Apigenin ameliorates hypercholesterolemic-induced kidney injury via modulating renal KIM-1, Fn1, and Nrf2 signaling pathways

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Abstract. – OBJECTIVE: Hypercholesterolemia (HC) is a devastating metabolic disorder that has a negative effect on the kidneys’ functional and structural modalities via oxidative stress and inflammation. The aim of this paper is to elaborate on the role of the flavonoid apigenin (Apg), considering its antioxidant, anti-inflammatory, and antiapoptotic capabilities, in alleviating hypercholesterolemia-induced kidney injury.

MATERIALS AND METHODS: Twenty-four adult Wister male rats were allocated into four equal groups and treated for eight consecutive weeks: a control group, supplemented with a normal pellet diet (NPD); the Apb group, supplemented with NPD and Apg (50 mg/Kg); the HC group, fed with NPD enriched with 4% cholesterol and 2% sodium cholate; and the HC/Apg group, concomitantly rendered hypercholesterolemic and gavaged with Apg. At the end of the experiment, serum samples were collected to measure renal function parameters, lipid profile, MDA, and GPX-1. Thereafter, the kidneys were processed for histological study and homogenized to assess IL-1β, IL-10, and the gene expression of kidney injury molecule-1 (KIM-1), fibronectin 1 (Fn1), and NF-E2-related factor 2 (Nrf2) via RT-qPCR.

RESULTS: HC disturbed the renal function, lipid profile, and serum redox balance. In addition, HC elicited a proinflammatory/anti-inflammatory imbalance, upregulating KIM-1 and Fn1 and downregulating the Nrf2 gene expression of the kidney tissue. Moreover, HC induced marked histopathological changes in the kidney cytoarchitecture. Efficaciously, upon concomitant Apg supplementation with a high-cholesterol diet, most functional, histological, and biomolecular impairments of the kidney were comparatively restored in the HC/Apg group.

CONCLUSIONS: Apg mitigated HC-induced kidney injury via modulating the KIM-1, Fn1, and Nrf2 signaling pathways, a promising effect that may be useful as an adjunct to antihypercholesterolemic medications to treat the devastating renal complications of HC.

Key Words: Hypercholesterolemia, Renal injury, Apigenin, KIM-1, Fn1, Nrf2.

Introduction

Cholesterol is an essential compound for the structural integrity of the cell membranes and plays a vital role as a precursor for steroid hormones and bile acids. Elevated blood cholesterol level, known as HC, is a well-established risk factor for cardiovascular, cerebrovascular, and peripheral vascular diseases. HC can be defined as LDL cholesterol greater than 190 mg/dL or greater than 160 mg/dL with one major risk factor or greater than 130 mg/dL with two major risk factors. These major risk factors include hypertension, diabetes mellitus, smoking, age (male: 45 years or older; female: 55 years or older), a positive family history of premature atherosclerosis (younger than 55 years in a male and younger than 65 years in a female) and low HDL cholesterol levels (less than 40 mg/dL in male and less than 55 mg/dL in a female).

HC is mainly due to primary (genetic or familial) or secondary (acquired) causes. The classical genetic disorder is familial hypercholesterolemia due to mutations in the LDL receptor gene, resulting in LDL-C greater than 190 mg/dL in heterozygotes and greater than 450 mg/dL in homozygotes. These genetic mutations of the LDL receptor gene account for 85% of familial causes. Other genetic causes of familial hypercholesterolemia include defective apolipoprotein B, resulting in a loss of ligands binding to the LDL receptor. Additionally, mutations in the proprotein convertase subtilisin/kexin type 9 (PCSK9) gene lead to an increased affinity of PCSK9 for the LDL receptor, which results in a more rapid clearance of the LDL receptor by targeting it to the lysosome for degradation in the liver, thus resulting in HC.
adaptor protein, resulting in defective endocytosis of the LDL receptors. However, the most common cause is polygenic HC, which results from an interaction of unidentified genetic factors compounded by a sedentary lifestyle and an increased intake of high saturated fats, trans-fatty acids, and cholesterol\(^2\). The acquired causes include diabetes mellitus, cholestatic liver diseases, hypothyroidism, metabolic syndrome, obesity, and nephrotic syndrome. Moreover, some drugs, such as thiazide diuretics, cyclosporine, olanzapine, and smoking have been linked with an increased risk of hypercholesterolemia\(^2\).

In 2020, the World Health Organization (WHO) revealed the global prevalence of HC to be around 11.9\%, accounting for 28.5 million adults aged ≥20 years. In Saudi Arabia, the data on the prevalence of hypercholesterolemia are scarce, and the local prevalence ranges widely from 8.5\% to 54.9\%\(^6\).

It was hypothesized that, if HC is not promptly treated, many general deleterious effects can be caused, including atherosclerosis, coronary heart disease, myocardial infarction, ischemic cardiomyopathy, sudden cardiac death, ischemic stroke, microvascular disease, erectile dysfunction, and acute limb ischemia\(^6\).

Determinately, HC is highly prevalent in patients with chronic kidney disease (CKD), and the relation between both issues remains unclearly understood. Hypercholesterolemia is a major risk factor for CKD, owing to HC-induced renal atherosclerosis and OS. It was found that elevated LDL cholesterol is associated with increased systemic levels of oxidized LDL (OxLDL), which is well-known to be involved in endothelial cell dysfunction\(^10\). Moreover, several experimental studies\(^3\) have demonstrated that, in addition to its well-known proatherogenic effects on renal microvasculature, hypercholesterolemia may exacerbate the renal vascular responses to ischemia/reperfusion injury (I/R injury) with subsequent renal dysfunction.

