as a multifactorial disease that usually arises when there is an excess intake of dietary needs with a low level of energy expenditure. In such a condition, the excess energy will be transformed into triglycerides, which will be then stored in adipose tissue. Over time, adipose tissue will increase in size (hypertrophy), causing the fat cells to increase in size, eventually evincing an increase in body weight (**Chooi *et al.*, 2018**). Abdominal fat is recognized as one of the major risk factors for obesity-related diseases such as: hypertension, dyslipidemia, metabolic syndrome, type 2 diabetes mellitus, coronary heart disease, stroke, non-alcoholic fatty liver disease, etc. (**Čabarkapa *et al.*, 2013** and **Balan *et al.*, 2024**).

One of the common factors leading to obesity could be the excessive intake of a high-fat diet (HFD). An HFD intake will stimulate the voluntary consumption of fat, leading to passive overeating. As so, in obese subjects, the regulation of fatty acid oxidation and lipid peroxidation will be dysregulated. Hence, the manifestation of elevated oxidative stress and inflammation with a repressed antioxidant defense system is common (**Jiang *et al.*, 2021**). Although the complications of obesity might differ from person to person, the consequences are similar for all subjects. This is true for the presence of oxidative stress and inflammation in all

obese patients. One of the common overlooked complications of obesity is cognitive impairments. Obesity-related oxidative stress can cause some pathological changes to arise, such as brain atrophy, as well as inflammation that includes brain atrophy, suppressed hippocampal neurogenesis, reduced microvascular density, blood–brain barrier dysfunction, and reactive astrogliosis, which may result in cognitive decline (**Nguyen *et al.*, 2014**).

Natural sources provide an advantageous basis for the development of novel anti-obesity products (**Karri *et al.*, 2019**). Food supplements and functional foods containing fruit extracts, vegetables, and herbal products are also gaining increased attention as an important natural alternative to prevent metabolic syndrome and obesity (**Engin *et al.*, 2018; Ngoc *et al.*, 2019 and Sander *et al.*, 2020**).

Matcha (*Camellia sinensis* L.) is finely milled green tea made from shaded young tea leaves and processed by low-temperature grinding (**Kolackova *et al.*, 2020**), which has been widely used as a health-beneficial food additive in cookies, candies, chocolates, ice cream, puddings, etc. (**Sutton, 2004**). Compared to green tea, matcha excels in high contents of chlorophyll and catechin due to the tea leaves being protected from direct sunlight before harvest. Moreover, with a different intake method from traditional green tea, both water-soluble and water-insoluble ingredients in matcha can be ingested and utilized. Previous studies suggested that matcha plays an important role as an antioxidant, anticarcinogen, anti-inflammatory, and anti-hypercholesterolemia (**Jeszka-Skowron *et al.*, 2015; Bonuccelli *et al.*, 2018 and Das *et al.*, 2019**). More relevantly, dietary matcha supplement could inhibit lipid

accumulation and ameliorate metabolic damage in obese mice induced by HFD (Willems *et al.*, 2018 and Zhou *et al.*, 2020).

Yerba mate (*Ilex paraguariensis*) is widely consumed by some South American populations and plays an important role in the culture and economy of Brazil, Argentina, Uruguay and Paraguay. Apart from the traditional South American mate beverages, yerba mate has been sold as an energy drink or as a weight reduction aid in different regions, as in Europe (Bracesco *et al.*, 2011). In the last 20 years, different studies have described potential therapeutic properties of mate tea, such as the capacity of modulating inflammation in several key tissues (hypothalamus, muscle, adipose tissue and liver) (Arcari *et al.*, 2009, 2011 and Pimentel *et al.*, 2012). These potential therapeutic effects of yerba mate are related to its bioactive compounds' composition, in particular to the presence of phenolic acids. The phenolic acid composition of mate tea is similar to that of coffee, chlorogenic acids being the most abundant compounds (Jaiswal *et al.*, 2010). Therefore, this study was conducted to evaluate the effect of matcha and yerba mate tea aqueous extract in the oxidative stress status on obese rats.

Materials and Methods

A. Materials

Matcha tea and Yerba Mate tea were obtained from the Biotechnology Unit, National Research Centre, Dokki, Giza, Egypt. Biochemical kits for blood analysis were purchased from Alkan Company for Biodiagnostic Reagents, Dokki, Cairo, Egypt. Lead acetate, casein, cellulose, choline chloride, D-L methionine, vitamins and mineral constituents were purchased from El-Gomhoriya Pharmaceutical Company, Cairo, Egypt. Starch, soy oil, and sucrose were obtained from the Egyptian local market. Thirty-five adult male albino rats (Sprague Dawley strain), weighing about

170 \pm 5 g b.wt. were obtained from the Laboratory Animal Colony, Agricultural Research Center, Giza, Egypt.

B. Methods

Preparation of an aqueous extract of matcha and yerba mate:

Hot water was added to the powder of matcha, stirred and soaked it overnight to obtain an aqueous extract 5% concentration (5 gram per 100 ml of water).

Induction of obesity:

Rats were fed four weeks on a basal diet according to **Reeves et al., (1993)**, with some modification in fat content containing: casein 14%, cellulose 5%, vitamin mixture 1%, mineral mixture 3.5%, sucrose 10%, (beef tallow 19% + corn oil 1%), l-cystine 0.18, choline bitartrate 0.25% and the remainder was starch, to induce obesity in rats (**Liu et al., 2004**).

Induction of oxidative stress:

Induction of oxidative stress in rats by using lead acetate powder which was prepared by diluting in double distilled water prior to use, in lead acetate group the animals were given a 6 ml lead acetate (200mg/L) in drinking water for four weeks according to (**Newairy and Abdou, 2009**).

Experimental Animal Design:

Thirty-five adult male albino rats were housed in well conditions and fed on basal diet in Research Labs, Agricultural Research Center, Giza, Egypt. After one week of acclimatization, the rats were randomly divided into two main groups as follows:

- **First main group:** Negative control group, rats (n=5) were fed on a basal diet only during the experimental period.
- **second main group:** Rats (n=30) were given lead acetate (200 mg/L) in drinking water for 4 weeks, and were fed on HFD for 4 weeks to induce obesity, then they were divided into six subgroups as follow:

Subgroup (1): Rats were fed on high fat diet and drinking water with 200mg/L lead acetate and served as positive control group.

Subgroup (2): Rats were fed on HFD and drinking water with 200mg/L lead acetate and were given 0.5 ml of Yerba Mate aqueous extract orally by gastric tube.

Subgroup (3): Rats were fed on HFD and drinking water with 200mg/L lead acetate and were given 1 ml of Yerba Mate aqueous extract orally by gastric tube.

Subgroup (4): Rats were fed on HFD and drinking water with 200mg/L lead acetate and were given 0.5 ml of Matcha aqueous extract orally by gastric tube.

Subgroup (5): Rats were fed on HFD and drinking water with 200mg/L lead acetate and were given 1 ml of Matcha aqueous extract orally by gastric tube.

Subgroup (6): Rats were fed on HFD and drinking water with 200mg/L lead acetate and were given 1 ml of Yerba Mate-Mate aqueous extract (1:1) orally by gastric tube.

