Silibinin attenuates methotrexate-induced pulmonary injury by targeting oxidative stress

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Abstract. The aim of the present study was to assess the protective effect of silibinin against methotrexate (MTX)-induced pulmonary toxicity. Rats were divided into four groups (MTX, MTX + silibinin, silibinin and control. MTX was injected intraperitoneally (i.p) into female Wistar rats (10 mg/kg/day for 3 days), which resulted in significant increases in the serum levels of alanine aminotransferase, aspartate aminotransferase and oxidant enzymes, including nitric oxide and myeloperoxidase. Furthermore, significant reductions were detected in the serum activity levels of the antioxidative enzymes, glutathione peroxidase and superoxide dismutase, when compared with the control group. However, administration of silibinin (100 mg/kg/day for 10 days, i.p.) was shown to ameliorate the MTX-induced pulmonary toxicity, as indicated by the normalization of the oxidative stress parameters. Furthermore, silibinin treatment was demonstrated to reduce the histopathological changes associated with MTX. In conclusion, silibinin exhibited protective effects against MTX-induced pulmonary toxicity, which may be attributed to its antioxidant activity.

Introduction

Methotrexate (MTX) is an antineoplastic agent that is associated with folic acid metabolism. MTX inhibits the synthesis of DNA, RNA, thymidylate and proteins. As a result of this activity, MTX is commonly used in cancer treatment, in addition to the treatment of non-neoplastic diseases, including rheumatoid arthritis and psoriasis (1). Previous studies have demonstrated that liver, kidney, brain and lung toxicities are potential side effects of MTX use (2). The predominant manifestation of MTX-induced lung toxicity is pulmonary fibrosis (3). In addition to pulmonary fibrosis, MTX treatment has been associated with acute interstitial pneumonitis (4-8).

Silibinin (C15H22O16; molecular weight, 482.44 g/mol) is isolated from the seeds of Silybum marianum L. (9). The compound is known to exhibit antioxidative activity, which has been previously reported to result in a number of biologically protective effects, including anti-inflammatory, antitumor and hepatoprotective effects (10-13). Silibinin has been demonstrated to be a potent antioxidant, supporting the capacity of cellular antioxidative agents, such as glutathione (GSH) and superoxide dismutase (SOD), to target reactive oxygen species. This activity may partially explain the effectiveness of silibinin in the prevention of hepatic injury, whether this injury is as a result of disease or exposure to toxins, since this antioxidant activity may mitigate the oxidative stress associated with hepatic injury, subsequently preventing the induction of lipid peroxidation (14). Furthermore, silibinin has been widely investigated for anticancer efficacy in a broad range of cancer models. As a consequence of the general anticancer properties associated with flavonoids collectively, there has been significant interest in the possibility of using silibinin as a chemopreventive agent (11,15). Silibinin has been traditionally used in folk medicine, and acute and chronic doses of silibinin administration in animals and humans have resulted in no significant toxicity. However, to the best of our knowledge, an LD50 for silibinin has not been reported in rodent studies. Silibinin intake has been demonstrated to be safe due to its wide usage as a dietary supplement, with sufficient tolerability and minimal toxicity (15). Therefore, the aim of the present study was to assess the possible protective effects of silibinin against MTX-induced pulmonary toxicity in rats.

Materials and methods

Animals. Animal experiments were approved by the Animal Ethical Committee of Suleyman Demirel University (B.30.2.SDU.0.05.06.00-196, 2012; Isparta, Turkey), and the study was conducted in accordance with the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals (8th Edition, 2011). In total, 32 female albino Wistar rats (age, 8-10 weeks) were obtained from the Experimental
Research Centre of Suleyman Demirel University and housed in an environmentally controlled room at 21±1°C and 75±5% humidity, under a 12-h light/dark cycle. The animals were acclimatized for 1 week prior to the study, and had free access to standard laboratory feed and water.

**Experimental protocol.** Rats were divided into four groups. Since silibinin was solubilized in dimethyl sulfoxide (DMSO), an equal amount of DMSO (10%/v/v) was included in the injections administered to the control group rats. The silibinin group rats received 100 mg/kg/day silibinin (Sigma-Aldrich, St. Louis, MO, USA), which was administered via intraperitoneal (i.p.) injection for 10 days (16). The MTX group rats were administered 10 mg/kg/day MTX (i.p.) on days 7-9, the commercially available form of the drug was used, it was taken from pharmacy with our own means (17). Finally, the MTX + silibinin group rats were cotreated with 100 mg/kg/day silibinin for 10 days and 10 mg/kg/day MTX for 3 days. In all the groups on day 14, anesthesia was induced by a single i.p. injection of 50 mg/kg ketamine (Ketalar®; Pfizer, Inc., Istanbul, Turkey) and 5 mg/kg xylasine (Rompun®; Bayer, Istanbul, Turkey). Blood samples were collected via cardiac puncture for serum analyses and the lungs were harvested for histological and immunohistochemical analysis.

