The protective effect of vildagliptin in chronic experimental cyclosporine A-induced hepatotoxicity

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Abstract: The study examined the effect of dipeptidyl peptidase-4 (DPP-4) inhibitor, vildagliptin, in cyclosporine (CsA)-induced hepatotoxicity. Rats were divided into 4 groups treated for 28 days: control (vehicle), vildagliptin (10 mg/kg, orally), CsA (20 mg/kg, s.c.), and CsA-vildagliptin group. Liver function was assessed by measuring serum levels of aspartate aminotransferase (AST), alanine aminotransferase (ALT), gamma glutamyltransferase (γGT), lactate dehydrogenase (LDH), and albumin, and histopathological changes of liver were examined. Oxidative stress markers were evaluated. Assessment of nuclear factor-kappa B (NF-κB) activity in hepatic nuclear extract, serum DPP-4, and expression of Bax and Bcl2 were also done. CsA-induced hepatotoxicity was evidenced by increase in serum levels of AST, ALT, γGT; a decrease in serum albumin; and a significant alteration in hepatic architecture. Also, significant increase in thiobarbituric acid reactive substance (TBARS) and decrease in superoxide dismutase (SOD), catalase (CAT), glutathione peroxidase (GPx), and glutathione (GSH) levels, increased expression of NF-κB and serum DPP-4 level were observed upon CsA treatment. Vildagliptin significantly improved all altered parameters induced by CsA administration. Vildagliptin has the potential to protect the liver against CsA-induced hepatotoxicity by reducing oxidative stress, DPP-4 activity, apoptosis, and inflammation.

Key words: vildagliptin, cyclosporine, oxidative stress, inflammation, apoptosis.

Introduction
Cyclosporine (CsA) is an immunosuppressive agent that is commonly used in the treatment of many autoimmune diseases and after organ transplantations (Kwak and Mun 2000; Rezzani et al. 2001). However, its use is limited due to several side effects, including nephrotoxicity, cardiotoxicity, hypertension, and hepatotoxicity. Impairments in liver architecture have been reported in rats given CsA for 2–13 weeks (Rezzani 2004; Josephine et al. 2008; Lee 2010). Cholestasis, hyper-bilirubinemia, hypoproteinemia, increased alkaline phosphatase and transaminase activities, bile salts in the blood, and inhibition of protein synthesis and disturbed lipid secretion in both human and experimental animals were found to characterize CsA-induced hepatotoxicity (McCashland et al. 1994; Rezzani 2004). Additionally, alterations in bile formation, the capacity of the liver to excrete organic anions and xenobiotics, and changes in the hepatic content of glutathione (GSH) are also prominent in CsA-provoked liver damage (Morán et al. 1998). Several mechanisms are involved in CsA-induced hepatotoxicity including reactive oxygen species (ROS) production, oxidative stress, and depletion of the hepatic antioxidant system (Rezzani et al. 2005).

Dipeptidyl peptidase-4 (DPP-4) is a membrane-associated peptidase known as CD26. Vildagliptin is a DPP-4 inhibitor that increases plasma level of active glucagon-like peptide-1 (GLP-1), improves glucose-dependent insulin secretion, improves β-cell function, improves insulin sensitivity, reduces inappropriate glucagon se-
creatinine, and reduces fasting and postprandial glucose (Barnett 2006). DPP-4 has widespread organ distribution throughout the body and exerts pleiotropic effects via its peptidase activity. DPP-4 is involved in immune stimulation, binding to and degradation of ECM, resistance to anti-cancer agents, and lipid accumulation (Itou et al. 2013). Tumor necrosis factor-α (TNF-α), interferons, retinoic acid, high insulin levels, glucose, and hypoxia are some of the factors that influence the production and release of DPP-4; however, the mechanisms of action involved remain unknown. DPP-4 activity levels in the serum have been shown to increase under conditions such as nonalcoholic fatty liver disease, rheumatoid arthritis, and inflammatory bowel disease (Das et al. 2014). Also, DPP-4 is highly expressed in the liver, and recent data suggested that DPP-4 is involved in the development of various chronic liver diseases such as hepatitis C virus infection, nonalcoholic fatty liver disease, and hepatocellular carcinoma. DPP-4 expression has similar lobular heterogeneity seen in the expression of cytochrome p450, gamma-glutamyl transpeptidase (GGT), and glutamine synthetase (Mentzel et al. 1996; Gorrell et al. 2001). This heterogeneous lobular distribution suggests that DPP-4 may be involved in the regulation of hepatic metabolism. Furthermore, DPP-4 occurs in hepatic stem cells and plays a crucial role in hepatic regeneration (Itou et al. 2013). Therefore, the present study was undertaken to assess the possible beneficial effects of vildagliptin in CsA-induced hepatotoxicity in rats.