Biological Evaluation

During the experiment period, the quantities of diet that were consumed and/or waste, were recorded every day. Water and basal diet had been introduced under hygienic conditions.

Feed intake was recorded daily and animals were weighed at the beginning and twice a week throughout the experimental period. Body weight gain percent (BWG%) and feed efficiency ratio were

determined according to **Chapman *et al.*, (1959)**, using the following equation:

$$\text{BWG}\% = \frac{\text{Final body weight} - \text{Initial body weight}}{\text{Initial body weight}} \times 100$$

$$\text{FER} = \text{Weight gain} / \text{Feed intake.}$$

Peritoneal fat pad was dissected from the rats, then weighed and stored at -20 °C according to the methods of **Azain *et al.*, (2000)**. The percentage of peritoneal fat pad was calculated as follows:

$$\text{Weight of peritoneal fat pad} / \text{Weight of rat} \times 100.$$

Biochemical Analysis of Serum

At the end of the experimental period (4 weeks), rats were fasted overnight before sacrificing and blood samples were collected from each rat and centrifuged at 3000 rpm for 15 min to obtain serum for biochemical analysis. Serum total cholesterol, triglyceride and high-density lipoprotein cholesterol were determined according to **Richmond, (1973); Wahlefeld, (1974) and Albers *et al.*, (1983)**, respectively. Low density lipoprotein cholesterol and very low-density lipoprotein cholesterol were calculated according to **Friedewald *et al.*, (1972)**. Serum leptin was determined according to the methods described by **Zhang *et al.*, (1995)**. Malondialdehyde (MDA), glutathione peroxidase (GPx) and nitric oxide (NO) were determined according to **Draper and Hadley, (1990); Hissin and Hilf, (1970) and Ding *et al.*, (1988)**, respectively. Serum aspartate aminotransaminase (AST), alanine aminotransaminase (ALT), urea and creatinine were determined according to the method described by **Young, (2001)**. Serum alkaline phosphates (ALP) and uric acid

were determined according to **Roy, (1970)** and **Milena, (2003)**, respectively.

Statistical Analysis

The obtained results were analyzed according to the SPSS program. An ANOVA test was used to compare results among groups and $P \leq 0.05$ were considered to be significant (**SPSS, 1989**).

Results and Discussion

Results in **Table 1** showed that feed intake increased in positive control rats compared with the negative control rats. On the other hand, BWG%, FER and PFP% significantly increased ($P < 0.05$) in the positive control group (+ve) when compared to the negative control group (-ve). Oral administration with yerba mate and matcha and their combination on obese rats caused a high reduction in the feed intake, BWG, FER and PFP% as compared with the positive control group. These findings were in the same line with **Han et al., (2022)**, who reported that excessive calorie intake results in the storage of energy in the form of fat within the body. An HFD, due to its elevated calorie content, is prone to inducing weight gain. Moreover, when the body exhibits a higher efficiency in utilizing food, the probability of obesity tends to increase.

The present results in **Table 1** were in agreement with **Borges et al., (2013)**, who noticed that there was a significant interaction between the HFD and mate tea consumption in terms of weight gain, because the administration of yerba mate promoted weight loss only in HFD-fed animals. Furthermore, in accordance with other studies **Arcari et al., (2011)**, reported that the administration of yerba mate aqueous extract could reverse, or at least attenuate, all of

the observed detrimental effects induced by the HFD. Not only did the consumption of mate tea reduce adipose tissue depots and improve body composition, but it also promoted weight loss in HFD-fed rats. The impact of yerba mate on body composition might be attributed to an increase in energy expenditure and fat oxidation, because yerba mate administration increases the expression of uncoupling proteins (Arcari *et al.*, 2009) and activates AMP-activated protein kinase-a in adipose tissue (Pang *et al.*, 2008).

In complement of the present results, green tea contains and is enriched with catechins, known for their antioxidant properties, including epigallocatechin-3-gallate (EGCG), epigallocatechin (EGC) and epicatechin gallate (ECG) (Zheng *et al.*, 2004). As an exceptional kind of green tea, matcha's special planting pattern and way of drinking made it different from the other green tea. Xu *et al.*, (2016) showed that intake of both water-soluble and water-insoluble parts of Matcha was more helpful in controlling weight that was elevated by HFD. Zhou *et al.*, (2021) established that dietary supplementation with matcha could effectively inhibit the weight gain and fat accumulation against HFD-induced obese mouse models. Green tea and its catechins, particularly EGCG, have been shown to lower body weight, adipose tissue, and blood lipid levels. The mechanism of action of EGCG includes a decrease in energy intake, an increase in energy expenditure, as well as changes in the activities of fat, liver, muscle and intestinal cells (Singla *et al.*, 2010 and Hena-Mejia *et al.*, 2012).

Data in **Table 2** revealed that positive control rats had a significant ($P < 0.05$) increases in serum levels of TC, TG and LDL-c, VLDL-c and a significant decrease ($P < 0.05$) in HDL-c when

compared to the negative control group. Rats that administrated with mate and matcha extract and their mixture had significant ($P<0.05$) reduction in the elevated serum TC, TG, LDL-c and VLDL-c levels and an increase in serum HDL-c when compared with positive control group. Also, data illustrated that rats in group 7 administrated with 1 ml of mate and matcha extract recorded the best results for modulating serum lipid profile.

Serum lipid profile confirmed the induction by HFD-LA, in which abnormalities in lipid metabolism, increasing cholesterol, triacylglycerol, LDL-c and VLDL-c, and depressed HDL-c, as compared to the control rat group. These findings were approved by **Bagabaldo *et al.*, (2022); Chu *et al.*, (2024); Diab *et al.*, 2024 and Hussein and Mustafa, 2024**). In the present study, obese rats that administrated with the extract of yerba mate and matcha significantly improved the lipid profile alterations. These findings were interpreted by **Taketa *et al.*, (2004)**, who reported that saponin, an important compound found in yerba mate, has been reported to interfere with cholesterol metabolism and provide a hypocholesteremic effect by inhibiting the passive diffusion of colic acid through the formation of micelles preventing absorption, anticancer and antiparasitic. Furthermore, chlorogenic acid, the main polyphenols in yerba mate, is thought to modulate the activity of glucose-6- phosphatase, which is involved in glucose metabolism and reduce the risk of cardiovascular disease by decreasing oxidation of LDL and cholesterol (**Klein *et al.*, 2011**).

On the other hand, the present results were in agreement with **El-Kholie *et al.*, (2022)** who found that matcha tea had the lowest cholesterol and triglyceride, LDL-c and VLDL-c levels, as well as the greatest levels of HDL-c, with significant differences.

Matcha green tea administration dramatically reduced triglyceride and total cholesterol levels in rats, according to **Basu *et al.*, (2019)**. These findings correspond with those of **Xu *et al.*, (2020)** who found that green tea drinking lowers LDL cholesterol and TC, in both normal weight and overweight/obese participants. Both in vitro and animal experiments have shown that green tea catechins can significantly reduce the levels of plasma triglycerides, total cholesterol and LDL-C (**Xing *et al.*, 2019**). Some randomized controlled trials and meta-analyses have suggested that green tea may affect the lipid profiles in subjects with cardiovascular-related diseases such as hypercholesterolemia, and glucose intolerance as well as in healthy individuals (**Ferreira *et al.*, 2017**). This might be explained by the fact that matcha tea contained higher fiber which could lower the levels of TC and TG in the blood, and also might be effective in controlling body weight, improving lipid levels and increasing the antioxidant status of mice and human beings (**Sokary *et al.*, 2023**).