**Biochemical evaluation.** All the biochemical analyses were performed in the Department of Biochemistry at Mugla Sıtkı Kocman University (Mugla, Turkey).

**Determination of alanine aminotransferase (ALT) and aspartate aminotransferase (AST) activity levels.** Serum activity levels of AST and ALT were calculated spectrophotometrically using Beckman Coulter kits and a UniCel DxC 800 Synchron autoanalyzer (Beckman Coulter, Inc., Brea, CA, USA).

**Determination of SOD activity.** Tissue samples were homogenized at 4,200 x g on ice in 5-10 ml cold buffer [20 mM HEPES buffer (pH 7.2) containing 1 mM EDTA, 210 mM mannitol and 70 mM sucrose per gram tissue]. Subsequently, the samples were centrifuged at 1,500 x g for 5 min at 4°C, and the supernatant was removed. In addition, the blood samples were centrifuged at 2,000 x g for 15 min at 4°C, after which the top yellow serum layer was pipetted off, without disturbing the white buffy coat. The serum was diluted 1:5 with sample buffer. The SOD activity was measured in the supernatant and serum using a SOD assay kit (Cayman Chemical Company, Inc.) with an ELx-800 absorbance reader (Bio-Tek Instruments, Inc., Winooski, VT, USA). The assay was based on the detection of superoxide radicals generated by xanthine oxidase and hypoxanthine. One unit of SOD was defined as the amount of enzyme required to induce 50% dismutation of the superoxide radical. The results are expressed in U/mg protein tissue for the liver tissue and U/ml for the serum.

**Determination of glutathione peroxidase (GPx) activity.** Tissue samples were homogenized in 5-10 ml cold buffer [50 mM Tris-HCl (pH 7.5), 5 mM EDTA and 1 mM DTT], and centrifuged at 10,000 x g for 15 min at 4°C, after which the supernatant was removed. In addition, the blood samples were centrifuged at 700-1,000 x g for 10 min at 4°C, and the plasma was removed. GPx activity was measured in the liver tissue and plasma samples using a GPx assay kit (Cayman Chemical Company, Inc.) with an ELx-50 microplate strip washer. GPx activity was measured indirectly by a coupled reaction with glutathione reductase, where the oxidized glutathione was produced upon the reduction of hydroperoxide by GPx.

**Determination of nitric oxide (NO) levels.** Tissue samples were homogenized in phosphate-buffered saline (pH 7.4) and centrifuged at 10,000 x g for 20 min to isolate the supernatant. A total NO assay was performed via spectrophotometry at 540 nm using a nitrate/nitrite colorimetric assay kit (Cayman Chemical Company, Inc.) with an ELx-50 microplate strip washer. The assay was based on nitrate and nitrite determinations, in which nitrate and nitrite were the stable end products of the reaction of NO with molecular oxygen. The total accumulation of nitrate and nitrite in the serum and liver tissue samples was evaluated, and expressed in µM/protein.

**Determination of myeloperoxidase (MPO) activity.** Quantitative detection of MPO activity was conducted using an enzyme-linked immunosorbent assay kit (MPO Instant ELISA; eBioscience, Inc., Vienna, Austria) in an ELx-50 microplate strip washer. The results are expressed in pg/ml/protein.

**Histopathological analysis of the lung tissue.** Histopathological analyses were performed in the Department of Pathology at Mugla Sıtkı Kocman University. Rat lung samples from the different groups were fixed in 10% neutral buffered formalin for 24 h. After washing with tap water, serial dilutions of alcohol (methyl, ethyl and absolute ethyl) were applied for dehydration. The specimens were cleared in xylene, and embedded in paraffin at 56°C in an oven for 24 h. Paraffin bees wax tissue blocks were prepared for sectioning at 5 µm thickness using a microtome. Lung tissues were stained with hematoxylin and eosin and observed under a BX46 light microscope (Olympus Corporation, Tokyo, Japan) for histopathological evaluation. Pulmonary damage was evaluated using six parameters, which included interstitial lymphocytic inflammation, interstitial fibrosis, type 2 pneumocyte infiltration, intraalveolar/interstitial macrophage existence, eosinophil existence and granuloma existence (18). Each parameter was scored semiquantitatively with regard to the severity using the following system: 0 (absent), no interstitial lymphocytic inflammation, fibrosis, type 2 pneumocyte infiltration, intraalveolar/interstitial macrophage existence, eosinophil existence and granuloma existence (18). Parametric methods were used