Materials and methods

Animals

Thirty-two male Sprague-Dawley rats (purchased from Urology and Nephrology Center, Mansoura University, Egypt), weighing between 200 and 250 g, were housed for 28 days in a well-ventilated room with a 12 h light – 12 h dark cycle and were allowed access to water ad libitum and fed a normal rat chow. The experimental protocol conducted in the study complies with the ethical guidelines and the principals of care, use, and handling of experimental animals adopted by The Research Ethics Committee, Faculty of Pharmacy, Mansoura University, Egypt, which is in accordance with “Principles of Laboratory Animals Care” (NIH publication No. 85-23, revised 1985).

Experimental design

Rats were randomly divided into 4 groups with 8 rats each and were treated as follows:

Group 1 (control group), rats received 2 ml/kg sunflower oil, refined bleached deodorized (RBD) (CsA vehicle) by s.c. injection and 2 ml/kg 0.5% carboxymethyl cellulose (vildagliptin vehicle) by oral gavage daily for 28 days;

Group 2 (vildagliptin group), rats treated with daily oral vildagliptin (10 mg/kg, Novartis Pharma AG, Basel, Switzerland) by oral gavage daily for 28 days;

Group 3 (CsA group), rats received a daily dose of 20 mg/kg of CsA (Novartis Pharma AG, Basel, Switzerland) by s.c. injection;

Group 4 (CsA-vildagliptin group), rats given daily doses of vildagliptin and CsA for 28 days. This treatment schedule and dose level were chosen based on the results obtained in pilot experiments using lower doses of vildagliptin (1 and 5 mg/kg), which failed to decrease the elevated serum ALT and AST levels (data not shown).

At the end of day 28 of the experiment, each rat was anesthetized by thiopental sodium solution at a dose of 30 mg/kg i.p. (Wixson et al. 1987). Blood was collected from the retro-orbital sinus and serum was separated by centrifugation at 2700 g for 20 min then used for analyzing the biochemical parameters.

The liver was rapidly excised; one part was used for histopathological examination of biological markers after homogenization and the other part was used for histopathological examination.

Preparation of liver homogenate

Excised livers were rinsed with ice-cold isotonic saline (0.9%), weighed, and then homogenized immediately in ice-cold 1.15% KCl solution to give a 10% (w/v) homogenate. Homogenates were centrifuged at 10 000 g for 20 min at 4°C to obtain the supernatant fractions, which were used for determination of oxidative stress parameters and Bax and Bcl2 expression.

Assessment of hepatic oxidative markers

Enzymatic antioxidants were determined in liver homogenates by estimating SOD, CAT, and GPx activity; non-enzymatic antioxidants by GSH; and lipid peroxidation by measuring thiobarbituric acid reactive substance (TBARS) levels using commercially available kits (Bio-diagnostic Company, Giza, Egypt).

Assessment of biochemical assays

Determination of serum levels of AST, ALT, γGT, LDH, and albumin were done using commercially available kits (bioMérieux, Marcy-l’Étoile, France) according to manufacturer’s instructions.

Hepatic nuclear factor-kappa B (NF-κB) activation

The transcription factor NF-κB activation was measured according to manufacturer’s instructions using a commercially available enzyme linked immunosorbent assay (ELISA) kit (Trans-AM NF-κB p65 Transcription Factor Assay Kit, Active Motif, California, USA). Nuclear protein extract was obtained using Nuclear Extract Kit (Active Motif) according to manufacturer’s instructions. Subsequently, 15 μg nuclear protein extract was used to assay for NF-κB activation. Values were represented as OD 450 nm.

Assessment of DPP-4 level

The amount of DPP-4 level in serum was measured using ELISA kit (antibodies-online Inc., Atlanta, USA) according to the manufacturer’s directions.

Assessment of Bax and Bcl-2 in hepatic tissue

Rat Bax ELISA kit and Rat Bcl-2 ELISA kit (antibodies-online Inc., Atlanta, USA) were used to evaluate the expression of Bax and Bcl-2 proteins in the rat liver homogenates according to manufacturer’s instructions. The absorbance was read at 450 nm.

Histopathology of liver tissue

Liver tissues were embedded in paraffin, and 4–5 μm thick sections were stained with hematoxylin and eosin. The tissues were examined under a light microscope by histologists blinded to the treatment groups.