Table 3 indicated that the higher level of serum leptin hormone in the obese control group as compared with the negative control group. Oral administration with mate and matcha green tea extracts and their extracts diminished the increasing in leptin levels in the treated group as compared with the positive control group. Group 7 that was treated with the combination extract of yerba mate and matcha was the best group to normalize the level of leptin hormone.

The present results found that HFD-Lead acetate in drinking water caused elevated leptin hormone levels in rats, these findings were in agreement with **Lindholm *et al.*, (2013)** and **Beier *et al.*, (2015)**. A common metabolic change observed in obesity is

increased leptin concentration in the plasma. Generally, white adipose tissue produces leptin and it exhibits the effects either via the nervous system or directly via the activity of autocrine signaling (**Martínez-Sánchez, 2020**). Leptin is a hormone that controls satiety. In obesity, an increased concentration of leptin is due to the excessive level of adipocytes that impairs the leptin signaling receptor, developing a condition known as leptin resistance. As a consequence, the ability of leptin to feel a sense of satiety is lost, leading to decreased energy expenditure and overconsumption of nutrients (**Andreoli et al., 2019**). Moreover, **Balsan et al., (2019)** noticed that leptin levels were decreased in the yerba mate group, as found in the present results. In addition, these findings were supported by **Wahyuni et al., (2023)** who found that green tea extract caused a significant decrease in leptin levels as compared to +ve group.

Results presented in **Table 4**, showed rats in the positive control group had a significant reduction ($P<0.05$) in serum levels of GPx while had a significant ($P<0.05$) elevation in serum NO and MDA levels when compared with the negative control group. On the other hand, oral administration of mate and matcha extract and their mixture to obese rats resulted in a significant increase ($P<0.05$) in GPx level and decrease in NO level, also this reduction applied to MDA except the low level of mate and matcha 0.5 ml (group 3 and 5) when compared to the positive control group. It was observed that the 1 ml mixture of mate and matcha considered the best result for elevating the antioxidant activity (GPx) and inhibit free radicals and oxidation.

According to present results, high fat diet-lead-acetate in drinking water led to oxidative stress by increasing in levels of MDA and NO and decreasing in antioxidant activity (glutathione). These results are supported by **Samarghandian *et al.*, (2013)** and **Chung *et al.*, (2018)**. Obesity is often characterized by high oxidative stress resulting from an imbalance between an excessive generation of reactive oxygen species (ROS) and insufficient antioxidant defense mechanisms such as GPx, which induces severe damage to DNA, protein and lipids in the cells (**Conceição *et al.*, 2013** and **Machado *et al.*, 2021**). Nitric oxide is one of the most important mediators of intra- and extra-cellular processes. It plays a complex role in humans, as it is sometimes beneficial and sometimes harmful. It is potentially toxic under oxidative stress conditions (**Halliwell and Gutteride, 2007**). **Zenaro *et al.*, (2014)** showed that the treatment with extract of yerba mate was able to reverse the oxidative stress on the serum by reduction in MDA level. **Branco *et al.*, (2013)** reported that the administration of mate avoided the increase in oxidative damage and the nitric oxide production, as found in the present study. Thus, the aqueous extract of yerba mate showed antioxidant action, being a potential protective mechanism against damage caused by oxidative stress. Also, the present results showed that yerba mate normalized GPx when compared to the +ve group, these results were found by **de Oliveira *et al.*, (2018)**. **Cho *et al.*, (2010)** demonstrated that the antioxidants in yerba mate, especially caffeic and chlorogenic acids, can contribute to protective effects against obesity-related oxidative damage. On the other hand, **Xu *et al.*, (2016)** confirmed that matcha could significantly lower the MDA level in serum; besides, the serum GPx activity indicated that the oxidative stress caused by HFD could be reversed by administration of matcha. **Nakagawa**

and Yokozawa, (2002) showed that green tea extract significantly inhibited nitric oxide production. Among the green tea components, tannin showed an inhibitory activity due to its excellent antioxidant properties.

Results in **Table 5** showed that serum levels of AST, ALT and ALP significantly increased ($P<0.05$) in obese control group when compared to negative control group. Rats that were administrated with mate and matcha and their mixture had a significant reduction ($P<0.05$) in serum levels of AST, ALT and ALP whereas AST of group 3 (0.5 ml mate) had no significant reduction when compared to positive control group. Moreover, group 7 (1 ml mate: matcha) was recorded as the best group for reducing serum liver enzymes.

Seum liver enzymes (AST, ALT and ALP) in the present results recorded a high elevation with HFD-LA, these results were confirmed by **Lasker et al., (2019)** and **Asgharian et al., (2022)**. According to **Abolfathi et al., (2012)**, liver function tests are commonly used in clinical practice to screen for liver disease, monitor the course of recognized disease, and evaluate the effects of potentially hepatotoxic medicines. In complement of the present results, **El mallah, (2015)** reported that oral administration of yerba mate tea to obese rats significantly reduced serum markers of liver functions AST, ALT and ALP. **Martins et al., (2009)** reported that, mate tea can protect unsaturated fatty acids from oxidation and may have selective protective effects within the body, especially on the liver. Additionally, yerba mate infusions were also characterized by a very high antioxidant activity (**Katarzyna et al., 2020**). Data suggested that mate tea has strong antioxidant and lipid-lowering

effects that prevents hepatic fatty deposition, and controls the expression of lipid metabolic regulators. It can therefore be used to reduce the risk of atherosclerosis (**Boaventura et al., 2012**). Moreover, **Zhou et al., (2021)** confirmed that the administration with aqueous extract of matcha te effectively decreased serum AST and ALT activities, as found in the present study. Matcha tea supplementation effectively prevented excessive visceral and hepatic lipid accumulation, elevated blood glucose, dyslipidemia, abnormal liver function, and steatosis hepatitis (**Li et al., 2020**). The hepatoprotective activities of green tea are attributed to its catechins that scavenge ROS/RNS in vitro (**Hasanein et al., 2012**). These results revealed that matcha tea and yerba mate could remit the pathological process in obesity-induced hepatic damage.

Table 6, revealed the effect of mate and matcha extract on serum kidneys functions of obese rats. Results showed that serum levels of urea, creatinine and uric acid significantly increased ($P<0.05$) in the obese control group when compared to the negative control group. Rats that received mate and matcha extract and their mixture had a significant reduction ($P<0.05$) in serum levels of urea, creatinine and uric acid when compared to the positive control group. Also, it was observed that there were no significant differences between group 5 and 7 as compared to negative control group in uric acid. Furthermore, group 7 that treated with 1 ml of mate and matcha mixture achieved the best level to reduce the elevated kidneys functions.

The present results indicated that the HFD-LA induced rats exhibited significant elevations in kidney function parameters (urea, creatinine and uric acid), as found by **Ahmed and Sabra, (2018)** and **Shahin and Aburaya, (2023)**. In the present study, yerba mate

reduced serum kidneys functions. These results were in accordance with other study which showed that yerba mate is able to influence the circulatory system, acting as a diuretic and hypotensive agent **Boaventura *et al.*, (2012)**. Also, these results agree with **Aboelnaga and Balouch, (2021)** who reported that yerba mate tea decreased kidneys functions level that elevated in the serum.

