Phytochemical Comparison Study for Evaluating the Hypolipidemic Effect Between Two Iraqi Pepper spp. in the Rats Model

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https://dx.doi.org/10.13005/bpj/2580

(Received: 03 October 2022; accepted: 14 November 2022)

Hypercholesterolemia is a predominant risk factor for atherosclerosis and cardiovascular disease (CVD). The World Health Organization (WHO) recommended reducing the intake of cholesterol and saturated fats. On the other hand, limited evidence is available on the benefits of vegetables in the diet to reduce these risk factors, so this research was conducted to compare the hypolipidemic effect between the extracts of two different types of Iraqi peppers, the fruit of the genus Capsicum traditionally known as red pepper extract (RPE), and Piper nigrum as black pepper extract (BPE), respectively, in different parameters and histology of the liver of the experimental animals. The red pepper was extracted by ethyl acetate, while the black pepper was extracted by 90% ethanol, then each extract was concentrated for further analysis. Reversed-phase high-performance liquid chromatography (RP-HPLC) results confirm the presence of a high percentage of alkaloids responsible for hypolipidemic activity studied against hypercholesterolemic rats divided into five rats into six groups for ten weeks. We measured different parameters such as the lipid profile, the oxidized low-density lipoprotein (LDL), and the serum level of the most important parameters of the kidney and liver function tests. We also compared body weight after ten weeks. This study showed a slight or nonsignificant decrease in cholesterol, TG, very low-density lipoprotein VLDL, low-density lipoprotein LDL, and oxidized LDL with a considerable increase in high-density lipoprotein HDL in group V (high-fat diet + black pepper extract). Study results were improved by histological changes induced in liver tissue. In conclusion, the results focused on consuming these vegetables and reducing cholesterol levels, as it is considered a risk factor for cardiovascular disease (CVD).

Keywords: Hypercholesterolemia; Histopathology; Lipid profile; Pepper extract; RP-HPLC.
must approve this to confirm its safety and meet the world population’s demand. Therefore, developing new and cost-effective herbal medications, such as cholesterol-lowering drugs, is preferable.

The fruit of the genus Capsicum has beneficial phytochemical compounds, including Capsaicinoids in red pepper as aromatic components [Fig. 1]. Among them, capsaicin (8-methyl-N-vanillyl-trans-6-none amid) and dihydrocapsaicin are the main pungent components; these compounds are now widely used in health foods and medical products. In addition, black pepper (Piper nigrum Linn. Piperaceae), known as the ‘King of spices’ in folklore, is used worldwide. Traditionally, black pepper was used to treat hyperlipidemia and coronary heart disease. Black pepper contains essential oils, piperine, chavicine, piperidine, and piperidines [Fig. 2] Piperine (1-piperoyl piperidine) is the main active constituent as an alkaloid containing nitrogen present in the fruits of many species of piper such as long pepper (Piper longum), black pepper (Piper nigrum), and others. In general, Piperine is considered safe by the United States Food and Drug Administration (FDA).

This research aimed to study the hypolipidemic effect of the main active constituents in two varieties of pepper red and black pepper extracts on the lipid profile, serum urea, creatinine, and aspartate transaminase (AST) and alanine transaminase (ALT), in addition to body weight.

MATERIALS AND METHODS

Materials
Plant materials and collection
Two varieties of pepper: one is the fresh red chili pepper (Capsicum annum, L.), and the second is the black pepper seeds (Piper nigrum Linn.). The red pepper was harvested from a green farm in the city of Youssoufia, June-July 2020, at full maturity, washed with tap water, and dried at room temperature in a shaded area for two weeks. At the same time, black pepper was bought from the market. Each was mixed to a fine powder with a blender, packed in perforated plastic bags stored in a dry, cool place until extraction.