In a crosstalk manner, CKD results in a profound dysregulation of several lipoprotein key enzymes that eventually contributes to increased LDL cholesterol, triglycerides, and triglyceride-rich apolipoprotein B (apoB), together with decreased HDL cholesterol\(^2\). With the progression of CKD, oxidative/nitrative stress participates in renal functional progression\(^13-15\). Oxidative/nitrative stress is an excess formation or insufficient removal of highly reactive molecules, such as reactive oxygen and/or nitrogen species (ROS and RNS), including superoxide, hydrogen peroxide, hydroxyl radical, and peroxynitrite\(^6\). Enzymatic sources for ROS formation include the mitochondrial respiratory chain, nicotinamide adenine dinucleotide phosphate (NADPH) oxidases, xanthine oxidase, cyclooxygenases, uncoupled nitric oxide synthase (NOS), and peroxidases, while antioxidative enzymatic systems include superoxide dismutase (SOD), catalase, glutathione peroxidase, gluthathione reductase, and heme oxygenase (HO)\(^7\).

To date, extensive research has been conducted to combat HC. However, no effective and clinically applicable preventive or curative compounds have been established. The major preventive emphasis strategy is directed primarily at exercise and changing lifestyles with strict diet regimens. In addition, several lipid-lowering drugs have been employed, such as statins (HMG-CoA reductase inhibitors), Ezetimibe (inhibits intestinal absorption of cholesterol), and PCSK9 monoclonal antibodies (increases LDL clearance from the circulation)\(^9\). Additionally, Gemcabene (enhances the clearance of VLDLs in plasma and the inhibition of the production of cholesterol and TGs in the liver through its liver-directed downregulation of hepatic apoC-III)\(^9\). Inclisiran increases LDL-C receptor recycling and expression on the hepatocyte cell surface, thereby increasing LDL-C uptake by hepatocytes and lowering LDL-C levels in the bloodstream\(^20\). Evinacumab inhibits angioptetin-like 3 (ANGPTL3), which is an inhibitor of lipoprotein and endothelial lipase that is responsible for increasing the levels of triglycerides and other lipids\(^21\). Vupanorsen reduces the production of the ANGPTL3 protein, which ultimately leads to a reduction in LDL-C levels\(^22\). ARO-ANG3 also inhibits ANGPTL3 and is designed to reduce triglycerides and decrease LDL-C in patients with mixed dyslipidemia\(^23\). Bempedoic acid inhibits ATP-citrate lyase, which is a component of the cholesterol synthesis pathway\(^24\).

However, the aforementioned synthetic drugs have several deleterious effects, and they are expensive and non-specific. One strategy that holds great promise in combating hypercholesterolemia is the use of flavonoid preparations derived from edible plants. These flavonoids are characterized by exhibiting high antioxidant activity, being well tolerated, cost-effective, of high therapeutic efficacy, and with almost no side effects.

In this study, attention was directed to Apigenin (Apg, one of the promising flavonoids). Apg (4′,5,7-trihydroxyflavone) is a natural product belonging to the flavone family and exists principally in vegetables (parsley, celery, and onions), fruits
Apigenin effect on kidney

(oranges), herbs (chamomile, thyme, oregano, and basil), and plant-based beverages. Apg is used as an alternative regimen with antioxidant, anti-apoptotic, and anti-inflammatory capabilities. Therefore, this study highlights and pays attention to the potentially mitigating role of Apg in the experimentally induced hypercholesterolemia of male rats.

Materials and Methods

Chemicals
All chemicals were purchased from Sigma-Aldrich Co. (St Louis, MO, USA).
- Cholesterol: White powder with purity >90% (CAS No: 57-88-5).
- Sodium cholate hydrate: (CAS No: 206986-87-0): Dried powder with purity ≥97.0%.
- Apigenin (Apg): Light yellow powder with purity >98% (CAS No: 520-46-5).
- Dimethyl Sulfoxide (DMSO): Derived in the form of a clear colorless liquid (CAS No: 76-86-5).
- Sodium pentobarbital: A vial of 2 ml solution (Virbac Animal Health, Milperra, NSW, Australia).

Study Setting
This study was conducted at Pharmacology Research Laboratory, Faculty of Pharmacy, King Abdulaziz University, Jeddah, Saudi Arabia, in April-June 2022.

Animals
Adult male Wister rats (n=24, aged approximately 12-16 weeks and weighing 190-220 g) were used in this study. The rats were obtained from the animal house of the Faculty of Pharmacy, King Abdulaziz University, Jeddah, Saudi Arabia. The rats were left to acclimatize without handling for one week prior to the start of the experiments. The animals were housed in well-ventilated cages at an ambient temperature of 23±2 °C, under the natural 12-hour day/night cycle with free access to a standard commercial rodent chow and tap water ad libitum throughout the entire study period.

Ethical Statement
The experimental protocol of this study was revised and approved by the local ethical guidelines established by the Research Ethics Committee, Faculty of Pharmacy, King Abdulaziz University, Jeddah, Saudi Arabia (Reference No: PH-1443-18). These institutional rules are in strict accordance with the international guiding principles for the care and use of laboratory animals.