Green tea can help to bring urea, creatinine, and uric acid levels back to normal. **El-Kholie *et al.*, (2022)** approved that matcha tea decreased serum kidneys functions (urea, creatinine and uric acid) as compared to positive group, as found in the present study. Also, these findings are consistent with those of **Yamabe *et al.*, (2009)**, who found that matcha tea therapy significantly reduced kidney advanced glycation end products AGE levels as well as serum thiobarbituric acid-reactive compounds. Match supplementation also resulted in lower levels of renal N (6)-(carboxymethyl) lysine (CML), N (6)-(carboxylethyl) lysine (CEL), and RAGE expression, as well as an increase in hepatic SREBP-2 expression, but not sterol regulatory element binding proteins (SREBP-1). These findings imply that matcha protects against hepatic and renal damage by inhibiting the buildup of AGE in the kidneys, lowering hepatic glucose, triglyceride, and total cholesterol levels, and acting as an antioxidant. Green tea is also promising as a nephroprotective drug against diethyl nitrosamine (DEN) and ferric nitrilotriacetate (Fe-NTA) caused nephrotoxicity in Wistar rats, according to (**El-Desouky *et al.*, 2019**).

In conclusion, the present findings illustrate that yerba mate and matcha tea consumption led to an improvement in the biochemical changes resulting from obesity. These therapeutic

effects of yerba mate and matcha tea can be attributed to their antioxidant properties due to containing phenolic compounds.

Table 1: Effect of Mate and Matcha Aqueous Extract on Feed Intake (FI), Body Weight Gain% (BWG%), Feed Efficiency Ratio (FER) and Peritoneal Fat Pad% (PFP%) of Obese Rats

Parameters Groups	FI (g/d/rat)	BWG%	FER	PFP%
G1: -Ve Control	20	23.49±1.09 ^{bc}	0.052±0.001 ^c	4.87±0.16 ^{cd}
G2: +Ve Control (HFD-LA)	22	47.42±1.86 ^a	0.101±0.002 ^a	11.55±0.79 ^a
G3: 0.5 ml Mate	21	26.29±1.58 ^b	0.087±0.001 ^b	7.85±0.33 ^b
G4: 1 ml Mate	20	25.27±0.52 ^{bc}	0.058±0.001 ^c	6.74±0.20 ^{bc}
G5: 0.5 ml Matcha	19.5	21.84±0.83 ^{bc}	0.079±0.001 ^b	5.58±0.26 ^{cd}
G6: 1 ml Matcha	19	22.65±0.53 ^{bc}	0.054±0.001 ^c	5.35±0.27 ^{cd}
G7: 1 ml Mate: Matcha	18	19.67±0.52 ^c	0.052±0.001 ^c	4.29±0.26 ^d

* Results are expressed as means ± SE.

*Mean values at the same column with the same superscript letters are not statistically significant at P<0.05.

* HFD = High Fat Diet. * LA = Lead Acetate.

Table 3: Effect of Mate and Matcha Aqueous Extract on Serum Lipid Profile of Obese Rats

Parameters Groups	TC	TG	HDL-c	LDL-c	VLDL-c
	mg/dl				
G1: -Ve Control	123.19±0.49 ^e	79.13±0.52 ^f	58.96±0.92 ^a	48.66±0.94 ^f	15.82±0.10 ^f
G2: +Ve Control (HFD-LA)	187.81±0.67 ^a	139.27±0.86 ^a	27.29±0.96 ^e	132.66±0.65 ^a	27.85±0.17 ^a
G3: 0.5 ml Mate	160.33±0.63 ^b	130.79±0.53 ^b	34.89±0.53 ^d	99.27±1.17 ^b	26.16±0.10 ^b
G4: 1 ml Mate	148.01±0.33 ^c	118.19±0.80 ^c	40.11±0.50 ^c	84.26±1.12 ^c	23.64±0.16 ^c
G5: 0.5ml Matcha	137.58±0.95 ^d	121.59±0.69 ^c	33.29±0.80 ^d	79.97±1.14 ^{cd}	24.32±0.13 ^c
G6: 1 ml Matcha	141.38±0.25 ^{cd}	112.59±0.96 ^d	42.29±0.51 ^c	76.57±1.15 ^d	22.52±0.34 ^d
G7: 1 ml Mate: Matcha	137.29±0.77 ^d	100.59±0.95 ^e	48.49±0.47 ^b	68.67±0.85 ^e	20.12±0.20 ^e

* Results are expressed as means ± SE.

*Mean values at the same column with the same superscript letters are not statistically significant at P<0.05.

* HFD = High Fat Diet. * LA = Lead Acetate.

Table 3: Effect of Mate and Matcha Aqueous Extract on Serum Leptin Hormone of Obese Rats

Parameters Groups	Leptin (ng/ml)
G1: -Ve Control	9.14±0.25 ^f
G2: +Ve Control (HFD-LA)	17.33±0.31 ^a
G3: 0.5 ml Mate	15.74±0.27 ^b
G4: 1 ml Mate	13.56±0.31 ^{cd}
G5: 0.5 ml Matcha	14.64±0.18 ^{bc}
G6: 1 ml Matcha	12.81±0.18 ^{de}
G7: 1 ml Mate: Matcha	11.73±0.26 ^e

* Results are expressed as means ± SE

*Mean values at the same column with the same superscript letters are not statistically significant at P<0.05.

* HFD = High Fat Diet. * LA = Lead Acetate.

Table 4: Effect of Mate and Matcha Aqueous Extract on Serum Malondialdehyde (MDA), Glutathione Peroxidase (GPx), and Nitric Oxide (NO) of Obese Rats

Parameters Groups	GPx (U/ml)	NO (μmol/L)	MDA (ng/ml)
G1: -Ve Control	136.62 \pm 0.70 ^a	62.57 \pm 0.74 ^f	118.78 \pm 0.52 ^e
G2: +Ve Control (HFD-LA)	83.23 \pm 0.40 ^e	98.05 \pm 0.55 ^a	402.25 \pm 1.68 ^a
G3: 0.5 ml Mate	94.76 \pm 0.35 ^d	92.45 \pm 0.41 ^b	393.81 \pm 1.89 ^{ab}
G4: 1 ml Mate	100.91 \pm 0.52 ^d	88.05 \pm 0.55 ^c	382.41 \pm 1.55 ^b
G5: 0.5 ml Matcha	115.34 \pm 0.57 ^c	89.37 \pm 0.45 ^c	391.14 \pm 2.47 ^{ab}
G6: 1 ml Matcha	121.18 \pm 0.63 ^c	81.75 \pm 0.60 ^d	302.19 \pm 2.74 ^c
G7: 1 ml Mate: Matcha	129.18 \pm 0.34 ^b	74.92 \pm 0.50 ^e	287.34 \pm 3.71 ^d

* Results are expressed as means \pm SE.

*Mean values at the same column with the same superscript letters are not statistically significant at P<0.05.

* HFD = High Fat Diet. * LA = Lead Acetate.