Chemicals
Cholesterol powder 500 mg (biochemical) BDH Chemicals Ltd Poole England, 10 mg tablet (micro-Labs Limited, India), All chemicals and reagents used were high quality. The diagnostic kits used were obtained from (Randox Laboratories Ltd) for the estimation of cholesterol, HDL, LDL, and VLDL, while the Rat Oxidized Low-Density Lipoprotein (Ox-LDL) ELISA Kit from (Cusabio-chain).

Experimental animals
Thirty adults, Wister Albino rats of both sexes weighing 200–250 g, were purchased from the laboratory animal house at Baghdad’s pharmacy.

Fig. 1. The structures of capsaicinoids. Adopted from (12,13)
college/university. Kept under specific pathogen-free conditions, provided with water and food ad libitum during a 12-hour light-dark cycle, and kept during the study at regular room temperature (18-21°C). Animals were handled throughout this process following the Declaration of Helsinki, which the University approved by Baghdad’s ethics committee for using animals in research. The Institutional Animal Ethics Committee accepted the experiment protocol.

Methods

Preparation of Plant Extract
About 2 kg of mature *Capsicum annum* (bird’s eye chili) gives 160 g of dry powder extracted by cold maceration with 100% ethyl acetate for 5 days, twice. After filtration, the clear filtrate is concentrated by a rotary evaporator for complete dryness to give 4 g of red pepper crude extract. At the same time, 150 g of dry black pepper seeds were extracted with 90% ethanol using the Soxhlet apparatus until exhausted. After filtration, the clear filtrate was concentrated by a rotary evaporator to give about 20.7 g of black pepper crude extract.

**RP-HPLC condition**
HPLC conditions for Capsaicin: C18 column (1506 mm, 5 µm).

**Gradient Elution**
A: 1% Acetic acid: Acetonitrile (50:50); B: Acetonitrile;

The program: (A/B), (100/0) from 0 min to 15 min; then (0/100) from 15.05 min to 20.00 min; 20.05 min (100/0) to 35.5 min; the injection volume 20 uL, at flow rate 1.0 ml/min, column temp 40°C; at wavelength 280 nm.

HPLC conditions for piperine: RP-HPLC was performed using methanol and water as mobile phase. The detection and quantification were performed at a wavelength of 345 nm.

Fig. 2. The structures of the main active constituents in black pepper adopted from (12,13)
Animals and diet
Thirty healthy rats weighing up to 200-250 g were used in the present study between March and May 2021. The animals were placed in cages and exposed to constant environmental conditions, and a high-fat diet was prepared by adding 1.5 g/kg to the daily animal diet. The plant extract was given by oral gavage to the animals, each one alone or in combination with two combinations of plant extracts.

The animals were divided into six groups, each of five animals:
(I): Control group either D.W. or normal saline only.

Fig. 3. HPLC chromatogram of the Capsicum annum fruit extract-analysis of capsaicinoids using the reversed-phase column under optimal conditions, and both capsaicin (tR 6.5 min) and dihydrocapsaicin (tR 7.7 min) were separated.

Fig. 4. HPLC chromatogram of Piper nigrum Linn seed extract. Piper nigrum Linn seed extract at 345nm, the standard piperine peak (tR 7.19 min).
Blood samples

Under control conditions, the analysis was performed at about 25°C. Two milliliters (2mL) of blood were drawn by heart puncture from each rat in the experiment after two weeks and ten weeks. The sample was collected in a plain tube, left for 15 min. for clotting at room temperature, then centrifuged for 15 min. at 3000 rpm for serum separation and kept at -20°C refrigerated for further analysis.

Biochemical analysis

Oxidized LDL was determined by enzyme-linked immunosorbent assay (ELISA) (Cusabio, China). At the same time, blood lipid profile, urea, creatinine, ALT, and AST were performed simultaneously after serum separation using the spectrophotometric method SPEC ORD 40 (Analytic Jena, Germany) at the Baghdad College of Medical Sciences labs.

Statistical analysis

Statistical analysis depended on mean±standard deviation (mean ±SD) and the Social Sciences statistical package (SPSS version 23.0). In contrast, the difference between the two means was assessed using an independent student t-test.