Preparation of the Hypercholesterolemic Diet (HCD)

The rats were kept on two dietary regimens, a standard commercial normal pellet diet (NPD) (for the control group) (Ashirwad Industries, Chandigarh, India) and a hypercholesterolemic diet (HCD) (for the tested groups). The NPD consisted of 48.8% carbohydrates, 21% protein, and 3% fat and 27.2% others (calcium, phosphorus, fibers, and ash). On the other hand, the HCD was reconstituted by enriching the NPD with 4% (weight/weight) cholesterol and 2% (weight/weight) sodium cholate. All ingredients of the HCD were thoroughly mixed with water and transformed into pellets and then oven-baked at 65 °C overnight for proper drying to avoid fungal contamination. HCD was freshly prepared every 3-4 days and stored at -20°C.

Experimental Groups
The rats were randomly allocated into four groups of six rats each:

I (Control group): Rats were supplemented with a standard commercial normal pellet diet (NPD) of 48.8% carbohydrates, 21% protein, and 3% fat and 27.2% others (calcium, phosphorus, fibers, and ash) and were gavaged with 0.5 ml DMSO by oral gavage, once daily for eight weeks.

II (Apg group): This group was supplemented with NPD and gavaged with Apg (50mg/Kg) dissolved in 0.5 ml DMSO by oral gavage, once daily for eight weeks.

III (HC group): In this group, the rats were fed orally with NPD enriched with 4% (weight/weight) cholesterol and 2% (weight/weight) sodium cholate for eight weeks. The development of hypercholesterolemia was confirmed by measuring the serum cholesterol, and rats with cholesterol levels ≤ 250 mg/dl were included in the study.

IV (HC/Apg group): The rats of this group were rendered hypercholesterolemic as in group III for eight weeks; during this time, they were concomitantly gavaged with Apg at the same dose as group II.

Experimental Protocol

At the end of the experiment (after eight weeks), the rats were weighed and fasted overnight. The following day, blood samples were withdrawn from the retro-orbital venous sinuses. The blood samples were left to clot at room temperature in non-heparinized tubes. The collected blood sam-
samples were centrifuged at (3000 rpm. for 15 min) to obtain the serum, which was stored at −20 °C until the time of biochemical analysis. Afterward, the rats were weighed and anesthetized using sodium pentobarbital (60 mg/kg) and sacrificed by cervical dislocation. Then, the abdomen was opened by a longitudinal incision, and the left kidney was excised, cleaned from the surrounding fat and connective tissue, and fixed for 48 hours in 10% neutral buffered formalin (NBF) to be processed later for histological study28. The right kidney was excised and homogenized in ice-cold 0.01 M Tris–HCl buffer, pH 7.4, to obtain a 10% homogenate29.

**Biochemical Analysis**

**Measurement of the renal function**

Serum urea, creatinine, uric acid, sodium, and potassium were measured. Additionally, the animals were kept for 24 hours in metabolic cages (starting at 08 pm and ending at 08 am the next day), and 24-hour urine samples were collected to measure the level of urinary creatinine. Finally, creatinine clearance was calculated to measure the glomerular filtration rate (GFR).

**Measurement of the serum lipid profile**

Serum TC, TG, and HDL were analyzed using commercial kits (Nanjing Jiancheng Bioengineering Institute, Nanjing, China). Serum LDL was calculated according to the equation of30 [LDL = TC− (HDL + TGs/5)].

**Measurement of serum lipid peroxidation and antioxidant enzymes**

Malondialdehyde (MDA, an indicator of lipid peroxidation) was determined spectrophotometrically using a thiobarbituric acid reactive substances (TBARS) assay kit (Sigma-Aldrich, Chemical Comp., MO, USA). MDA was expressed as (nmol/ml). In addition, the serum antioxidant enzyme, Glutathione peroxidase-1 (GPX-1), was measured spectrophotometrically using a commercial assay kit (Abcam, UK). GPX-1 was expressed as (U/ml).

**Measurement of the renal tissue pro-inflammatory and anti-inflammatory cytokines**

In the kidney tissue homogenate, the pro-inflammatory cytokine (IL-1β) and the anti-inflammatory cytokine (IL-10) were measured using rat ELISA kits following the manufacturer’s instructions. IL-1β and IL-10 were expressed as pg/µg protein.

**Gene Expression Via Real Time-Quantitative Polymerase Chain Reaction (RT-QPCR)**

In the kidney tissue homogenate, the gene expressions of the kidney injury molecule-1 (KIM-1), fibronectin 1 (Fn1), and NF-E2-related factor 2 (Nrf2) were assessed by RT-qPCR. Briefly, the total RNA was extracted from the homogenate using a standard TRizol RNA isolation method (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA); then, cDNA was synthesized using a reverse transcription kit (Reverse Transcriptase Master Mix, Roche Diagnostics) according to the manufacturer’s instructions. RT-qPCR analysis was performed in a mixture containing 5 µL cDNA, 10 µL Eva green mix (Jena Bioscience), and 0.6 µL primers (10 µM), with a PCR grade water up to 20 µL. The amplification was carried out using Real-time PCR (Strata gene Mx 3005 P-qPCR System). The amplification conditions were 2 min at 50°C, 10 min at 95°C, and 40 cycles of denaturation for 15 s and annealing/extension at 60°C for 10 min. The relative gene expression of each target cDNA sample relative to the reference sample was calculated. The fold-change of each target gene expression was normalized to the Ct value of a reference gene of glyceraldehyde 3-phosphate dehydrogenase (GAPDH) using the comparative Ct method ($2^{ΔΔCT}$) method. The sequence of the primers of the target studied genes and that of GAPDH are shown in Table I.