Statistical analysis

Data are expressed as mean ± standard deviation (SD). Statistical analysis was carried out by one-way analysis of variance (ANOVA) followed by post hoc Tukey-Kramer multiple comparisons test. Statistical calculations were carried out using Instat-2 computer program (GraphPad Software Inc. version 2.04, San Diego, California, USA).

Results

Effect of vildagliptin on CsA-induced hepatotoxicity

As shown in Table 1, structural damage of the liver after treatment with CsA is evidenced by a significant elevation of ALT, AST, γGT, and LDH and a significant decrease of serum albumin levels as compared with the control. Vildagliptin treatment significantly improved all measured liver function parameters as compared with CsA.

Effect of vildagliptin on CsA-induced hepatic pathological changes

Livers from normal rats administered vehicle or vildagliptin showed normal hepatic structure. Upon treatment with CsA, liver changes were evident including enlarged hepatocytes, numerous

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vacuoles, inflammatory cell infiltrate, and dilated sinusoids with congestion. Rats that received both CsA and vildagliptin showed apparently less marked damage in liver tissue than that of liver tissue from CsA-treated rats (Figs. 1A–1D).

**Effect of vildagliptin on CsA-induced oxidative stress**

CsA caused significant elevation of hepatic oxidative stress as compared with the control. This is evident by high levels of TBARS, a marker of increased lipid peroxidation. Also, CsA treatment as compared with the control caused significant reduction of hepatic enzymatic antioxidant (SOD, CAT, and GPx) activity and non-enzymatic antioxidants (GSH) as shown in Table 2. Administration of vildagliptin with CsA to rats significantly attenuated CsA-induced oxidative stress. Vildagliptin caused significant reduction in TBARS, elevation in SOD, CAT, and GPx activities, and restored GSH levels as compared with the CsA-treated group (Table 2).

**Effect of vildagliptin on serum DPP-4 level**

Serum DPP-4 level was significantly increased by CsA administration indicating the high activity of DPP-4 enzyme compared with control non-treated rats. Upon vildagliptin concomitant treatment with CsA, vildagliptin was able to significantly reduce DPP-4 level to the normal value compared with the CsA-treated group (Table 1).

### Table 1. Effect of vildagliptin administration on serum biological changes in CsA-induced hepatotoxicity in rats.

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Vildagliptin</th>
<th>CsA</th>
<th>CsA+vildagliptin</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALT (U/L)</td>
<td>50.04±15.27</td>
<td>49.07±14.17</td>
<td>101.7±31.14†</td>
<td>63.7±18.24</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td>145.7±41.75</td>
<td>140.7±58.75</td>
<td>254±73.26†</td>
<td>177±5.06†</td>
</tr>
<tr>
<td>γGT (U/L)</td>
<td>22.34±6.25</td>
<td>24.27±6.99</td>
<td>42.71±11.65*</td>
<td>28.76±8.23‡</td>
</tr>
<tr>
<td>LDH (U/L)</td>
<td>667.70±19.12</td>
<td>678.67±19.43</td>
<td>1375.70±383.82*</td>
<td>785.7±225.43‡</td>
</tr>
<tr>
<td>Albumin (g/dL)</td>
<td>4.77±1.27</td>
<td>4.67±1.19</td>
<td>2.68±0.76</td>
<td>4.41±1.1§</td>
</tr>
<tr>
<td>DPP-4 (ng/mL)</td>
<td>34.05±10.32</td>
<td>29.09±8.43</td>
<td>109.06±33.66*</td>
<td>43.6±13.49‡</td>
</tr>
</tbody>
</table>

**Note:** Data are expressed as mean ± SD, n = 8. ALT, alanine transaminase; AST, aspartate aminotransferase; γGT, gamma-glutamyl transferase; LDH, lactate dehydrogenase; DPP-4, dipeptidyl peptidase-4; CsA, cyclosporine.

*p < 0.01, †p < 0.001 statistically different from mean value of control group using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

‡p < 0.05, †p < 0.01, §p < 0.001 statistically different from mean value of CsA-treated groups using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

**Fig. 1.** Effect of vildagliptin on pathological changes in rat liver induced by cyclosporine (CsA) (hematoxylin and eosin): (A) control group (magnification x100), (B) vildagliptin group (magnification x100), (C) CsA group (magnification x200), (D) CsA + vildagliptin (magnification x200). (A color version of this figure is available through the journal Web site at http://www.nrcresearchpress.com/doi/10.1139/cjpp-2015-0336.)
mulatelarge amounts of calcium (Fournier et al. 1987). These reduced calcium efflux, which allows the mitochondria to accu-
marked by a significant decreased in the level of Bax protein and significant decrease of the antiapoptotic protein Bcl2. Vildagliptin ameliorated CsA-induced apoptosis as apoptotic protein Bax with significant decrease of the antiapoptotic protein Bcl2.