Histological Examination

The animals were anesthetized by cervical dislocation. Liver tissues were quickly removed from rats, fixed in a 4% paraformaldehyde solution, and then embedded in paraffin. Sections were obtained and stained with hematoxylin and eosin (H&E) for histological examination under a microscope. The samples were treated with an ultrathin microtome and stained with uranyl citrate-lead citrate. Finally, the ultrastructure of the hepatocytes was observed and photographed with a transmission electron microscope.
RESULTS

Chromatographic results

Optimal chromatographic conditions should be used to obtain a good resolution of adjacent peaks within a short analysis time. Using the reversed-phase column, various mobile phases have been used to analyze capsaicinoids and piperine. For qualitative and quantitative analysis of the main active constituents in the plant extract, the high-performance liquid chromatography (HPLC) technique depends on the tR of the peak. The area under the peak depends on the concentration of the active phytochemical constituents in the sample, as shown in Figure (3) for the analysis of capsaicinoids using the reversed-phase column under optimal conditions, both capsaicin (tR 6.5 min) and dihydrocapsaicin (tR 7.7 min) were separated. Since capsaicin and dihydrocapsaicin are very similar in molecular structures, the maximum absorption wavelengths determined are nearly the same and are found to be 280.8 and 279.6 nm, respectively. Figure (4) HPLC chromatogram of the Piper nigrum Linn seed extract at 345nm, the standard piperine peak (tR 7.19 min).

Effects on Serum Lipid Profiles

Table (1) of the present study did not show significant differences in serum levels of (TG, VLDL, LDL, cholesterol, and oxidized LDL) s. TG level s. VLDL, s. LDL s. cholesterol and s. oxidized

Table 3. Kidney Function Test (KFT) & Liver Function Test (LFT) parameters of the animal groups after the second week feeding

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control</th>
<th>HFD</th>
<th>HFD + atorvastatin</th>
<th>HFD + black pepper</th>
<th>HFD + red pepper</th>
<th>HFD + comb</th>
</tr>
</thead>
<tbody>
<tr>
<td>Urea (mg/dl)</td>
<td>35.6 ± 5.9</td>
<td>19.2 ± 4.1</td>
<td>16.0 ± 3.7</td>
<td>32.8 ± 5.1**</td>
<td>29.1 ± 2.8**</td>
<td>45.4 ± 4.7**</td>
</tr>
<tr>
<td>Creatinine (mg/dl)</td>
<td>0.4 ± 0.06</td>
<td>0.4 ± 0.05</td>
<td>0.4 ± 0.05</td>
<td>0.4 ± 0.05</td>
<td>0.3 ± 0.04</td>
<td>0.4 ± 0.05</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td>41.0 ± 2.9</td>
<td>39.5 ± 2.8</td>
<td>50.0 ± 13.8</td>
<td>43.2 ± 4.7</td>
<td>40.0 ± 3.9</td>
<td>39.5 ± 2.8</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td>115.6 ± 5.1</td>
<td>121.5 ± 11.4</td>
<td>156.4 ± 33.5</td>
<td>127.0 ± 10.9</td>
<td>158.1 ± 48.8</td>
<td>121.0 ± 11.3</td>
</tr>
</tbody>
</table>

* compared to control rats; ** compared to rats fed a high-fat diet. *P < 0.05, **P < 0.01 and ***P < 0.001

Table 4. Kidney Function Test (KFT) & Liver Function Test (LFT) parameters of the animal groups after the 10th week of feeding