**Histological Study**

The formalin-fixed kidneys were dehydrated in ascending grades of ethanol, cleared in xylene, and embedded in paraffin wax; then, serial sections of 5 µm thick were cut using a microtome (Leica RM 2125, Leica Biosystems Nussloch GmbH, Germany).

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**Table I.** Primer sequences of the studied genes for RT qPCR.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward</th>
<th>Reverse</th>
</tr>
</thead>
<tbody>
<tr>
<td>KIM-1</td>
<td>5'-TGGCAGTCGTGACATCTCAGA-3'</td>
<td>5'-GCAACGGGACATGCCAACATA-3'</td>
</tr>
<tr>
<td>Fn1</td>
<td>5'-CTCGTCCTTGGACCTCCACCA-3'</td>
<td>5'-TCTCCCTTCTGCTGCTGACT-3'</td>
</tr>
<tr>
<td>Nrf2</td>
<td>5'-CTTGCCGAGGACGCGCCATGACT-3'</td>
<td>5'-CTGAGCGAGGACATGGTGGTACT-3'</td>
</tr>
<tr>
<td>GAPDH</td>
<td>5'-TCAAGAAGGTGGTGAAGCAG-3'</td>
<td>5'-AGGTTGGAGAATGGGAGTGGT-3'</td>
</tr>
</tbody>
</table>

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and mounted on glass slides. Finally, some sections were stained with Hematoxylin and Eosin (H&E) stain “for a routine histological examination of the renal cortex”31. Finally, the stained H&E sections were examined under a light microscope.

Statistical Analysis

In the present study, the data were analyzed using the statistical package for social sciences program [SPSS v.23 (IBM, Armonk, NY, USA)] and the GraphPad prism (GraphPad 9 statistical Software, Inc., San Diego, CA, USA). The data are expressed as mean±SD. The variables were checked for normality using Shapiro-Wilk test. The significance of differences between groups was determined by one-way analysis of variance (ANOVA) followed by post-hoc Tukey’s test. p-value ≤0.05 was considered statistically significant.

Results

Effect of Apg on the Renal Function

Table II depicts a statistically significant increase in the mean serum urea, uric acid, and creatinine together with a significant decrease in the creatinine clearance of the HC group compared to that of the control and Apg groups. Upon concomitant Apg administration to the high-cholesterol-fed rats in the HC/Apg group, the deteriorated renal function parameters were returned nearly to a level comparable to that of the control and Apg groups.

Effect of Apg on the Lipid Profile

Table III depicts a statistically significant increase in the mean serum cholesterol, TG, and LDL, together with a significant decrease in serum HDL in the HC group compared to that of the control and Apg groups. In the HC/Apg group, these lipid profile parameters were mitigated compared to those of the HC group.

Effect of Apg on Serum Lipid Peroxidation and Antioxidant Enzymes

There was a statistically significant increase in the mean serum MDA and a decrease in serum GPX-1 in the HC group in comparison with the control and Apg groups. In the HC/Apg group, the mean serum MDA decreased, and serum GPX-1 increased significantly upon concomitant supplementation of Apg with HFD (Table IV).

Table II. Renal function tests in all animal groups (Mean±SD).

<table>
<thead>
<tr>
<th></th>
<th>Control group</th>
<th>Apg group</th>
<th>HC group</th>
<th>HC/Apg group</th>
<th>F-test</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Urea (mg/dl)</td>
<td>13.85±0.948</td>
<td>15.70±3.77</td>
<td>76.00±8.50</td>
<td>34.60±8.44</td>
<td>131.4</td>
<td>.000</td>
</tr>
<tr>
<td>BUN (mg/dl)</td>
<td>6.46±0.44</td>
<td>7.30±1.73</td>
<td>35.51±3.97</td>
<td>15.80±3.63</td>
<td>135.7</td>
<td>.000</td>
</tr>
<tr>
<td>Creatinine (mg/dl)</td>
<td>0.58±0.20</td>
<td>0.56±0.23</td>
<td>2.49±0.57</td>
<td>1.14±0.34</td>
<td>35.4</td>
<td>.000</td>
</tr>
<tr>
<td>Creatinine clearance (ml/min)</td>
<td>1.18±0.25</td>
<td>1.15±0.20</td>
<td>0.41±0.14</td>
<td>0.76±0.15</td>
<td>20.5</td>
<td>.000</td>
</tr>
</tbody>
</table>

Table III. Serum lipid profile in all animal groups (Mean±SD).

<table>
<thead>
<tr>
<th></th>
<th>Control group</th>
<th>Apg group</th>
<th>HC group</th>
<th>HC/Apg group</th>
<th>F-test</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>118.50±5.61</td>
<td>124.16±7.57</td>
<td>292.66±39.08</td>
<td>193.83±23.6</td>
<td>35.1</td>
<td>.000</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>75.66±3.9</td>
<td>81.83±6.21</td>
<td>160.33±31.06</td>
<td>84.50±11.72</td>
<td>33.2</td>
<td>.000</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>46.83±3.76</td>
<td>47.33±4.63</td>
<td>23.50±3.64</td>
<td>37.83±8.03</td>
<td>22.5</td>
<td>.000</td>
</tr>
<tr>
<td>LDL (mg/dl)</td>
<td>71.66±7.39</td>
<td>76.83±7.44</td>
<td>265.83±45.65</td>
<td>156.00±30.95</td>
<td>33.4</td>
<td>.000</td>
</tr>
</tbody>
</table>

Values are expressed as Mean±SD of 6 rats of each group. Statistical analysis was carried out using One-way ANOVA followed by Tukey’s test. Statistical significance was set at p-value<0.05. *Significant p-value when compared to the control group. p-value when compared to the HC group.
Effect of Apg on Renal Tissue Pro-Inflammatory and Anti-Inflammatory Cytokines

There was a statistically significant increase in the mean renal tissue IL-1β and a decrease in IL-10 in the HC group in comparison with the control and Apg groups. In the HC/Apg group, the mean renal tissue IL-1β decreased and serum IL-10 increased significantly upon concomitant supplementation of Apg with HFD (Table V).