Table 5: Effect of Mate and Matcha Aqueous Extract on Serum Liver Functions of Obese Rats

Parameters Groups	AST (μ /L)	ALT (μ /L)	ALP (mg/dl)
G1: -Ve Control	18.18 \pm 0.25 ^e	36.53 \pm 0.40 ^f	115.18 \pm 1.04 ^f
G2: +Ve Control (HFD-LA)	47.18 \pm 0.98 ^a	95.93 \pm 1.04 ^a	172.58 \pm 1.56 ^a
G3: 0.5 ml Mate	40.38 \pm 0.74 ^{ab}	81.93 \pm 0.50 ^b	164.18 \pm 1.31 ^b
G4: 1 ml Mate	32.38 \pm 0.97 ^{cd}	70.91 \pm 0.45 ^{cd}	160.38 \pm 1.13 ^b
G5: 0.5 ml Matcha	33.39 \pm 0.69 ^{cd}	75.13 \pm 0.61 ^c	144.58 \pm 0.83 ^c
G6: 1 ml Matcha	38.78 \pm 0.39 ^{bc}	66.53 \pm 0.55 ^{de}	134.38 \pm 1.14 ^d
G7: 1 ml Mate: Matcha	30.38 \pm 0.89 ^d	61.73 \pm 0.34 ^e	125.38 \pm 1.02 ^e

* Results are expressed as means \pm SE.

*Mean values at the same column with the same superscript letters are not statistically significant at P<0.05.

* HFD = High Fat Diet. * LA = Lead Acetate.

Table 6: Effect of Mate and Matcha Aqueous Extract on Serum Kidneys Functions of Obese Rats

Parameters Groups	Urea	Creatinine	Uric Acid
	(mg/dl)		
G1: -Ve Control	24.12±0.40 ^e	0.70±0.01 ^f	2.65±0.07 ^d
G2: +Ve Control (HFD-LA)	49.53±0.66 ^a	1.68±0.02 ^a	5.92±0.16 ^a
G3: 0.5 ml Mate	42.52±0.78 ^b	1.48±0.03 ^b	4.79±0.25 ^b
G4: 1 ml Mate	34.64±0.64 ^c	1.30±0.01 ^{cd}	3.53±0.17 ^c
G5: 0.5 ml Matcha	39.48±0.59 ^b	1.40±0.02 ^{bc}	3.08±0.03 ^{cd}
G6: 1 ml Matcha	33.97±0.89 ^c	1.20±0.03 ^d	3.50±0.19 ^c
G7: 1 ml Mate: Matcha	28.21±0.55 ^d	0.94±0.04 ^e	2.93±0.04 ^{cd}

* Results are expressed as means ± SE.

*Mean values at the same column with the same superscript letters are not statistically significant at P<0.05.

* HFD = High Fat Diet. * LA = Lead Acetate.

REFERENCES

Aboelnaga, S. and Balouch, F. (2021): Protective Effect of Yerba Mate on Rats Fed on High Cholesterol Diet- An Experimental Study. JCDR., 15: 9-13.

Abolfathi, A., Mohajeri, D., Rezaie, A. and Nazeri, M. (2012): Protective Effects of Green Tea Extract against Hepatic Tissue Injury in Streptozotocin-Induced Diabetic Rats. Evidence-Based Complementary and Alternative Medicine, 1: 1-10.

Ahmed, N. and Sabra, H. (2018): Reno- protective Effect of Graviola (Annona Muricata) Leaves Against Lead Acetate Toxicity on experimental Albino Rats. Biochemistry letters, 13: 1-13.

Albers, N., Benderson V. and Warnick G. (1983): Enzymatic determination of high-density lipoprotein cholesterol, Selected Methods. Clin. Chem., 10: 91-99.

Andreoli, M., Donato, J., Cakir, I. and Perello, M. (2019): Leptin resensitisation: A reversion of leptin-resistant states. J. Endocrinol, 241: 81-96.

Arcari, D., Bartchewsky, W., dos Santos, T., Oliveira, K., Funck, A. and Pedrazzoli, J. (2009): Antiobesity effects of yerba mate extract (*Ilex paraguariensis*) in high-fat diet-induced obese mice. Obesity 17:2127–2133.

Arcari, D., Bartchewsky, W., dos Santos, T., Oliveira, K., Oliveira, C. and Gotardo, E. (2011): Anti-inflammatory effects of yerba mate extract (*Ilex paraguariensis*) ameliorate insulin resistance in mice with high fat diet-induced obesity. Mol Cell Endocrinol 335: 110-115.

Asgharian, S., Lorigooini, Z., Bijad, E., Hosseinkhani, H., Abbasian, Z. and Rafieian-Kopaei, M. (2022): Protective effect of Rheum ribes extract against lead-induced hepatotoxicity in male rats. Braz. J. Pharm. Sci., 58: 1-9.

Azain, M., Hausman, D., Sisk, M., Flat, W. and Jewell, D. (2000): Dietary conjugated Linoleic acid reduces rat adipose tissue cell size rather than cell number. J.Nutr., 130: 1548-1554.

Bagabaldo, P., Atienza, L. and Castillo-Israel, K. (2022): ‘Saba’ banana (*Musa acuminata* x *balbisiana* BBB Group) peel pectin supplementation improves biomarkers of obesity and associated blood lipid disorders in obese hypercholesterolemic mice. *Curr Res Food Sci.*, 5: 251-260.

Balan, A., Halațiu, V. and Scridon, A. (2024): Oxidative Stress, Inflammation, and Mitochondrial Dysfunction: A Link between Obesity and Atrial Fibrillation. *Antioxidants (Basel)*, 17: 117.

Balsan, G., Pellanda, L., Sausen, G., Galarraga, T., Zaffari, D., Pontin, B. and Portal, V. (2019): Effect of yerba mate and green tea on paraoxonase and leptin levels in patients affected by overweight or obesity and dyslipidemia: a randomized clinical trial. *Nutrition Journal*, 18: 1-10.

Basu, A., Du, M., Sanchez, K., Leyva, M., Betts, N., Blevins, S., Wu, M., Aston, C. and Lyons, T. (2019): Green tea minimally affects biomarkers of inflammation in obese subjects with metabolic syndrome. *Nutrition*, :27: 206-213.

Beier, E., Inzana, J., Sheu, T., Shu, L., Puzas, J. and Mooney, R. (2015): Effects of Combined Exposure to Lead and High-Fat Diet on Bone Quality in Juvenile Male Mice. *Environmental health perspectives*, 123: 935-943.

Boaventura, B., di Pietro, P., Stefanuto, A., Klein, G., de Moraes, E., de Andrade, F. and Wazlawik, E. (2012): Association of mate tea (*Ilex paraguariensis*) intake and dietary intervention and effects on oxidative stress biomarkers of dyslipidemic subjects. *Nutrition*, 28: 657-664.

Bonuccelli, G., Sotgia, F. and Lisanti, M. (2018): Matcha green tea (MGT) inhibits the propagation of cancer stem cells (CSCs), by targeting mitochondrial metabolism, glycolysis and multiple cell signalling pathways. *Aging*, 10: 1867-1883.