**Statistical analysis.** SPSS software, version 21.0 (IBM SPSS, Armonk, NY, USA) and PAST software (http://folk.uio.no/ohammer/past/) were used for data analysis. For comparisons among multiple independent groups one-way analysis of variance (ANOVA) was used. If significance was identified in the test we used the Least Significant Difference (LSD) test for post-hoc analysis. Parametric methods were used
for the analysis of the data with a normal distribution, while non-parametric methods were used for the analysis of the variations without a normal distribution. For comparisons among multiple independent groups, ANOVA (Robust test: Brown-Forsythe), one of the parametric methods was used, while for post-hoc analysis, Fisher’s LSD test was used. For the comparison of categorical data, Pearson’s $\chi^2$ test was conducted using the Monte Carlo simulation technique.

Quantitative data are expressed as the mean ± standard deviation, and categorical data are expressed as a number and percentage. Data were analyzed in the 95% confidence level, where P<0.05 was considered to indicate a statistically significant difference.

Results

Biochemical analysis. Serum levels of ALT and AST were significantly increased in the MTX group when compared with the control and silibinin groups (P<0.05; Table I). In the MTX + silibinin group, the ALT and AST levels in serum decreased significantly compared with the MTX group (P<0.05). Furthermore, in the MTX group, the SOD levels decreased significantly when compared with the control group (P<0.05). The levels of GPx and SOD increased in the MTX + silibinin group when compared with the MTX group, while the MPO values decreased significantly (P<0.05).

Histopathological alterations. When compared with the control (Fig. 1) and silibinin groups (Fig. 2), the pulmonary damage appeared to be significantly increased in the MTX group (Fig. 3). Similarly, when the MTX + silibinin group was compared with

Table I. Effect of silibinin on the biochemical parameters associated with MTX-induced pulmonary toxicity.

<table>
<thead>
<tr>
<th>Group</th>
<th>ALT (U/L)</th>
<th>AST (U/L)</th>
<th>SOD (U/ml)</th>
<th>GPx (U/ml)</th>
<th>NO (µm/g)</th>
<th>MPO (ng/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>20.86±3.34</td>
<td>62.75±6.24</td>
<td>0.89±0.14</td>
<td>0.88±0.14</td>
<td>2.25±0.36</td>
<td>2.63±0.76</td>
</tr>
<tr>
<td>Silibinin</td>
<td>16.25±2.60</td>
<td>49.38±7.90</td>
<td>3.54±0.57</td>
<td>0.24±0.04</td>
<td>3.05±0.49</td>
<td>2.70±0.53</td>
</tr>
<tr>
<td>MTX</td>
<td>55.38±8.86</td>
<td>86.88±13.90</td>
<td>0.48±0.08</td>
<td>0.19±0.03</td>
<td>4.49±0.40</td>
<td>4.3±0.56</td>
</tr>
<tr>
<td>MTX + silibinin</td>
<td>31.25±5.01</td>
<td>60.25±9.64</td>
<td>5.40±0.86</td>
<td>0.63±0.10</td>
<td>1.06±0.17</td>
<td>2.33±0.37</td>
</tr>
</tbody>
</table>

Results are presented as the mean ± standard deviation. a P<0.05, vs. control group; b P<0.05, vs. silibinin group; c P<0.05, vs. MTX group. MTX, methotrexate; ALT, alanine aminotransferase; AST, aspartate aminotransferase; SOD, superoxide dismutase; GPx, glutathione peroxidase; NO, nitric oxide; MPO, myeloperoxidase.

Table II. Histopathological damage parameters detected in the rat pulmonary tissue samples.