Effect of Apg on the Gene Expression of KIM-1, Fn1, and Nrf2 in the Renal Tissue

The results of the qRT-PCR demonstrated that the gene expression of KIM-1 and Fn1 increased significantly by ~6-fold, and the expression of Nrf2 protein decreased by ~10-fold in the kidney tissue of the HC group compared with those of the control and Apg groups. The level of these three renal biomarkers was rescued upon Apg supplementation (Figures 1-3).

Table IV. Serum Lipid peroxidation and antioxidant enzymes in all animal groups (Mean±SD).

<table>
<thead>
<tr>
<th></th>
<th>Control group</th>
<th>Apg group</th>
<th>HC group</th>
<th>HC/Apg group</th>
<th>F-test</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDA (nmol/ml)</td>
<td>3.33±0.46</td>
<td>3.08±0.67</td>
<td>14.63±0.77*</td>
<td>70.78±1.13*</td>
<td>273.3</td>
<td>.000</td>
</tr>
<tr>
<td>GPX-1 (U/L)</td>
<td>176.33±10.76</td>
<td>181.33±12.48</td>
<td>86.50±15.57*</td>
<td>175.50±10.78*</td>
<td>79.4</td>
<td>.000</td>
</tr>
</tbody>
</table>

Values are expressed as Mean±SD of 6 rats of each group. Statistical analysis was carried out using One-way ANOVA followed by Tukey’s test. Statistical significance was set at p-value <0.05. *Significant p-value when compared to the control group. #Significant p-value when compared to the HC group.

Table V. Renal tissue pro-inflammatory and anti-inflammatory cytokines in all animal groups (Mean±SD).

<table>
<thead>
<tr>
<th></th>
<th>Control group</th>
<th>Apg group</th>
<th>HC group</th>
<th>HC/Apg group</th>
<th>F-test</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-1β (pg/µg protein)</td>
<td>5.96±1.02</td>
<td>5.26±1.58</td>
<td>14.60±0.68*</td>
<td>7.85±0.39*</td>
<td>104.0</td>
<td>.000</td>
</tr>
<tr>
<td>IL-10 (pg/mg protein)</td>
<td>5.43±1.08</td>
<td>6.23±1.59</td>
<td>2.28±0.84*</td>
<td>13.98±1.81*</td>
<td>76.8</td>
<td>.000</td>
</tr>
</tbody>
</table>

Values are expressed as Mean±SD of 6 rats of each group. Statistical analysis was carried out using One-way ANOVA followed by Tukey’s test. Statistical significance was set at p-value <0.05. *Significant p-value when compared to the control group. #Significant p-value when compared to the HC group.

Figure 1. Gene expression of KIM-1 in the renal tissue in different animal groups (Mean±SD). Values are expressed as Mean±SD of 6 rats of each group. Statistical analysis was carried out using One-way ANOVA followed by Tukey’s test. Statistical significance was set at p-value <0.05. *Significant p-value when compared to the control group. #Significant p-value when compared to the HC group.

Figure 2. Gene expression of Fn1 in renal tissue in different animal groups (Mean±SD). Values are expressed as Mean±SD of 6 rats of each group. Statistical analysis was carried out using One-way ANOVA followed by Tukey’s test. Statistical significance was set at p-value <0.05. *Significant p-value when compared to the control group. #Significant p-value when compared to the HC group.
Apigenin effect on kidney

**Effect of Apg on the Histology of the Kidney Tissue**

The H&E-stained section of the kidney of a control rat (group I) and Apg rat (group II) revealed a normal cytoarchitecture of both cortical and juxta-medullary (JM) nephrons, having renal corpuscles encompassing normal Bowman’s capsule and glomeruli with well-organized juxtaglomerular apparatus. Additionally, both normal proximal convoluted tubules (PT) and distal convoluted tubules (DT) and renal microvasculature were observed (Figures 4-5).

On the other hand, the H&E-stained section in the kidney of the HC rat (group III) displayed subcapsular congestion and hemorrhage of the renal capsules. Both cortical and JM nephrons exhibited wide Bowman’s spaces, dilated congested glomerular capillaries, and a slight expansion of the mesangial matrix. Moreover, dilated congested microvasculature with focal areas of hemorrhage in the renal interstitium was observed. Additionally, disorganized juxtaglomerular apparatus and dilated congested glomerular afferent and efferent arterioles. The cells of PT and DT were laden with many lipid vacuoles and appeared more acidophilic. The DT showed a dilated lumen with some cells having pyknotic nuclei, and other cells appeared shortened than the control rats (Figure 6).

The H&E-stained section in the kidney of HC/Apg rat (group IV) showed apparently normal cytoarchitecture of both cortical and JM nephrons with an intact visceral layer of Bowman’s capsule and the glomerular basement membrane. Moreover, the renal microvasculature revealed mild congestion and minute areas of hemorrhage in the renal interstitium (Figure 7).

