The effect of taxifolin on high-dose-cisplatin-induced oxidative liver injury in rats

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D – writing the article; E – critical revision of the article; F – final approval of the article

Abstract

Background. Cisplatin is a non-specific platinum-based (derivative) chemotherapeutic agent that causes an increase in free radicals activity in the liver. Antioxidant activity of taxifolin has been demonstrated previously, and it has been reported that taxifolin inhibits the hydroxyl, radical in experimental studies.

Objectives. No studies were found in the current literature examining the protective effect of taxifolin on cisplatin-induced oxidative liver damage. We aimed to determine the protective effect of taxifolin on cisplatin-induced hepatotoxicity in an experimental study.

Materials and methods. In total, 18 albino Wistar male rats were assigned into 3 groups: healthy controls (HC group), 5 mg/kg of cisplatin administered for 8 days (CIS group) and 50 mg/kg of taxifolin + 5 mg/kg of cisplatin administered for 8 days (TCG group). Malondialdehyde (MDA), total glutathione (tGSH), total oxidant (TOS), and total antioxidant (TAS) capacity levels were measured in the extracted liver tissue.

Results. Liver tissue MDA and TOS levels were significantly higher in the CIS group. In contrast, tGSH and TAS levels were significantly lower in the CIS group, administered cisplatin alone (p < 0.001), compared to other groups. In the TCG group, administered cisplatin + taxifolin, MDA and TOS levels were significantly lower, whereas tGSH and TAS levels were significantly higher than in the CIS group (p < 0.001).

Conclusions. These results suggest that taxifolin may be useful in preventing cisplatin-related liver injury.

Key words: cancer, hepatotoxicity, cisplatin, taxifolin, antioxidant
Background

Cisplatin is a non-specific platinum-based (derivative) chemotherapeutic agent used to treat stomach, testis, ovarian, bladder, kidney, ureterovesical, head, and neck cancer. It has been reported to cause serious toxic effects in many organs and systems during treatment. The chemotherapeutic efficacy of cisplatin increases together with the dose. However, the increase in dose causes side effects such as nephrotoxicity, ototoxicity, neurotoxicity, hepatotoxicity, nausea, vomiting and, in 67% of patients, diarrhea, limiting its clinical use. Cisplatin also causes an increase in free radicals activity in the liver, leading to oxidative stress. Intravenously administered cisplatin (180–480 mg/m²) can also reach high concentrations in the liver, except in kidney and intestine, and may cause significant toxicity. Therefore, it has been reported that the most critical side effects are hepatotoxicity and nephrotoxicity. Hepatotoxicity manifests as an increase in serum transaminase (alanine aminotransferase (ALT) and aspartate aminotransferase (AST)) levels. Even the treatment dose used as a tumor suppressor has been shown to cause hepatotoxicity. Although the mechanism of injury of cisplatin is not well known, some evidence suggests that it is caused by oxidative stress, due to reactive oxygen species (ROS) activity. The oxidative stress induced by ROS causes a decrease in glutathione (GSH), which is an endogenous antioxidant.

Furthermore, ROS affect cell membrane lipid peroxidation and lead to occurrence of more toxic products such as malondialdehyde (MDA). This suggests that antioxidant therapy may help to prevent or reduce cisplatin-induced hepatotoxicity. Studies have shown that various antioxidants are protective against cisplatin hepatotoxicity. However, hepatotoxicity caused by cisplatin remains one of the crucial reasons for limiting its use at the desired dose. Therefore, several kinds of research to alleviate cisplatin hepatotoxicity are being pursued. In our study, we investigated whether taxifolin (3,3',4',5,7-pentahydroxiflavanone), a flavonoid contained in onions, milk thistle, French maritime bark, and Douglas fir bark, has a protective effect against the hepatotoxicity of cisplatin. Antioxidant activity of taxifolin has been documented. In experimental studies, taxifolin has been reported to inhibit the hydroxyl radical (•OH), a ROS in the cell, and protects cellular DNA from oxidative damage by antioxidant activity. These data suggest that taxifolin will protect liver tissue against the oxidative damage caused by cisplatin.