<table>
<thead>
<tr>
<th>Histopathological parameter</th>
<th>Control</th>
<th>Silibinin</th>
<th>MTX</th>
<th>MTX + silibinin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Interstitial lymphocytic inflammation</td>
<td>0.125</td>
<td>0.375</td>
<td>1.25$^{ab}$</td>
<td>0.25$^c$</td>
</tr>
<tr>
<td>Interstitial fibrosis</td>
<td>0</td>
<td>0</td>
<td>0.875$^{ab}$</td>
<td>0$^c$</td>
</tr>
<tr>
<td>Type 2 pneumocyte infiltration</td>
<td>0</td>
<td>0</td>
<td>0.5$^{ab}$</td>
<td>0$^c$</td>
</tr>
<tr>
<td>Intraalveolar/interstitial macrophage existence</td>
<td>0</td>
<td>0.375</td>
<td>0.5</td>
<td>0.125$^c$</td>
</tr>
<tr>
<td>Eosinophil existence</td>
<td>0</td>
<td>0.5</td>
<td>0.875$^a$</td>
<td>0.125$^c$</td>
</tr>
<tr>
<td>Granuloma existence</td>
<td>0</td>
<td>0</td>
<td>0.25</td>
<td>0</td>
</tr>
</tbody>
</table>

Results are presented as the mean ± standard deviation. a P<0.05, vs. control group; b P<0.05, vs. silibinin group; c P<0.05, vs. MTX group. MTX, methotrexate.
the MTX group, the damage was observed to significantly decrease (Fig. 4 and Table II). MTX administration was shown to induce interstitial lymphocytic inflammation, interstitial fibrosis, type 2 pneumocyte hyperplasia and eosinophil infiltration in the pulmonary tissue. However, the MTX + silibinin group exhibited significantly decreased scores for interstitial lymphocytic inflammation, interstitial fibrosis and type 2 pneumocyte hyperplasia when compared with the MTX group, indicating that silibinin exerted beneficial effects (Table II).

Discussion

The present study evaluated the interactions between silibinin and MTX, and their effects on lung tissue. The results indicate that treatment with silibinin ameliorated MTX-induced alterations in the serum ALT and AST levels. In addition, silibinin significantly mitigated the oxidation in the serum induced by MTX, as manifested by the decreased MPO and NO levels, accompanied by enhanced SOD and GPx activity. Furthermore, histopathological analysis demonstrated that administration of silibinin mitigated a number of the histopathological changes induced by MTX.

A previous study indicated that MTX exposure activates components of the mitogen-activated protein kinase (MAPK) signaling pathway (19). Activation of the MAPK and Akt pathways may in turn activate the transcription factors, activator protein-1 and nuclear factor-κB, which are crucial for the regulation of inflammation (20). Inflammatory cells that infiltrate into the tissue are known to cause tissue damage through activating oxidation systems, such as MPO. Furthermore, inflammatory cells are known to induce cell death by generating DNA damage via oxidation systems (21).

Silibinin possesses marked antioxidative, anticancer, anti-inflammatory and cancer chemopreventive properties (22,23). Previous studies have reported that silibinin targets multiple signaling pathways, including those associated with oxidative stress and inflammation, to subsequently prevent tissue injuries and cancer by genotoxicity and other agents, which are similar to the pathways triggered following vesicant exposure (22,24,25). Therefore, silibinin was hypothesized to exhibit notable efficacy in attenuating MTX-induced lung injury, since MTX is known to trigger oxidative stress. The results of a previous study by Tewari-Singh et al (21) indicated that vesicant drugs exert a protective effect on the skin, combined with the anti-inflammatory and antioxidative effects of silibinin via the MAPK pathway. In addition, the results of the present study indicated that silibinin administration in combination with MTX decreased the activation of serum oxidation enzyme systems, while increasing antioxidant enzyme levels (Table II). Furthermore, silibinin appeared to reduce inflammatory cell infiltration and MTX-induced changes in the lung tissue (Table II). In accordance with previous literature (14), the current results indicated that silibinin is able to decrease pulmonary oxidative stress, possibly via the MAPK pathway.

The potential for hepatic toxicity remains a concern for physicians who are increasingly using MTX (26). The preventive effect of silibinin against liver damage has been demonstrated in a previous study (10). In the present study, cotreatment with MTX and silibinin was demonstrated to result in significantly decreased serum levels of AST and ALT when compared with MTX treatment alone (P<0.05; Table I). These findings indicate that silibinin decreases MTX-associated tissue damage, which may provide an advantage for clinicians regarding the use of this drug.

In conclusion, silibinin was demonstrated to protect the lung tissue against MTX-induced pulmonary toxicity in rats. The antioxidant activity of silibinin may be the primary factor responsible for such pulmonary protective effects. Therefore,
silibinin represents a potential candidate agent for the prevention of lung injury, which is a major and dose-limiting side effect of MTX therapy.

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References