![Graph showing gene expression of Nrf2 in renal tissue in different animal groups](image)

**Figure 3.** Gene expression of Nrf2 in renal tissue in different animal groups (Mean±SD). Values are expressed as Mean±SD of 6 rats of each group. Statistical analysis was carried out using One-way ANOVA followed by Tukey’s test. Statistical significance was set at \( p\)-value<0.05. *Significant \( p\)-value when compared to the control group. #Significant \( p\)-value when compared to the HC group.

![Photomicrographs of H&E-stained sections](image)

**Figure 4.** A photomicrograph of the H&E-stained section of the kidney of a control rat. a, Normal appearance of the renal capsule (Cap) and cortical nephrons reveal conventional renal corpuscles and renal tubules. The cortical renal corpuscle (C) exhibits Bowman’s capsule (BC) with average Bowman’s space (BS) and the glomerulus (G), which is in intimate contact with the visceral layer of BC (white arrow). At the vascular pole of the glomerulus, there is a well-organized juxtaglomerular apparatus (interrupted arrow). The cortical renal tubules reveal proximal convoluted tubules (PT) lined by acidophilic tall cuboidal epithelium with a narrow lumen, and distal convoluted tubules (DT) lined by smaller cuboidal epithelium with a wide lumen. b, Normal juxtamedullary nephrons with a normal aspect of renal corpuscles and renal tubules. The corpuscle (C) has the same histological features as the cortical corpuscle, with a distinct visceral layer of Bowman’s capsule (white arrow). Also, the proximal convoluted tubules (PT) and distal convoluted tubules (DT) appear normal, besides the common arcuate artery (AA) and arcuate vein (AV). Scale bar =50 μm. H&E x400.
**Figure 5.** A photomicrograph of the H&E-stained section of the kidney of an Apg-treated rat. 

- **a.** The renal capsule (Cap) and cortical nephrons appear normal. The cortical renal corpuscle (C) exhibits common Bowman’s capsule (BC) with average Bowman’s space (BS) and the glomerulus (G), which is in intimate contact with the visceral layer of BC (white arrow). The cortical renal tubules reveal regular proximal convoluted tubules (PT) and distal convoluted tubules (DT).
- **b.** The juxtamedullary nephron exhibits normal renal corpuscle (C) having the same histological appearance as the cortical nephrons with a definite visceral layer of Bowman’s capsule (white arrow). Also, PT and DT appear normal, besides the arcuate vein (AV) at the cortico-medullary junction. Scale bar = 50µm. H&E x400.

**Figure 6.** A photomicrograph of the H&E-stained section in the kidney of an HC-treated rat. 

- **a.** A renal capsule (Cap) with subcapsular hemorrhage and congestion (double arrows) are noticed. Also, interstitial vascular congestion with focal areas of hemorrhage (long arrows) is inspected. The cortical nephrons reveal nephropathy, whereas the cortical renal corpuscles (C*) display wide Bowman’s space (BS*), dilated congested glomerular capillaries (arrowheads), and a slight expansion of mesangial matrix (short arrows). The cells of some proximal convoluted tubules (PT*) are laden with large lipid vacuoles. The cells of other proximal convoluted tubules (PT***) appear more acidophilic. The distal convoluted tubules (DT*) reveal dilated lumen with some cells having pyknotic nuclei and other cells appear shortened than that of the control rats.
- **b.** The juxtamedullary nephrons exhibit affected renal corpuscles (C*) that reveal dilated congested glomerular capillaries (arrowheads), and a slight expansion of mesangial matrix (short arrow). Also, disorganized juxtaglomerular apparatus (interrupted arrow). The cells of the distal convoluted tubules (DT*) are loaded with lipid vacuoles alongside dilated congested glomerular arteriole (GA). Some cells of the proximal convoluted tubules appear more acidophilic (PT***). Moreover, dilatation and congestion of the arcuate artery (AA*) with leukocytic infiltration (curved arrow) besides, focal areas of hemorrhage (long arrows) are noticed in the interstitium. Scale bar = 50µm. H&E x400.
Discussion

Hypercholesterolemia (HC) is a metabolic disorder that increases the risk of renal diseases. The renal burden depends on the degree and duration of exposure to raised cholesterol levels. In this study, the rats supplemented with a high-cholesterol diet (HC group) had oxidative stress (OS) that was intermingled with disturbed biomarkers of oxidant/antioxidant balance, whereby the serum MDA, a marker of lipid peroxidation product, increased, while the serum GPX-1 activity decreased in the HC rats. It was reported that HC caused the oxidative modification of LDL, protein glycation, and glucose-autodoxation, thus leading to an excess production of lipid peroxidation products, particularly MDA, which may cause an elevation of OS in hyperlipidemic subjects. Equivalently, it was found that MDA increased in the renal tissue of hypercholesterolemic rats. Moreover, Green et al. have found that diet-induced hypercholesterolemia increased ROS levels, contributing secondarily to a decrease in the peroxidase activity of the enzymatic antioxidant glutathione in rats.