Borges, M., Vinolo, M., Nakajima, K., Castro, I., Bastos, D., Borelli, P., Fock, R., Tirapegui, J., Curi, R. and Rogero, M. (2013): The effect of mate tea (*Ilex paraguariensis*) on metabolic and inflammatory parameters in high-fat diet-fed Wistar rats. *International Journal of Food Sciences and Nutrition*, 1: 1-9.

Bracesco, N., Sanchez, A., Contreras, V., Menini, T. and Gugliucci, A. (2011): Recent advances on *Ilex paraguariensis* research: minireview. J. Ethnopharmacol., 136: 378–384.

Branco, C., Scola, G., Rodrigues, A., Cesio, V., Heinzen, H., Godoy, A., Funchal, C., Coitinho, A. and Salvador, M. (2013): Organic and Conventional Yerba Mate (*Ilex paraguariensis* A. St. Hil) Improves Metabolic Redox Status of Liver and Serum in Wistar Rats. Antioxidants, 2: 100-109.

Čabarkapa, V., Đerić, M., Stošić, Z., Sakač, V., Davidović, S. and Eremić, N. (2013): Determining the relationship between homocysteinemia and biomarkers of inflammation, oxidative stress and functional kidney status in patients with diabetic nephropathy J Med Biochem., 32: 131.

Chapman, D., Gastilla, R. and Campbell, J. (1959): Evaluation of protein in foods: 1- A Method for the determination of protein efficiency ratio. Can. J. Biochem. Phys; 37: 679- 686.

Cho, A., Jeon, S., Kim, M., Yeo, J., Seo, K. and Choi, M. (2010): Chlorogenic acid exhibits anti-obesity property and improves lipid metabolism in high-fat diet-induced-obese mice. Food Chem Toxicol., 48: 937-943.

Chooi, Y., Ding, C. and Magkos, F. (2018): The epidemiology of obesity. Metabolism, 92: 6-10.

Chu, W., Yang, L., Lin, W. and Chien, Y. (2024): Evaluation of the Anti-Obesity Effect of a Vegan Complex Supplement (Sinetrol, Green Tea, Bergamot and Chromium Yeast) in High-Fat Diet-Induced Obese Rats. Obese Res Open J., 10: 6-13.

Chung, A., Gurtu, S., Chakravarthi, S., Moorthy, M. and Palanisamy, U. (2018): Geraniin Protects High-Fat Diet-Induced Oxidative Stress in Sprague Dawley Rats. Front. Nutr., 5: 17.

Conceição, E., Franco, J., Oliveira, E., Resende, A., Amaral, T. and Peixoto-Silva N. (2013): Oxidative stress programming in a rat model of postnatal early overnutrition-role of insulin resistance. J Nutr Biochem., 24: 81-87.

Das, P., Kim, Y., Hong, S. and Eun, J. (2019): Profiling of volatile and non-phenolic metabolites-Amino acids, organic acids, and sugars of green tea extracts obtained by different extraction techniques. Food Chem., 296: 69-77.

de Oliveira, E., Lima, N., Conceição, E., Peixoto-Silva, N., Moura, E. and Lisboa, P. (2018): Treatment with *Ilex paraguariensis* (yerba mate) aqueous solution prevents hepatic redox imbalance, elevated triglycerides, and microsteatosis in overweight adult rats that were precociously weaned. *Brazilian Journal of Medical and Biological Research*, 51: 1-10.

Diab, A., Al-Mathal, E., Zahra, M., Dowidar, M., El-Sherbeny, S. and Abu El-Magd, M. (2024): Ameliorative Effect of *Costus speciosus* Extracts on Toxic Effects of Lead Acetate on Liver and Kidney of Male Rats. *Egyptian Journal of Veterinary Sciences*, 55: 895-901.

Ding, A., Nathan, C. and Stuchr, D. (1988): Release of reactive nitrogen intermediates and reactive oxygen intermediate from mouse peritoneal macrophages. Comparison of activating cytokines and evidence for independent production. *J. Immune*, 141: 2407-2412.

Draper, H. and Hadley, M. (1990): Malondialdehyde determination as index of lipid peroxidation. *Methods Enzymol.* 186: 421-431.

El Mallah, M. (2015): Influence of yerba mate tea (*Ilex paraguariensis*) in improving some lipolytic enzymes of high-fat diet-induced obese rats. *Egypt. J. of Nutrition and Health*, 10: 27-48.

El-Desouky, M., Mahmoud, M., Riad, B. and Taha, Y. (2019): Nephroprotective effect of green tea, rosmarinic acid and rosemary on N-diethylnitrosamine initiated and ferric nitrilotriacetate promoted acute renal toxicity in Wistar rats. *Interdiscip. Toxicol.*, 12: 98-110.

El-Kholie, E., Afifi, T. and Abdelal, N. (2022): Potential Anti-Obesity Effects of Matcha Tea in Rats Fed on a High Fat Diet. *JHE.*, 32: 94-108.

Engin, A., Tsatsakis, A., Tsoukalas, D. and Engin, A. (2018): Do flavanols-rich natural products relieve obesity-related insulin resistance? *Food Chem. Toxicol.*, 112: 157-167.

Ferreira, M., Gomes, A. and de Moraes, A. (2017): Green tea extract outperforms metformin in lipid profile and glycaemic control in overweight women: a double-blind, placebo-controlled, randomized trial. *Clin. Nutr.*, 22: 1-6.

Friedewald, W., Leve, R. and Fredrickson, D. (1972): Estimation of the concentration of low-density lipoprotein cholesterol in plasma without use of the preparative ultracentrifuge. *Clin Chem*, 18: 499-502.

Halliwell, B. and Gutteride, J. (2007): Free radicals in biology and medicine (4th ed.). New York: Oxford University Press Inc. (Chapters 4 and 10).

Han, X., Choi, S., Men, X., Lee, S., Jin, H., Oh, H., Kim, E., Kim, J., Lee, B. and Lee, O. (2022): Anti-obesity activities of standardized ecklonia stolonifera extract in 3T3-L1 preadipocytes and high-fat-diet-fed ICR mice. *Applied Sciences*, 12: 5115.

Hasanein, A. Gawad, H. and El-Megeid, A. (2012): Effect of water extract prepared from green tea, black tea and cinnamon on obese rats suffering from diabetes, *World Applied Sciences Journal*, 20: 976-987.

Hena-Mejia, J., Elinav, E., Jin, C. and Hao, L. (2012): Inflammasome-mediated dysbiosis regulates progression of NAFLD and obesity. *Nature*, 482: 179-185.

Hissin, P. and Hilf, R. (1970): A fluorometric method for determination of oxidized and reduced glutathione in tissues. *Anal Biochem*, 74: 214-226.

Hussein, A. and Mustafa, N. (2024): Impact of a high-fat diet on dyslipidemia and gene expression of low-density lipoprotein receptors in male rats. *Iraqi Journal of Veterinary Sciences*, 38: 133-138.

Jaiswal, R., Sovdat, T. and Vivan, F. (2010): Profiling and characterization by LC-MSn of the chlorogenic acids and hydroxycinnamoylshiki-mate esters in mate (*Ilex paraguariensis*). *J Agric Food Chem.*, 58: 5471-5484.

Jeszka-Skowron, M., Krawczyk, M. and Zgola-Grzeskowiak, A. (2015): Determination of antioxidant activity, rutin, quercetin, phenolic acids and trace elements in tea infusions: Influence of citric acid addition on extraction of metals. *J. Food Compos. Anal.*, 40: 70-77.

