Curcumin improves liver damage in male mice exposed to nicotine

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1. Introduction

A large and increasing number of patients in the world use medicinal plants and herbs for health purposes. Curcuma longa (turmeric; 薑 黃), one of the oldest plants, belongs to the Zingiberaceae family, which has long been used in traditional medicine for blood purification, digestion, arthritis treatment, liver protection, and as an anti-inflammatory agent. The color of turmeric is because of a substance called curcumin, which comprises 3–4% of turmeric. Numerous studies have reported the antioxidant properties, anti-mutation and antitumor effects, and carcinogenic characteristics of curcumin. Curcumin affects the metabolism of arachidonic acid by inhibiting the phosphorylation of phospholipase A2 (PLA2), decreasing the expression of cyclooxygenase (COX)-2 gene and inhibiting the catalytic activity of COX-5. These effects induce the anti-inflammatory activity of curcumin. In addition, curcumin decreases the expression of different inflammatory cytokines such as interleukin (IL)-1, tumor necrosis factor (TNF)-α, IL-6, and chemokines. Curcumin shows efficacy in promoting wound healing as well as treating liver ailments, urinary tract diseases, and hepatitis. Pharmacologically, curcumin exhibits a wide range of effects including anti-inflammatory, hypo-cholesterolemic, and anti-infection activities and as well as anticarcinogenic effects. Although approximately 4000 components are present in the cigarette, nicotine is a highly toxic organic compound containing nitrogen and alkaloid, which are mostly found in tobacco, and it is responsible for some of the deleterious effects of smoking. Nicotine affects a variety of cellular processes including altered gene expression. Exposure to nicotine produces oxidative stress, and it is thought that nicotine-induced oxidative stress plays a key role in the development of nicotine-related diseases.
tissue injuries in the mouse, often resulting in a depletion of glutathione content and a decrease in the activity of some oxygen free radical scavengers, such as catalase and superoxide dismutase. The liver is considered to be the major site of nicotine biotransformation, and nicotine exerts a number of adverse physiological effects on the liver. Nicotine is absorbed through the lungs during smoking and is rapidly metabolized in the liver, which induces three major adverse effects on the liver: toxic (direct or indirect), immunological, and oncogenic. Smoking causes liver cell injury and exerts genotoxic effects on rat liver. Current studies on curcumin effects have not reported on the protective effect of curcumin against nicotine; therefore, the current study was conducted to analyze the protective effect of curcumin to offset the damage induced by nicotine in the liver of male mice.

2. Materials and methods

2.1. Chemicals

Curcumin (4-hydroxy-3-methoxyphenyl)-1,6-heptadiene-3,5-dione; C21H20O6) powder (Merck, Germany) was purchased. The powder was dissolved in absolute ethanol (C2H5OH) and diluted with normal saline (0.9%) to prepare different doses. Also, the nicotine solution (1-metyl-2-3-pyridel-pirolidin; C10H14N2, Merck, Germany) was also purchased and was diluted by normal saline (0.9%) for administration (Fig. 1).

2.2. Experimental protocol

The mice were randomly divided into eight groups of six: control group (normal saline; 1 mL distilled water/daily); nicotine-treated group (2.5 mg/kg); 3) nicotine + curcumin 10 mg/kg treated group; nicotine + curcumin 30 mg/kg treated group; nicotine + curcumin 60 mg/kg treated group; nicotine + curcumin 10 mg/kg treated group; curcumin 30 mg/kg treated group; and curcumin 60 mg/kg treated group. Nicotine was administered intraperitoneally once a day for 4 weeks. Curcumin and nicotine plus curcumin were administered intraperitoneally in animals.

2.3. Animals

Forty-eight Balb/c male mice with weight range of 27–30 g were purchased from Tehran Razi Institute. Animals were kept at the temperature of 22 ± 2°C, under controlled environmental conditions, 12/12 hours light/dark cycle and free access to water and food. Maintenance and care of experimental animals complies with National Institutes of Health guidelines. Experiments were designed to conform with the International Guiding Principles for Biomedical Research Involving Animals (1985).

2.4. Liver weight and collection of blood serum

At the end of the experimental period, all animals were deeply anesthetized with ether. Blood was collected from the right ventricle, and serum separated and stored at −80°C for measurement of nitric oxide. They were then killed. Livers were removed and weighed on a microbalance sensitive to 0.001 mg (Precisa 125A, Switzerland) and recorded.

2.5. Histological analysis

For the histological evaluation of the hepatic structures, the lower 1-cm-long part of the right lobe of the liver in transverse pieces was removed, washed in saline, and fixed in 10% formalin at room temperature for 72 hours. After tissue fixation, it was thoroughly washed under running water and dehydrated in ascending concentration ethanol, cleared in xylene, and then embedded in soft paraffin. Thin sections (5 μm) were cut using a microtome (Leica RM 2125, Leica Microsystems Nussloch GmbH, Germany) and stained with hematoxylin and eosin. The preparation was examined with an Olympus BX-51T-32E01 research microscope connected to a DP12 Camera with 3.34-million pixel resolution and Olysys Bio software (Olympus Optical Co. LTD, Tokyo, Japan).