Previous studies have shown that HC is an independent risk factor for the onset and progression of renal disease via the induction of OS in the kidneys. This effect of the HC diet added a pro-oxidant burden to the naturally occurring one in the kidneys due to its intertwined renal vasculature that often causes renal ischemia-reperfusion injury (IRI) with the subsequent production of overwhelming ROS, which are common mediators underlying various kidney diseases. In line with this, patients with chronic kidney disease (CKD) often experience an imbalance between oxygen reactive species (ROS) production and antioxidant defenses, leading to cell and tissue damage. Earlier studies have shown a positive correlation between the imbalance in pro-oxidant and antioxidant activities and renal dysfunction in humans. Strikingly, OS has been found even in the early stages of CKD, and it seems to increase as CKD progresses and correlates significantly and inversely with the level of glomerular filtration rate (GFR) as reflected by an increased systemic marker of OS plasma 8-isoprostane levels. Additionally, Witko-Sarsat et al. found the same results using other markers of OS, including MDA and GPX.

Therefore, the modulation of HC-induced renal OS is a feasible approach to prevent and/or treat deleterious renal consequences. This mod-

Figure 7. A photomicrograph of the H&E- stained section in the kidney of HC/Apg-treated rat. a, The renal capsule (Cap) appears normal without vascular congestion, and the cortical nephrons display an improvement, whereas the renal corpuscles (C*) exhibit slight congestion (arrowhead) in the glomerular capillaries and a minimal proliferation of the mesangial matrix (short arrow). Moreover, some renal tubules (T*) are dilated while the majority of the tubules, proximal (PT) and distal convoluted tubules (DT), are conserved without deposition of fat vacuoles. b, The juxtamedullary nephrons illustrate some preserved renal corpuscles (C) with a slightly defined visceral layer of Bowman’s capsule with the glomerulus (white arrow), while few renal corpuscles (C*) exhibit slight congestion (arrowhead). However, most of the renal tubules, proximal (PT) and distal convoluted tubules (DT), are preserved. The arcuate artery (AA) and vein (AV) appear normal without congestion. Scale bar =50µm. H&E x400.
ulation can be approached in at least three different ways. First, cholesterol-lowering therapies have indirect secondary antioxidant properties. Second, the induction of endogenous enzymatic antioxidant systems or the inhibition of the pro-oxidant enzymes may be feasible. The third novel approach is the application of antioxidant molecules, particularly those of natural products such as flavonoids.

In this regard, the third approach was chosen in this study, where the concomitant administration of the flavonoid Apigenin (Apg) alongside the high-cholesterol diet could successfully mitigate the oxidant/antioxidant imbalance and restore the MDA and GPX-1 to approximately their basal control levels. Many studies have reported the antioxidant activity of Apg. Recently, an in vitro study and another in vivo study have documented an efficacious antioxidant activity of Apg against renal tubular epithelial cell injury.

In this study, the obtained HC-induced redox imbalance may be beyond the histological changes that were clearly contemplated under the light microscopic examination of the kidney tissue. Obviously, microscopic changes involving both glomeruli and renal tubules of the cortical and juxtamedullary nephrons were observed and pointed mostly to the development of glomerulosclerosis and segmental necrosis in the renal tissues of the HC group. Moreover, dilatation in glomerular capillaries and Bowman’s space were detected; in addition, many necrotic interstitial cells in the kidneys of the HC rats were observed. The aforementioned renal histopathological changes were partially ameliorated in HC/Apg group.

Redox imbalance and inflammation of the renal tissue was encountered in HC rats where a significant concomitant increase in pro-inflammatory markers (IL-1β) and a decrease in the anti-inflammatory markers (IL-10) in the renal tissue homogenate of the HC group was detected compared with those of the control one. Based on the histological evidence, the inflammation of the renal tissue in the HC group was evident and confirmed under a light microscope, where mononuclear cell infiltration in the renal cortices was contemplated. Inflammatory cells, which are locally activated in the renal microenvironment or migrate from hematopoietic organs, further produce ROS. Thus, the crosstalk between OS and inflammation represents a key aggravating factor for the initiation and progression of acute kidney diseases, including diabetic nephropathy, hypertension-associated kidney disease, and toxic-induced nephropathy.

Pathologically, HC represents an inconvenience mechanism that drives the associated renal injury. The high-cholesterol diet frequently known as the western diet (WD), which is characterized by highly processed foods with high fat content, is one of the major contributors to the increased incidence of CKD, which is disrupted lipid profile, kidney function, and structural changes. In this respect, disrupted lipid profile, kidney function, and structural changes were employed in diet-induced hypercholesterolemia in both rat models and mice models. Population-based studies have determined that high cholesterol is a strong independent risk factor for end-stage renal disease (ESRD), even after adjusting for other major risk factors associated with this condition.

Comparatively, the HC/Apg group of rats exhibited a significant drop in IL-1β and a rise in IL-10 in the renal tissue homogenate, a distinctive finding that implies the anti-inflammatory capability of Apg. In previous studies, Apigenin has shown several valuable bioactive properties, such as antioxidant and anti-inflammatory activities. Apigenin has been repeatedly reported to attenuate the nephrotoxicity caused by several toxic agents via its antioxidant and anti-inflammatory attributes.

In this work, in conjunction with the renal tissue inflammation, a derangement of the lipid profile was encountered in HC rats where there was a significant increase in serum cholesterol and triglyceride, and LDL, together with a decrease in HDL. This HC-associated dyslipidemia worsened the renal function biomarkers with a rise in serum urea, BUN, and creatinine aligned with decreased GFR as employed by decreased creatinine clearance. These findings might point to the possibility of the development of CKD following a chronic high-cholesterol diet.