Objectives

No data were found in literature reviews regarding the protective effects of taxifolin on cisplatin-induced liver toxicity. Therefore, our study aimed to investigate, through biochemical analysis, whether or not taxifolin has a protective effect on cisplatin-induced hepatotoxicity in rats.
Malondialdehyde analysis

Malondialdehyde measurements were based on the method used by Ohkawa et al., involving spectrophotometrically measured absorbance of the pink-colored complex formed by thiobarbituric acid (TBA) and MDA. Briefly, 25 µL of tissue homogenate was added to a solution containing 25 µL of 80 g/L sodium dodecyl sulfate and 1 mL of mixture solution (20 g/L of acetic acid + 1.06 g 2-thiobarbituric acid + 180 mL of distilled water). The mixture was incubated at 95°C for 1 h. Upon cooling, the mixture was centrifuged for 10 min at 4000 rpm. The absorbance of the supernatant was measured at 532 nm. The standard curve was obtained by using 1,1,3,3-tetramethoxypropane.19

Total glutathione (GSH) analysis

The amount of GSH in the total homogenate was measured with the method used by Sedlak et al. and Baker et al., with some modifications. The principle of the method is that the color intensity of dark yellow 5-thio-2-nitrobenzoic acid (TNB) is released through the reduction of Ellman’s reagent (5,5’-dithiobis(2-nitrobenzoic acid) – DTNB) by free thiol groups, and through the reduction of Ellman’s reagent (5,5’-dithiobis(2-nitrobenzoic acid) – DTNB) by free thiol groups, and measured with the method used by Sedlak et al. The amount of GSH in the total homogenate was measured at 412 nm using a spectrophotometer. The TOS method is based on bleaching the characteristic color of the groups in terms of TOS levels (p < 0.05). The levels significantly were compared in the study groups, there was a statistically significant difference among the groups (p < 0.001), but no significant difference was found compared to the HC group (p > 0.05). When the mean tGSH levels were compared in the study groups, there was a statistically significant difference among the groups (p < 0.05).

Statistical analyses

Statistical analyses were performed using IBM SPSS software v. 21.0 (IBM Corp., Armonk, USA). The estimated power (1-beta) test value was calculated as 0.99 with the G-Power Program (https://www.psychologie.hhu.de/arbeitsgruppen/allgemeine-psychologie-und-arbeitspsychologie/gpower). Numerical variables were expressed as a median (min–max). For the analysis of continuous variables, the Kruskal–Wallis test was performed. For between 2 groups, Dunn’s test (a post hoc comparison test) was performed. The minimum criterion for statistical significance was set at p < 0.05 for all comparisons.

ALT and AST analysis

Serum ALT and AST analysis were performed on a Roche Cobas 8000 autoanalyzer using the spectrophotometric method and kits from the same company (REF: 207649557-322, 20764949-322, respectively; Roche Diagnostics, Basel, Switzerland). The principle of both measurements is based on measuring the absorbance change of NADH at 340 nm.

Results

As shown in Table 1, serum ALT and AST activity were significantly higher in the CIS group compared with the HC and TCG groups (p < 0.001).

When mean MDA levels were compared, statistically significant differences were found between the study groups (p < 0.05). The amount of MDA in the TCG group was significantly decreased compared to the CIS group (p < 0.001), but no significant difference was found compared to the HC group (p > 0.05). When the mean tGSH levels were compared in the study groups, there was a statistically significant difference among the groups (p < 0.05). In the TCG group, tGSH levels were significantly higher (p < 0.001) compared to the CIS and HC groups (Table 1).

There was a statistically significant difference among the groups in terms of TOS levels (p < 0.05). The levels of a more stable ABTS (2,2’-azino-bis (3-ethylbenzothiazoline-6-sulfonic acid)) radical cation by antioxidants, and measurements are performed at 660 nm. The results are expressed as nmol Trolox equivalent/L. In the TOS method, the oxidants present in the sample oxidized the ferrous ion-o-dianisidine complex to ferric ion. The oxidation reaction was enhanced by glycerol molecules, which are abundantly present in the reaction medium. The ferric ion produced a colored complex with xylenol orange in an acidic medium. The color intensity, which could be measured at 530 nm spectrophotometrically, was related to the total amount of oxidant molecules present in the sample. The results are expressed as nmol hydrogen peroxide (H₂O₂) equivalent/L.
of TOS in the liver tissue of the TCG group was significantly lower compared to the CIS group (p < 0.001). There was no significant difference between the TCG and HC groups in terms of TOS level (p > 0.05). When the mean TAS levels were compared in the study groups, statistically significant differences were found (p < 0.05). The level of TAS in the liver tissue of the TCG group was significantly higher compared to the CIS group (p < 0.05) (Table 1). However, the difference between TAS levels in TCG and HC groups was insignificant (p > 0.05) (Table 1).