2.6. Morphometric measurements

For each hepatocyte, the total cellular area was measured. The outline of each hepatocyte was measured after taking an image with a 40 × objective lens. The longest and shortest axis were measured in the drawing of each hepatocyte in order to estimate the mean diameter (mean axis). At least 50 hepatocytes from each zone (total, 100) were measured in each liver. A separate measurement for central hepatic vein was performed, using the same methodology.

2.7. Griess assay

Nitric oxide was measured based on Griess colorimetric assay. Accordingly, 1-N-1-naphthylenediamine (NEED), sulfonamide solutions, and nitrite standards were prepared. To measure nitrite concentration in serum, after defreezing the serum samples, 100 μL of the sample serum was deproteinized by zinc sulfate and transferred to the wells. One hundred microliters of chloride vanadium, 50 μL sulfonamide, and 50 μL NEED solutions were added afterward. The samples were incubated in the temperature of 30°C in darkness. The optical density of samples was measured using an enzyme-linked immunosorbent assay reader (Hyperion, Germany) at the wavelength of 540 nm.

2.8. Biochemical analysis

The liver was minced and homogenized (10% w/v) in ice-cold 0.1M sodium phosphate buffer (pH 7.4). The homogenate was centrifuged twice at 10,000 rpm for 15 to 20 min at 4°C to obtain enzyme fraction. The resultant supernatant was used for various biochemical assays. Activities of aspartate aminotransferase (AST) and alanine aminotransferase (ALT) were assayed using the method of Reitman and Frankel. Activities of alkaline phosphatase (ALP) were determined according to the protocol described in the laboratory practical manual.

2.9. Statistical analysis

All the quantitative data were presented as mean ± standard deviation. One-way analysis of variance followed by least significant difference post hoc test were performed to determine the statistical significance between different groups using SPSS software package 16.0 (Statistical Package for the Social Sciences, version 16.0, SPSS Inc, Chicago, Illinois, USA). A p value <0.05 was considered significant.

Fig. 1. Structure of nicotine and curcumin.
Fig. 2. *Significant decrease of liver weight in the nicotine group compared with the saline group (p < 0.05). **Significant increase in all groups of curcumin compared with the nicotine group (p < 0.05). ***Significant increase all groups of curcumin-nicotine compared with the nicotine group (p < 0.05).

Fig. 3. (E) *Significant increase of the mean hepatocytes diameter in the nicotine group compared with the saline group (p < 0.05). **Significant decrease in all groups of curcumin administration compared with the nicotine group (p < 0.05). ***Significant decrease in all groups of curcumin-nicotine administration compared with the nicotine group (p < 0.05).

(A) Saline group. (B) Nicotine group. (C) Curcumin 60 mg/kg. (D) Curcumin (60 mg/kg) plus nicotine (magnification 100×).
3. Results

3.1. Liver weight

In the current study, the effective dose of nicotine (2.5 mg/kg) caused a significant decrease in the liver weight of the mice compared with the saline group \( (p < 0.05) \). Moreover, liver weight was significantly decreased in animals treated with curcumin and curcumin plus nicotine in all doses in comparison with the nicotine group \( (p < 0.05) \) (Fig. 2).

3.2. Morphometric measurements

The mean diameter of hepatocytes and central hepatic vein was substantially increased in the nicotine administration group in comparison with the saline group \( (p < 0.05) \). Further, curcumin and curcumin plus nicotine caused a significant decrease in the mean diameter of hepatocytes and central hepatic vein in all treated groups in comparison with nicotine group administration \( (p < 0.05) \) (Figs. 3 and 4).

3.3. Nitric oxide

The mean nitric oxide in blood serum increased significantly \( (2.5 \text{ mL/kg}) \) in the nicotine group in comparison with the saline group \( (p < 0.05) \). Also, the mean nitric oxide in blood serum decreased significantly in curcumin and curcumin plus nicotine in all groups in comparison with the nicotine group (Fig. 5).

3.4. Biochemical analysis

Nicotine \( (2.5 \text{ mL/kg}) \) caused a significant increase in the mean of ALT, AST, and ALP enzymes compared with the saline group \( (p < 0.05) \). In addition, the mean of ALT, AST, and ALP enzymes

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Fig. 4. (E) *Significant increase of the mean central hepatic vein diameter in the nicotine group compared with the saline group \( (p < 0.05) \). **Significant decrease in all groups of curcumin administration compared with the nicotine group \( (p < 0.05) \). ***Significant decrease in all groups of curcumin-nicotine administration compared with the nicotine group \( (p < 0.05) \). (A) Saline group. (B) Nicotine group. (C) Curcumin 60 mg/kg. (D) Curcumin (60 mg/kg) plus nicotine (magnification 100×).
decreased significantly in curcumin and curcumin plus nicotine in all groups in comparison with the nicotine group \((p < 0.05)\) (Figs. 6, 7, and 8).

Liver damage induced by nicotine administration included histological changes of liver confirmed by hematoxylin and eosin staining (Fig. 9).