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control</th>
<th>HFD</th>
<th>HFD + atorvastatin</th>
<th>HFD + black pepper</th>
<th>HFD + red pepper</th>
<th>HFD + comb</th>
</tr>
</thead>
<tbody>
<tr>
<td>Urea (mg/dl)</td>
<td>41.6 ± 6.5</td>
<td>41.8 ± 3.4</td>
<td>31.8 ± 5.1***</td>
<td>22.0 ± 5.4***</td>
<td>18.0 ± 2.1***</td>
<td>21.9 ± 3.7***</td>
</tr>
<tr>
<td>Creatinine (mg/dl)</td>
<td>0.34 ± 0.05</td>
<td>0.34 ± 0.05</td>
<td>0.36 ± 0.05</td>
<td>0.36 ± 0.05</td>
<td>0.28 ± 0.04</td>
<td>0.3 ± 0.07</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td>45.2 ± 3.6</td>
<td>43.2 ± 3.6</td>
<td>36.5 ± 3.00</td>
<td>35.2 ± 11.7</td>
<td>42.5 ± 7.7</td>
<td>43.7 ± 11</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td>113.5 ± 10.3</td>
<td>136.9 ± 17.3</td>
<td>163.6 ± 44.5</td>
<td>135.7 ± 25.3</td>
<td>98.1 ± 20.0</td>
<td>127.9 ± 28.6</td>
</tr>
</tbody>
</table>

* compared to control rats; ** compared to rats fed a high-fat diet. *P < 0.05, **P < 0.01 and ***P < 0.001

Table 5. Reduced body weight of animal groups after 10 weeks of feeding

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>Body weight (Kg) (Mean ± STD)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control vs HFD</td>
<td>269.8 ± 11.4 Vs 310.3 ± 27.2</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>HFD vs atorvastatin</td>
<td>310.3 ± 27.2 Vs 274.9 ± 13.7</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>HFD vs black pepper</td>
<td>310.3 ± 27.2 Vs 242.5 ± 4.3</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>HFD Versus red pepper</td>
<td>310.3 ± 27.2 Vs 260.4 ± 7.8</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>HFD vs. combination</td>
<td>310.3 ± 27.2 Vs 244.6 ± 7.0</td>
<td>P &lt; 0.001</td>
</tr>
</tbody>
</table>
LDL between the HCD group and all other animal groups. However, there is a significant increase in s. cholesterol and HDL in the (HCD +BP) and (HCD +statin) groups, compared to the HCD group (p < 0.05) after the second week of feeding.

At the end of the research, the serum level of lipid profiles (s.TG, s. cholesterol, s. LDL, s. VLDL, oxidized LDL) of the group (HCD + RP) and the group (HCD combination of BP&RP) group (p< 0.05) significantly decreased in comparison to HCD group.

In addition, there is a significant decrease in the serum level of lipid profiles (s.TG, s. cholesterol, s. LDL, s. VLDL) and a highly significant decrease in the serum level of oxidized LDL of the group (HFD +BP) group as compared to the HFD group as shown in Table (2).

In Table (2), the HDL level is significantly increased in both groups (the HFD +RP), and (the HFD +combination of BP&RP) groups (p< 0.05) after the 10th week of feeding compared to the HFD group. At the same time, there is no significant
increase in the group (HFD +BP) group (p > 0.05) as compared to the HFD group. Furthermore, there is a significant decrease in serum lipid profiles (s.TG, s. cholesterol, LDL-c, s. VLDL, s. HDL) and oxidized LDL of the group (HFD +statin) group (p < 0.001), with a nonsignificant decrease in HDL-c level (p > 0.05) as compared to the HFD group after 10 weeks of feeding.

After the second feeding, there was a significant increase in the KFT parameter (serum urea) in all animal groups (p < 0.05), except the group (HCF +statin) group, which showed no significant change in serum urea (p > 0.05) as compared to the HCF group, as shown in Table 3.

After the 10th week of feeding, there was a highly significant decrease in the KFT parameter (serum urea) in all animal groups (p < 0.001) except between the control group and HCF, as shown in Table 4.

There are no significant changes in the KFT parameter (serum creatinine) (p > 0.05) in all animal groups after the second and 10th week of feeding, as shown in Tables 3 and 4, respectively.