Effect of vildagliptin on hepatic Bax and Bcl2 level

Increased apoptosis by CsA was evident by the increase in proapoptotic protein Bax with significant decrease of the antiapoptotic protein Bcl2. Vildagliptin ameliorated CsA-induced apoptosis as marked by a significant decreased in the level of Bax protein and increased that of Bcl2 (Table 4).

Discussion

The current study demonstrated the hepatotoxic effect of CsA when administered to rats for 4 weeks. Rats treated with CsA showed significant elevation of hepatic marker enzymes, AST, ALT, γGT, and LDH, and a significant decrease in albumin level as compared with the control. These changes indicate damage of the structural integrity of the liver, which was confirmed by the histopathological changes of the liver in CsA-treated rats that revealed enlarged hepatocytes, numerous vacuoles, inflammatory cell infiltrate, and dilated sinusoids with congestion as compared with the control. Vildagliptin ameliorated significantly the elevation of hepatic enzymes and the pathological changes of hepatic architecture induced by CsA.

ROS such as hydroxyl radicals, superoxide anions, and hydrogen peroxide are normally generated in the liver, but this is counteracted by a detoxification system of endogenous antioxidants such as GSH, SOD, and CAT that scavenge the produced ROS. The imbalance between intracellular prooxidants and antioxidants leads to an increase in both DNA damage and peroxidation of membrane lipids (Czaja 2007). In our study, liver damage evidenced by changes in biochemical markers (AST, ALT, γGT, LDH, and albumin) and tissue pathology of the liver in CsA-treated group was associated with increased oxidative stress. CsA-treated rats showed a significant elevation in hepatic TBARS and a decline in both enzymatic and non-enzymatic antioxidants, SOD, CAT, GPx, and GSH, as compared with the control concentrations. Oxidative damage has an important role in the pathogenesis of CsA-induced hepatotoxicity (El-Sokkary et al. 2005). Excess production of ROS by CsA is thought to be related to 2 important mechanisms, the first is due to its effect on mitochondria; CsA block the mitochondrial permeability transition pore (Nicoll et al. 1995) causing reduced calcium efflux, which allows the mitochondria to accumulate large amounts of calcium (Fournier et al. 1987). These events affect the mitochondrial respiratory chain and cause uncoupling of oxidative phosphorylation (Salducci et al. 1992) with subsequent increase in ROS production. The second mechanism is through CsA metabolism by cytochrome P-450 3A that can also generate ROS (Serino et al 1994). Previous studies found that CsA treatment inhibits the expression and activity of antioxidant en-
zymes like SOD, CAT, and GPx; and that exogenous antioxidants could inhibit the adverse actions of CsA (Lee 2010). To eliminate ROS from the cellular system, SOD and CAT function to remove superoxide radicals. GSH also plays an important role in the removal of ROS and protection of thiols in cell macromolecules. Depletion of GSH has been associated with CsA hepatotoxicity (Duruibe et al. 1989). The antioxidant effects of GSH are directly related to the key enzymes GPx and GSH reductase that help protect cells from free radical generated toxicity (Ghadermarzi and Moosavi-Movahedi 1996). Reduced hepatic GSH might account for the inhibition of GSH synthesis in addition, high levels of peroxides may explain the inhibition of CAT activity induced by CsA.

Table 2. Effect of vildagliptin administration on hepatic oxidative markers changes in CsA-induced hepatotoxicity in rats.

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Vildagliptin</th>
<th>CsA</th>
<th>CsA-Vildagliptin</th>
</tr>
</thead>
<tbody>
<tr>
<td>TBARS (nmol/g tissue)</td>
<td>39.7±5±1.65</td>
<td>41.2±5±1.66</td>
<td>11.2±6±3.39*</td>
<td>57.7±5±17†</td>
</tr>
<tr>
<td>SOD (U/g tissue)</td>
<td>21.87±5.71</td>
<td>20.75±5.69</td>
<td>9.75±2.77*</td>
<td>17.7±4.72‡</td>
</tr>
<tr>
<td>CAT (µmol/min/g tissue)</td>
<td>98.07±28.2</td>
<td>95.79±27.46</td>
<td>26.7±37.1*</td>
<td>75.7±621.6§</td>
</tr>
<tr>
<td>GPx (µmol/min/g tissue)</td>
<td>10.7±5.31</td>
<td>11.07±3.48</td>
<td>3.57±1.22*</td>
<td>8.7±2.57†</td>
</tr>
<tr>
<td>GSH (µmol/g tissue)</td>
<td>2.67±0.76</td>
<td>2.42±0.57</td>
<td>0.75±0.23‡</td>
<td>2.02±0.62‡</td>
</tr>
</tbody>
</table>

Table 3. Effect of vildagliptin administration on hepatic nuclear extracts of NF-κB in CsA-induced hepatotoxicity in rats.