Jiang, S., Liu, H. and Li, C. (2021): Dietary Regulation of Oxidative Stress in Chronic Metabolic Diseases. *Foods*, 10: 1854.

Karri, S., Sharma, S., Hatware, K. and Patil, K. (2019): Natural anti-obesity agents and their therapeutic role in management of obesity: A future trend perspective. *Biomed. Pharmacother.*, 110: 224-238.

Katarzyna, J., Karolina, J., Agnieszka, L., Irena, B., Ewa, R. and Karolina, D. (2020): Effect of the Yerba mate (*Ilex paraguariensis*) brewing method on the content of selected elements and antioxidant potential of infusions J of Chemical. Technology, 22: 54-60.

Klein, G., Stefanuto, A., Boaventura, B., de Morais, E., Cavalcante Lda, S., de Andrade, F., Wazlawik, E., Di Pietro, P., Maraschin, M. and da Silva, E. (2011): Mate tea (*Ilex paraguariensis*) improves glycemic and lipid profiles of type 2 diabetes and pre-diabetes individuals: a pilot study. J Am Coll Nutr., 30:320-32.

Kolackova, T., Kolofikova, K., Sytarova, I., Snopek, L., Sumczynski, D. and Orsavova, J. (2020): Matcha Tea: Analysis of Nutritional Composition, Phenolics and Antioxidant Activity. Plant Food Hum. Nutr., 75: 48-53.

Lasker, S., Rahman, M., Parvez, F., Zamila, M., Miah, P., Nahar, K., Kabir, F., Sharmin, S., Subhan, N., Ahsan, G. and Alam, M. (2019): High-fat diet-induced metabolic syndrome and oxidative stress in obese rats are ameliorated by yogurt supplementation. Sci Rep., 27: 1-15.

Li, Y., Rahman, S., Huang, Y., Zhang, Y., Ming, P., Zhu, L., Chu, X., Li, J., Feng, S. and Wang, X. (2020): Green tea polyphenols decrease weight gain, ameliorate alteration of gut microbiota, and mitigate intestinal inflammation in canines with high-fat-diet-induced obesity. J. Nutr. Biochem., 78.

Lindholm, C., Ertel, R., Bauwens, J., Schmuck, E., Mulligan, J. and Kurt, W. Saupe, K. (2013): A high-fat diet decreases AMPK activity in multiple tissues in the absence of hyperglycemia or systemic inflammation in rats. J Physiol Biochem., 69: 165-175.

Liu, M., Ling, S., Yin, L., Stephen, C. W., Randy, J., David, D. and Patrick, T. (2004): Obesity induced by a high-fat diet downregulates apolipoprotein A-IV gene expression in rat hypothalamus. Am. J. Physiol. Endocrinol Metab., 287: 366-370.

Machado, M., Banin, R., Thomaz, F., de Andrade, I., Boldarine, V., Figueiredo, J., Hirata, B., Oyama, L., Lago, J., Ribeiro, E. and Telles, M. (2021): Ginkgo biloba extract (gbe) restores serotonin and leptin receptor levels and plays an antioxidative role in the hippocampus of ovariectomized rats. Molecular Neurobiology volume, 58: 2692-2703.

Martínez-Sánchez, N. (2020): There and back again: Leptin actions in white adipose tissue. *Int. J. Mol. Sci.*, 21: 6039.

Martins, F., Suzan, A., Cerutti, S., Arçari, D., Ribeiro, M. and Bastos, D. (2009): Consumption of mate tea (*Ilex paraguariensis*) decreases the oxidation of unsaturated fatty acids in mouse liver. *Br J Nutr.*, 101: 527-32.

Milena, J. (2003): Predrag Djurdjevic and DejanStankov. *JSCS*. 68(8-9): 691-698.

Nakagawa, T. and Yokozawa, T. (2002): Direct scavenging of nitric oxide and superoxide by green tea. *Food and Chemical Toxicology*, 40: 1745-1750.

Naomi, R., Teoh, S., Embong, H., Balan, S., Othman, F., Bahari, H. and Yazid, M. (2023): The Role of Oxidative Stress and Inflammation in Obesity and Its Impact on Cognitive Impairments—A Narrative Review. *Antioxidants*, 12: 1-20.

Newairy, A. and Abdou, H. (2009): Protective role of flax lignans against lead acetate induced oxidative damage and hyperlipidemia in rats. *Food chem. Toxicol.*, 47: 813-818.

Ngoc, L., Man, H., Besselink, H., Cam, H., Brouwer, A. and Burg, B. (2019): Identification of PPAR-activating compounds in herbal and edible plants and fungi from Vietnam. *Ind. Crops Prod.*, 129: 195-200.

Nguyen, J., Killcross, A. and Jenkins, T. (2014): Obesity and cognitive decline: Role of inflammation and vascular changes. *Front. Neurosci.*, 8: 375.

Pang, J., Choi, Y. and Park, T. (2008): *Ilex paraguariensis* extract ameliorates obesity induced by high-fat diet: potential role of AMPK in the visceral adipose tissue. *Arch Biochem Biophys*, 476: 178–185.

Pérez-Torres, I., Castrejón-Téllez, V., Soto, M., Rubio-Ruiz, M., Manzano-Pech, L. and Guarner-Lans, V. (2021): Oxidative Stress, Plant Natural Antioxidants, and Obesity. *Int. J. Mol. Sci.* 2021, 22: 1-26.

Pimentel, G., Lira, F., Rosa, J., Caris, A., Pinheiro, F., Ribeiro, E., do Nascimento, C. and Oyama, L. (2012): Yerba mate extract (*Ilex paraguariensis*) attenuates both central and peripheral inflammatory effects of diet-induced obesity in rats. *J. Nutr. Biochem.*, 4: 16.

Reeves, R., Nielsen, F. and Fahey, G. (1993): AIN-93 Purified Diets for Laboratory Rodents. *J. Nutr.*, 123: 1939-1951.

Richmond, N. (1973): Colorimetric determination of total cholesterol and high-density lipoprotein cholesterol (HDL-c). *Clin. Chem.*, 19: 1350-1356.

Roy, S. (1970): colorimetric method of serum alkaline phosphatase. *Journal of Clinical Chemistry*, 16: 431-432.

Samarghandian, S., Borji, A., Afshari, R., Delkhosh, M. and gholami, A. (2013): The effect of lead acetate on oxidative stress and antioxidant status in rat bronchoalveolar lavage fluid and lung tissue. *Toxicology Mechanisms and Methods*, 23: 432-436.

Sander, G., Konig, A., Wallner, M. and Weghuber, J. (2020): Functional foods-dietary or herbal products on obesity: Application of selected bioactive compounds to target lipid metabolism. *Curr. Opin. Food Sci.*, 34: 9-20.

Shahin, K. and Aburaya, S. (2023): The Effect of Arabic Gum and High-Fat Diets on Obese Rats. *J Home Econ.*, 33: 103-111.

Singla, P., Bardoloi, A. and Parkash, A. (2010): Metabolic effects of obesity: a review. *World J. Diabetes*, 1: 76-88.