The present study did not show significant changes in LFT (Serum levels of AST and ALT) of the animal groups compared to the HFD group (p > 0.05) after the second and 10th weeks of feeding, as shown in Tables 3 and 4. Only the group (HFD +atorvastatin) group showed a significant decrease in serum ALT levels (p < 0.05), and the (HFD + RB) group showed a significant decrease in serum AST levels (p < 0.05) as compared to the group HFD.
after 10th week of feeding, as shown in Table 4.

Body weight in all animal groups is significantly reduced after the 10th week of feeding compared to the HFD group (p<0.001), as shown in Table 5.

There is a highly significant decrease in serum cholesterol and serum between the 2nd and 10th week of the feeding of the group (HFD + BR) group, (HFD + RP) group, and group (HFD + Combination) group (p < 0.01). Furthermore, there is a significant decrease in serum cholesterol between the second and tenth weeks of the feeding of the (HFD + statin) group (p<0.05), as shown in figure (5).

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**Fig. 9.** Changes in serum LDL level in different animal groups between 2nd and 10th weeks of feeding. *P < 0.05, **P < 0.01 and ***P < 0.001

**Fig. 10.** Changes in serum oxidized LDL levels in different animal groups between the 2nd and 10th week of feeding. *P < 0.05, **P < 0.01 and ***P < 0.001.
Although serum TG & serum VLDL were a highly significant decrease between the second week and 10th week of the feeding of the (HFD + RP) group (p< 0.001) & a significant decrease in serum levels (p< 0.05) between the second week and 10th week of feeding of the (HFD + combination) group (p < 0.01) as shown in figure (6) & figure (7) respectively.

The present study showed a significant increase in serum HDL between the 2nd week and the 10th week feeding of the group (HFD + BP) group and (HFD + statin) (p<0.05), as shown in figure (8).

Both serum LDL and serum oxidized LDL levels showed a highly significant decrease between the second week and 10th-week feeding of the group (HFD + RP) group (p< 0.001), while the (HFD+ combination) group showed a significant decrease in serum LDL between the second week and 10th-week feeding, in addition, the group (HFD + BP) group showed a significant decrease in serum oxidized LDL (p<0.05) as shown in figure (9) & (10) respectively.

**Histopathological examination**

The effects of the plant extract on the ultrastructure of liver tissue and the analysis of the surface area of the lipid droplets by microscopically examining the histopathological sections of liver tissue are shown in Figs. (11-16). In the control group, the hepatic lobular structure was intact.
and clear. There was a hepatic cord of arranged hepatocytes. In the cytoplasm, the endoplasmic reticulum (ER) was dense and clear, while in the hepatocytes, a few lipid droplets with a large number of mitochondria did not show swelling. (Fig.11). While in the HFD group, liver cells had partly necrosis, liver tissues were disordered, and there were a large number of lipid droplets but a small number of mitochondria in the cytoplasm. The mitochondria were swelling and few ER or glycogen granules were found in the liver cells. After treatment with pepper extract, liver tissues returned to normal status. In the cytoplasm, the lipid droplets reduced significantly, and the mitochondria were rich but still swelling. At the same time, the number of ER and glycogen granules in the cytoplasm increased clearly (Figs.12-16). The results suggested that the plant extract reduced fatty liver and lipid droplets, repaired liver damage caused by a high-fat diet, and increased the number of mitochondria and ER in the cytoplasm.

**DISCUSSION AND CONCLUSION**

Plant extract is usually a mixture of different classes of organic compounds soluble in the solvents used\(^{11}\). Since many factors play an important role in the extraction of capsaicinoids and piperine in Capsicum and piper plants, such as the extraction method, its chemical nature, the size of the sample particles, the storage time, and the presence of impurities\(^{12}\). The medicinal effects of chilies and peppers are related to different constituents such as capsaicin, fixed oil, thiamine, and others\(^{13}\) which are responsible for a hot sensation in the tongue\(^{14}\) and are used for the treatment of many disorders such as hyperlipidemia and obesity\(^{15}\).