As part of a vicious circle, all patients with CKD experience a secondary form of dyslipidemia characterized by increased serum triglycerides with elevated VLDL, small dense LDL particles, and low HDL cholesterol. All these particles are characterized by triglyceride-rich apolipoprotein B (apoB)-containing complex lipoproteins, which have a significant atherogenic potential. CKD results in the profound dysregulation of several key enzymes and metabolic pathways that eventually contributes to dyslipidemia. With the progression of CKD, these metabolic derangements may be further worsened and participate in atherogenic diathesis and possibly further progression of renal dysfunction.

Previous animal studies have shown a positive correlation between the presence of an atherogenic lipid profile and the onset of glomerulosclerosis and endothelial dysfunction. In this respect, disrupted lipid profile, kidney function, and structure were employed in diet-induced hypercholesterolemia in both rat models and mice models.
Apigenin effect on kidney

Consistent with the experimental models, dyslipidemia in humans is usually associated with the development and progression of renal dysfunction. Among human studies relating dyslipidemia to the renal outcome, one study found that a higher total cholesterol and a lower HDL cholesterol were significantly associated with an increased risk of developing renal dysfunction.

In this study, Apig significantly restored the distorted lipid profile upon its concomitant supplementation with 4% cholesterol and 2% sodium cholate. Inconsistent with these results, Apig regulated cholesterol metabolism in vivo and antilipidemic effect. Moreover, Apig improved lipid metabolism in vitro. These collective results shifted the attention gradually in recent years to Apig as a novel antilipidemic natural plant flavonoid with a high efficacy and few adverse effects rather than the available antilipidemic drugs.

Profound dyslipidemia in the HC group could explain the accumulation of fat droplets in the renal tubules upon the light microscopic examination of the kidney tissue. This finding was in line with a previous work in which HC caused the accumulation of fat droplets in the tubular cells of rats. The accumulation of fat droplets in tubular cells initiated lipotoxicity that determinately affects the cell metabolism with the overproduction of OS and increased cell apoptosis. Szeto et al. have shown that excess fat droplets in the kidney lead to mitochondrial damage in all renal cell types, with 50% cell loss, reduced size, and loss of cristae membranes that contain the protein complexes of the electron transport chain; thus, a loss of ATP production was employed. Another drawback of excess fat droplets in the kidney is an increased inflammatory cell infiltration in the histological analysis of the kidney tissue. Indistinguishably, this drawback was strikingly achieved in this study.

The disclosed HC-induced lipid accumulation in the kidney tissue was associated with an increased expression of the immune-related biomarkers genes, viz., kidney injury molecule-1 (KIM-1) and fibronectin 1 (Fn1), in the renal tissue of the HC group. Biologically, KIM-1 is a type 1 transmembrane protein that serves as a urinary marker in acute renal tubular injury. In a human study, high urinary KIM-1 levels and an extensive expression of this protein were detected in biopsies from the proximal tubule of patients with acute tubular necrosis. In a somewhat analogous manner to KIM-1, fibronectin 1 (Fn1) is an extracellular matrix protein that is linked to some renal diseases characterized by proteinuria, hematuria, and considerable fibronectin deposits in the mesangium and subendothelial space, subsequently leading to end-stage renal failure. Upregulated Fn1 gene has been identified in several human diseases, including renal carcinoma. In previous studies, Fn1 mRNA expression was significantly higher in renal cell carcinoma (RCC) compared to that in normal renal tissue, indicating its possible role in renal carcinogenesis and its progression. Recently, it has been intriguing to speculate that the ubiquitous expression of Fn1 is an unfortunate common histological endpoint of progressive CKD.

Another critical gene that was unfortunately downregulated in the kidney tissue of the HC group was the transcription factor Nrf2. This factor normally serves to protect the cells from harmful oxidative stress, playing an integrative role in inducing the expression of genes encoding enzymes involved in antioxidant (e.g., glutathione and NADPH) production and the reduction in pro-oxidants (e.g., heme and quinonoids). It was determined that the loss of Nrf2 aggravates tissue damage and fibrosis in kidney disease models, such as autoimmune nephritis, diabetic nephropathy, toxic injury, and IRI. Additionally, clinical trials investigating Nrf2-activating compounds in kidney disease patients are ongoing, and beneficial effects are being obtained. Successfully, Apig modulated the HC-induced KIM-1, Fn1, and Nrf2 gene expression in the kidney tissue.

**Limitations**

We acknowledge that this study has some limitations. Kidney function in rats could not be assessed via monitoring the passage of an extracellular contrast agent in conjunction with MRI, which is a gold standard method that could enable us to better understand the relative contributions of HC to the detrimental renal functional outcomes and subtle structural abnormalities, as it is not available in our lab.

**Conclusions**

The results reveal that Apig ameliorated HC-induced renal injury by suppressing not only oxidative stress and inflammation, but also it has a beneficial role in modulating the HC-induced KIM-1, Fn1, and Nrf2 gene expression. Therefore, Apig is expected to emerge as a first-in-class innovative natural flavonoid strategy for preventing HC-induced kidney injury and its future transfer to clinical applications must be a paramount concern.
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Conflicts of Interests
The author declares no competing interests.

Availability of Data and Materials
The data supporting this study’s findings are available from the author.

Ethics Approval
The experimental protocol of this study was revised and approved by the local ethical guidelines established by the Research Ethics Committee, Faculty of Pharmacy, King Abdulaziz University, Jeddah, Saudi Arabia (Reference No; PH-1443-18). These institutional rules are in strict accordance with the international guiding principles for the care and use of laboratory animals.

References
Apigenin effect on kidney