Sokary, S., Al-Asmakh, M., Zakaria, Z. and Bawadi, H. (2023): The therapeutic potential of matcha tea: A critical review on human and animal studies. *Curr Res Food Sci.*, 6: 1-10.

SPSS (1986): Statistical package for social science, version 19. SPSS Inc., II. U.S.A.

Sutton, J. (2004): New tastes in green tea: A novel flavor for familiar drinks, dishes, and desserts. *Libr. J.*, 129: 110.

Taketa, A., Gnoatto, G., Gosmann, V., Schenkel, P. and Guillaume, D. (2004): Triterpenoids from Brazilian *Ilex* species and their in vitro antitrypanosomal activity. *Journal of Natural Products* (67): 1697-1700.

Wahlefeld, A. (1974): Methods of Enzymatic Analysis. Academic Press, Chapter, 5: 1831-1835.

Wahyuni, E., Maryatun, M., Veri, N., Susilawati, E., Firrahmawati, L., Sri Wahyuni, E. and Wulandari, R. (2023): Green Tea Extract has a Protective Effect on Leptin and Lipid Profile Levels Due to the Induction of Depot Medroxyprogesterone Acetate. *Med Arch.*, 77: 173-177.

Willems, M., Sahin, M. and Cook, M. (2018): Matcha Green Tea Drinks Enhance Fat Oxidation During Brisk Walking in Females. *Int. J. Sport Nutr. Exerc.* 28: 536-541.

Xing, L., Zang, H., Qi, R., Tsao, R. and Mine, Y. (2019): Recent advances in the understanding of the health benefits and molecular mechanisms associated with green tea polyphenols. *J. Agric. Food Chem.*, 67: 1029-1043.

Xu, P., Ying, L., Hongb, G. and Wang, Y. (2016): The effects of the aqueous extract and residue of Matcha on the antioxidant status and lipid and glucose levels in mice fed a high-fat diet. *Food Funct.*, 7: 294-300.

Xu, R., Yang, K., Li, S., Dai, M. and Chen, G. (2020): Effect of green tea consumption on blood lipids: a systematic review and meta-analysis of randomized controlled trials. *Nutrition Journal*, 19 48: 2-15.

Yamabe, N., Kang, K., Hur, J. and Yokozawa, T. (2009): Match, a powdered green tea, ameliorates the progression of renal and hepatic damage in type 2 diabetic OLETF rats. *J. Med. Food*, 12: 714-721.

Young, D. (2001): Effect of disease on clinical lab Tests, 4th ed. AACC press.

Zenaro, L., Andrade, L., Santos, P. and Locatell, C. (2014): Effects of Aqueous Extract of Yerba Mate (*Ilex Paraguariensis*) on the Oxidative Stress in Rats Fed a Cafeteria Diet. *International Journal of Natural Sciences Research*, 2: 30-43.

Zhang, Y., Proenca, R., Maffe, i., M., Barone, M., Leopold, L. and Friedman, J. (1995): positional cloning of the mouse obese gene and its human homologue, *Nature*. 72: 425-432.

Zheng, G., Sayama, K., Okubo, T., Juneja, L. R. and Oguni, I. (2004): Anti-obesity effects of three major components of green tea, catechins, caffeine and theanine, in mice. *In Vivo*, 18: 55-62.

Zhou, J., Lin, H., Xu, P., Yao, L., Xie, Q., Mao, L. and Wang, Y. (2020): Matcha green tea prevents obesity-induced hypothalamic inflammation via suppressing the JAK2/STAT3 signaling pathway. *Food Funct.*, 11: 8987-8995.

Zhou, J., Yu, Y., Ding, L., Xu, P. and Wang, Y. (2021): Matcha Green Tea Alleviates Non-Alcoholic Fatty Liver Disease in High-Fat Diet-Induced Obese Mice by Regulating Lipid Metabolism and Inflammatory Responses. *Nutrients*, 13: 1-12.

الملخص العربي

تأثير المستخلص المائي لشاي الماتشا وشاي المته على حالة إجهاد تأكسدي في الفئران المصابة بالسمنة

سوسن عادل نور، مايسة محمد الملاح، هاني جابر المصري
قسم التغذية وعلوم الأطعمة - كلية الاقتصاد المنزلي - جامعة حلوان

استهدف هذا البحث دراسة تأثير المستخلص المائي لشاي الماتشا والمته على حالة الإجهاد التأكسدي الفئران المصابة بالسمنة. تم إجراء التجربة على عدد 35 من ذكور الفئران البالغة البيضاء بشكل عشوائي إلى 7 مجموعات متساوية كل منها 5 فئران على النحو التالي: المجموعة 1: الضابطة السالبة، تم تغذيتها على نظام غذائي أساسي. أعطيت المجموعات 2-7 خللات الرصاص (200 ملجم / لتر) في مياه الشرب لمدة 4 أسابيع للبحث على الإجهاد التأكسدي. ثم أصبحت الفئران بدينة عن طريق التغذية على غذاء عالي في محتواه من الدهون لمدة 4 أسابيع. تم الاحتفاظ بالمجموعة 2 كمجموعة ضابطة موجبة. تم إعطاء المجموعتين 3 و 4 عن طريق الفم مع 0.5 و 1 مل من المستخلص المائي لشاي المته، في حين تم إعطاء 0.5 و 1 مل من المستخلص المائي لشاي الماتشا للمجموعتين 5 و 6، على التوالي. تم إعطاء المجموعة 7 عن طريق الفم مع 1 مل من خليط المستخلص المائي لشاي المته والماتشا لمدة 4 أسابيع. في نهاية التجربة، تم ذبح الفئران وجمع المصل لإجراء التحاليل البيوكيميائية. أظهرت النتائج أن المجموعة الضابطة الموجبة أدى إلى ارتفاعات ملحوظة في زيادة وزن الجسم، ونسبة كفاءة التغذية، ونسبة الدهون البريتونية، والكوليسترول الكلي في الدم، والدهون الثلاثية، وكوليسترول البروتين الدهني منخفض الكثافة، كوليسترول البروتين الدهني منخفض الكثافة بشدة (VLDL-C)، الليبتين، أكسيد النيتريك، المالونديالدهيد، انزيمات الكبد (AST, ALT, ALP)، حمض البوليك، اليوريا ومستويات الكرياتينين بالمقارنة إلى المجموعة الضابطة السالبة، وأيضا انخفضت مستويات الكوليسترول الدهني عالي الكثافة في الدم، ونشاط الجلوتاثيون المختزل بشكل ملحوظ، من ناحية أخرى، أدى تناول مستخلص المته والماتشا ومزيجهما عن طريق الفم إلى تخفيف هذه التأثيرات الضارة و التغيرات البيوكيميائية مقارنة بالمجموعة الضابطة الموجبة. في الختام، يُظهر شاي المته والماتشا نشاطاً مضاداً للأكسدة وخافضاً لدهون الدم وهو فعال في تقليل وزن الجسم لدى الفئران السمنة. توصي الدراسة بأن استهلاك شاي المته والماتشا قد يكون مفيداً للأفراد الذين يعانون من السمنة وفرط دهون الدم.

الكلمات المفتاحية: شاي الماتشا، شاي المته، أسيتات الرصاص، الإجهاد التأكسدي، السمنة، الفئران.