The main reason for dyslipidemia during the HFD period is that hepatic tissue increases TG formation by the conversion of glucose to FFA, then transported by the bloodstream as VLDL-C and stored as fat in adipose tissue. In experimental animal studies, long-term exposure to HFD resulted in hyperlipidemia, leading to increased visceral fat deposition by facilitating the development of a positive energy balance, leading to abdominal obesity\(^{16}\).

According to the Hwan-Hee Jang et al. 2020 study, the first two weeks included in this study are considered an acclimatization period. Therefore, there were no significant differences in serum s.TG level s. VLDL, s. LDL s. cholesterol and s. oxidized LDL between the (HFD) group, and all other animal groups except the (HFD +BP) group showed a significant increase in s. cholesterol, compared to the HFD group (p < 0.05). HDL increased significantly in the group (HFD +statin) group only after the second week of
feeding compared to the control group, as shown in table (1)\textsuperscript{17}.

As shown in Table (2) of the present study, the effect of red pepper consumption on lipid profile after the 10th week of feeding was observed may be due to capsaicinoids, which are a group of pungent compounds found in chilies, contain capsaicin, dihydrocapsaicin, nor-dihydrocapsaicin, homodihydrocapsaicin, and homocapsaicin. Furthermore, the hypocholesterolemia effect of red pepper may be due to a combined effect of capsaicinoids with other phytochemicals such as vitamins (ascorbic acid, tocopherol, carotenoids) and phenolic compounds present in chili\textsuperscript{18}. A previous animal study demonstrating the hypolipidemic effect of red pepper in rats on a high-fat diet for 8 weeks showed a significant decrease in serum TG level\textsuperscript{19}.

Another two previous animal studies in Wistar rats showed a significant decrease in serum total cholesterol, TG, LDL-c, and significantly elevated serum HDL-c\textsuperscript{20-21}. Recently 2020, Al-Jumayi et al. The results of their study introduced scientific evidence that red chili pepper ethanol extract possesses antiobesity potentials and hypolipidemic effects through decreased levels of TG, total cholesterol, LDL, and VLDL with increased HDL levels and weight loss of high-fat dieted female albino Wistar rats\textsuperscript{16}.

Furthermore, there was a highly significant decrease in serum cholesterol and serum between the 2nd and 10th weeks of the feeding of the group (HFD + BR) group, (HFD + RP) group, and group (HFD + Combination) group (p < 0.01). Furthermore, there is a significant decrease in serum cholesterol between the second and tenth weeks of the feeding of the (HFD + statin) group (p<0.05), as shown in figure (5).

The possible explanation for the hypolipidemic effect of capsaicin, which is the active component of red pepper, may be related to reduced intestinal absorption of cholesterol and increased excretion of cholesterol and bile acid in the feces or due to activation of the peroxisome proliferator-activated receptor á (PPARá)\textsuperscript{22}. It was also known that lecithin, one of the components of chili seeds, is a natural fat emulsifier that prevents it from settling in the arteries by keeping cholesterol suspended in the blood\textsuperscript{23}.

Another study revealed the efficacy of capsaicinoids in decreasing total plasma cholesterol and improving the lipid profile by decreasing cholesterol absorption and reducing aortic plaque under high-cholesterol-fed conditions\textsuperscript{24}.

According to the results of the study mentioned in Table (2), the possible explanation for the hypolipidemic effect of pepper extracts is that the lipolytic enzymes and the up-regulated mRNA expressions increased significantly by Piperine. Additionally, the anti-obesity effect of Piperine is related to the regulation of lipogenic and lipolytic genes, which is more effective in visceral fat than in subcutaneous fat\textsuperscript{25}.

A previous study by Nogaraa L, Naberb N, Pate E, et al., 2016 concluded that pepper extracts suppressed the increase in plasma lipids that might result from inhibition of pancreatic lipase activity that reduced fat and cholesterol absorption\textsuperscript{26}.