### Discussion

In our study, the effect of taxifolin on cisplatin-induced oxidative liver injury in rats was investigated biochemically. There are many studies showing cisplatin-induced liver damage. In our research, in parallel with the literature, serum ALT and AST levels increased after exposure to high doses of cisplatin. These values decreased after administration of taxifolin. Our results showed that cisplatin caused an increase in MDA and TAS levels and a decrease of TSH and TAS levels in animal liver tissue. Oxidant-antioxidant parameters are used to determine oxidative damage. Increased MDA production in the CIS group suggests that lipid peroxidation is exacerbated in liver tissue cells in this group. The ROS lead to the peroxidation of lipids in the cell membrane and result in the secretion of more toxic products, such as MDA, from lipids. Malondialdehyde can cause damage in membrane proteins by inactivating receptors and membrane-bound enzymes in membranes, leading to cross-linking and polymerization of membrane components. The LPO reaction is either terminated by scavenging antioxidant reactions or continues with auto-catalytic spreading reactions. Studies have suggested that ROS also play a role in the pathogenesis of the hepatotoxic effect of cisplatin. Taxifolin used against cisplatin-induced oxidative liver damage significantly inhibited an increase in MDA and TOS levels, and decreased TSH and TAS levels. It has been documented in previous studies that taxifolin reduced intracellular free radical levels, inhibited MDA and prevented the consumption of antioxidants. In another study, taxifolin was found to increase the effect of GSH.

### Table 1. The results of biochemical evaluations among groups

<table>
<thead>
<tr>
<th>Variables</th>
<th>CIS (median (min–max) (n = 6))</th>
<th>TCG (median (min–max) (n = 6))</th>
<th>HC (median (min–max) (n = 6))</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDA (µmol/g protein)</td>
<td>8.65 (7.9–9.1)</td>
<td>2.50 (2.2–3.7)</td>
<td>3.05 (2.8–3.9)</td>
<td>0.002</td>
</tr>
<tr>
<td>tGSH (nmol/g protein)</td>
<td>1.65 (1.4–2.1)</td>
<td>4.85 (4.2–5.1)</td>
<td>5.45 (5.1–5.7)</td>
<td>0.001</td>
</tr>
<tr>
<td>TAS (nmol/diet exq/mg protein)</td>
<td>5.45 (4.7–6.1)</td>
<td>12.00 (10.0–16.0)</td>
<td>14.50 (12.0–17.0)</td>
<td>0.002</td>
</tr>
<tr>
<td>TOS (nmol H2O2 eq/mg protein)</td>
<td>18.00 (15.0–21.0)</td>
<td>7.00 (6.1–8.0)</td>
<td>6.25 (5.1–7.1)</td>
<td>0.001</td>
</tr>
<tr>
<td>AST (IU/L)</td>
<td>200.06 (178.5–235.4)</td>
<td>48.71 (38.1–56.3)</td>
<td>40.43 (34.7–48.8)</td>
<td>0.001</td>
</tr>
<tr>
<td>ALT (IU/L)</td>
<td>100.12 (89.2–116.8)</td>
<td>38.07 (34.5–41.4)</td>
<td>30.66 (23.6–34.5)</td>
<td>0.001</td>
</tr>
</tbody>
</table>

HC – healthy control; CIS – 5 mg/kg cisplatin group; TCG – 50 mg/kg taxifolin + 5 mg/kg cisplatin group; MDA – malondialdehyde; tGSH – total glutathione; TAS – total antioxidant status; TOS – total oxidative status; ALT – alanine aminotransferase; AST – aspartate aminotransferase.

a, b – statistically significantly different compared with CIS; b, c – statistically significantly different compared with TCG.
As noted above, taxifolin is a flavonoid. Previous studies have expressed that flavonoids demonstrate antioxidant activity by inhibiting lipid peroxidation and enzymatic reactions responsible for the formation of free radicals. In a recent study, it was suggested that taxifolin produces a hepatoprotective effect with antioxidant activity by inhibiting the lipid peroxidation pathway. Viewed in total, data from the reviewed literature are in agreement with our experimental results.

**Limitations**

In our study, it was necessary to investigate the effect of taxifolin on pro-inflammatory cytokine levels, known to play a role in the pathogenesis of cisplatin toxicity, in order to clarify its protective effect against hepatotoxicity caused by cisplatin. In addition, histopathological findings should support all biochemical findings. The small number of animals in the groups could be considered as another limitation.

**Conclusions**

As a result, cisplatin caused oxidative damage in the liver tissue of animals. In the CIS group, the antioxidant–antioxidant balance deteriorated against the oxidants. Taxifolin prevented the increase in oxidants and decreased antioxidants in the liver tissue injured due to cisplatin. Taxifolin prevented the change in favor of oxidants in the antioxidant–antioxidant balance. These findings suggest that taxifolin could be a clinically beneficial agent for treating hepatotoxicity resulting from chemotherapy procedures.

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