4. Discussion

The liver is an organ that plays a significant role in the oxidation of fatty acids and carbohydrates.\(^1^3\) Curcumin, with its high antioxidant power, exerts its protective effect on the liver.\(^2^7\) Moreover, curcumin can have protective effects on lipid peroxidation in rat liver in oxidative stress conditions, can collect free radicals, and can increase intracellular glutathione levels.\(^2^7\) The current study investigated the protective effects of curcumin against the disorders induced by nicotine administration, which affected liver weight, diameter of hepatocytes and central vein, serum level of liver enzymes (AST, ALT, and ALP), and nitric oxide secretion level. In the current study, nicotine administration significantly increased the serum level of AST, ALT, and ALK enzymes in comparison with the control group, which mostly approached the normal level with curcumin administration. Necrosis or cell membrane damage can cause the release of these enzymes into blood. However, the serum level of these enzymes is associated with liver performance.\(^2^8\) It seems that nicotine causes the destruction of hepatic cell membrane, thereby releasing cytosomal enzymes of hepatocytes and finally increasing their serum level.\(^2^9\) In the groups that received curcumin, the amount of these enzymes was reduced, which indicates the protective effects of curcumin against the toxicity induced by nicotine. The obtained results may be due to the antioxidant effects and reduction of oxidative stress in curcumin.\(^3^0\) Further, because TNF-\(\alpha\) and IL-1 play a role in the induction of hepatic necrosis, curcumin can reduce the effects of toxicity by inhibiting the secretion of TNF-\(\alpha\) and IL-1 by macrophages.\(^3^1\) The findings of the current study were in line with the results of Fu et al,
which reported curcumin as a factor to reduce the increased hepatic enzyme due to Chemokine (C–C motif) ligand 4 (CCl₄) administration. The results of liver weight analysis showed a significant decrease in liver weight between the nicotine and control (saline) groups; however, liver weight increased after curcumin administration in comparison with the nicotine group. The metabolic disorders in rats caused by nicotine administration might cause the weight loss, which was largely compensated by curcumin administration

The results of the current study were also in agreement with the findings of Chuang et al, who reported curcumin as the factor to increase liver weight. Moreover, the results confirmed the findings of the study carried out by Willis et al, in which nicotine was reported to reduce the weight of different organs, including the liver. The histopathological results obtained in this study revealed the increase in diameter of mean hepatocytes and central veins due to nicotine administration. The change in the size of hepatocytes and central veins seems to be the result of an increase in the metabolic activity of cells to excrete toxins from the body during the toxification process. It seems that free radicals induced by nicotine metabolism trigger lipid peroxidation and reaction to DNA and membrane proteins, thereby causing cell damage. Antioxidant compounds such as curcumin can induce inhibitory effects on cytochrome P450, prevent nicotine metabolism, and consequently reduce the production of free radicals. Tetrahydro curcumin is one of the most important metabolites of curcumin and an active biological component of turmeric (薑薑 jiang huang) in the cytosol of hepatic cells. This substance seems to have extensive antioxidant activities in both in vitro and in vivo conditions. The findings of this study were also in line with the results of the study by Suñer et al, in which they reported nicotine administration resulted in increased size and vascularity of choroidal neovascularization. The results obtained from the measurement of nitric oxide in blood indicated a significant increase between the nicotine and control (saline) groups; however, nicotine administration reduced nitric oxide level. Nitric oxide is a free radical that is produced in mammalian cells and interferes with
regulation of physiological processes, and its production increase is associated with induction of various diseases. The hydroxyl radicals produced by nitric oxide seem to interfere with the genesis process and hepatic toxicity. Nicotine can induce the release of noradrenalin in paraventricular nucleus and amygdala through direct effect on the nuclei of solitary tract and N-methyl-D-aspartate receptors, which in turn stimulate the production of nitric oxide and noradrenergic activity of neurons. Nicotine absorption in the body seems to be followed by the increase of serum nitric oxide level and oxidative stress. Curcumin is structurally a polyphenol and exerts a protective effect on lipid oxidation in the liver in oxidative stress conditions, and acts as a collector of free radicals of oxygen and enhancer of intracellular glutathione. Thus, it can destroy the nitric oxide system (protein enzymes, substrates, and cofactors) and reduce nitric oxide production. The results of the study conducted by Zhao et al indicated that nicotine can decrease angiogenesis in gastric mucosa through inhibition of nitric oxide production and prevention of new cells’ production process, which are in contrast with the findings obtained in the current study. However, the results of the current study confirm the findings of Muller and Schepers, showing the increase of peroxynitrite formation due to cigarette smoke, an element known to damage DNA.

5. Conclusion

The current investigation demonstrates that nicotine has the potential to cause substantial damage in liver tissues and curcumin may improve some of this liver damage in mice. Therefore, curcumin could be useful to protect the liver from nicotine toxicity. The antioxidant effects of curcumin may be a major reason for its positive effect on liver parameters. However, further studies are required to define its exact mechanism of action.

Conflicts of interest

All authors have no conflicts of interest to declare.

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