As shown in Table (2) there is a highly significant decrease in the serum level of oxidized LDL of the (HFD +BP) (HFD + Combination of BP&RP) group compared to the HFD group. Agbor et al. 2012 in their study had been reported that Piper species must prevent oxidation of LDL-c and enhance an antioxidant protection mechanism in the progression to prevent atherosclerosis in diet-fed hamsters. The phytochemicals, such as alkaloids, tannins, and polyphenols, which are the main constituents of Piper species, can be responsible for their therapeutic efficacy\textsuperscript{21}.

The present study showed a highly significant decrease in body weight of the animal groups compared to the HFD group, as shown in Table (5)

Due to the side effects of synthetic antioxidants and anti-obesity drugs, scientists are now focusing on natural products that produce similar effects to synthetic chemicals\textsuperscript{27}.

Possible mechanisms of the antiobesity effect of capsaicin by a reduction in body weight, induction of lipolysis in adipocytes, satiety, and reduction of energy and fat intake\textsuperscript{28}. In their study, Tremblay et al. 2018 reported that capsaicinoids accentuate the impact of caloric restriction on body weight loss; Capsaicin supplemented in diets attenuated increased hunger and decreased fullness, energy expenditure, and fat oxidation\textsuperscript{29}.

The previous study showed that capsaicin
in C. annuum contributed to thermogenesis, increasing energy expenditure\(^{29}\). When capsaicin application was tested on 3T3-L1 adipocytes, it decreased intracellular lipid content and was involved in thermogenesis\(^{30}\).

The possible explanation for the anti-obesity effect of Piperine is that the number of calories burned by muscle isolated from rabbits increased after adding Piperine to their diet\(^{31}\). The previous study explained another possible mechanism of the antiobesity effect of Piperine by preventing and slowing fat cell production\(^{32}\).

Enzymes, in general, are the most important suitable diagnostic tools for tissue damage, and their levels alter in the bloodstream under diseased conditions. Many reports showed that the effect of HFD on liver damage in rats could be diagnosed by elevated AST, ALT, and ALP activities in blood during liver damage\(^{33}\).

Sharma et al., 2012 showed a reduction in AST, ALT and ALP activities compared to the HFD control group, which could diminish the hepatotoxic effect caused by HFD-induced obesity\(^{28}\).

Preclinical studies showed a direct correlation between a high-fat diet and overweight and obesity among different animal groups\(^{34, 35}\). But these studies do not often discuss the relationship between overweight and obesity caused by a high-fat diet on kidney function, which is an important aspect of such studies. Ridwan R et al. 2019 observed the alteration in renal cellular morphology caused by a high-fat diet that induced overweight and obesity without any changes in blood levels of urea and creatinine\(^{32}\).

The previous study showed that red chili pepper did not cause significant changes in the functional biochemical parameters of the liver and kidney. However, excessive consumption of red pepper can induce liver and kidney damage that can be diagnosed by histological examination of both\(^{36}\).

In conclusion, the (HFD + RP) group and the (HFD + Combination) group showed the best hypolipidemic and antiobesity effects compared to the HFD group after 10 weeks of feeding. After observing the ultrastructure of liver tissue, lipid droplets were significantly reduced after treatment with pepper extracts, and the number of ER in the cytoplasm increased. These results suggested that the pepper extract exerted hypolipidemic effects and may have a protective effect against atherosclerosis, similar to a statin, the positive drug. Furthermore, after 10 weeks of gavage, the weight gain of the treated group was significantly lower than that of the HFD group, indicating that pepper extracts may have anti-obesity effects.

**Conflict of Interest**

There are no conflict of interest.

**REFERENCES**


32. Ridwan R, Nazir R, Rasyid H et al. Decreased Renal Function Induced by High-Fat Diet in Wistar Rat: The Role of Plasma Angiotensin-