<table>
<thead>
<tr>
<th></th>
<th>NF-κB activation (OD 450 nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.45±0.09</td>
</tr>
<tr>
<td>Vildagliptin</td>
<td>0.35±0.065</td>
</tr>
<tr>
<td>CsA</td>
<td>2.01±0.65*</td>
</tr>
<tr>
<td>CsA-vildagliptin</td>
<td>0.76±0.29††</td>
</tr>
</tbody>
</table>

Table 4. Effect of vildagliptin administration on hepatic protein level of Bax and Bcl2 in CsA-induced hepatotoxicity in rats.

<table>
<thead>
<tr>
<th></th>
<th>Bax ng/g tissue</th>
<th>Bcl2 ng/g tissue</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>2.05±0.45</td>
<td>5.57±1.44</td>
</tr>
<tr>
<td>Vildagliptin</td>
<td>1.95±0.57</td>
<td>5.07±1.16</td>
</tr>
<tr>
<td>CsA</td>
<td>8.07±2.18*</td>
<td>2.97±0.71*</td>
</tr>
<tr>
<td>CsA-vildagliptin</td>
<td>3.09±1.19†</td>
<td>4.52±0.99††</td>
</tr>
</tbody>
</table>

Note: Data are expressed as mean ± SD, n = 8. TBARS, thiobarbituric acid reactive substance; SOD, superoxide dismutase; GSH, glutathione; CAT, catalase; GPx, glutathione peroxidase; CsA, cyclosporine.

†p < 0.05, ‡p < 0.01 statistically different from mean value of control group using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

§p < 0.001 statistically different from mean value of CsA-treated groups using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

Note: Values are represented as mean ± SD, n = 8. NF-κB, nuclear factor kappa B; CsA, cyclosporine.

†p < 0.05, †p < 0.001 statistically different from mean value of control group using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

§p < 0.001 statistically different from mean value of CsA-treated groups using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

Note: Values are represented as mean ± SD, n = 8. CsA, cyclosporine.

†p < 0.001 statistically significant as compared with the control using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.

§p < 0.001 statistically different from mean value of CsA-treated group using one-way ANOVA test followed by Tukey–Kramer multiple comparison test.
The protective effect of vildagliptin in the present study may be partly explained by its ability to restore the prooxidant–antioxidant balance by enhancing hepatic radical scavenging activities, increase the antioxidant defense and prevent the formation of free radicals originating from peroxidation products and from metabolism of CsA. Vildagliptin maintained hepatic enzymatic (SOD, CAT, and GPx) and non-enzymatic antioxidants (GSH) in spite of CsA administration. Vildagliptin was found to ameliorate oxidative stress and pancreatic beta cell destruction in type 1 diabetic rats both by reducing levels of the stress marker TBARS, as well as by increasing activity of enzymatic antioxidants SOD and CAT (Avila Dde et al. 2013).

As a DPP-4 inhibitor, vildagliptin hinders degradation of bioactive incretins GLP-1 and GIP and amplifies their biological action. Vildagliptin treatment protects diabetic rats from the loss of renal vascular reactivity and the development of glomerulosclerosis perhaps secondary to a reduction in oxidative stress (Vavrinc et al. 2014). Liu et al. (2012) found that vildagliptin prevented the progression of diabetic nephropathy in rats by the enhancement of active GLP-1 level, which activated GLP-1R leading to antioxidative and antiapoptotic effects. Proapoptotic Bcl2 family members, such as Bax and Bak, promote formation of specific cytochrome c release channels in the mitochondrial outer membrane (Wei et al. 2001). Cytochrome c released after mitochondrial swelling initiates caspase-dependent apoptotic signaling (Jiang and Wang 2004).

Because oxidative stress is a well-known apoptosis mediator, many apoptosis inhibitors have an antioxidant activity or increase antioxidative defense mechanisms. By ameliorating oxidative stress induced by CsA, vildagliptin might have decreased CsA-induced apoptosis as evidenced by increased antiapoptotic protein Bcl2 and decreased that of proapoptotic protein Bax expression. Chronic administration of CsA in rats (2 weeks, 15 mg/kg/day) causes increased number of apoptotic cells in liver tissues when counted using the terminal deoxynucleotidyl transferase (TdT)-mediated dUTP nick end labeling (TUNEL) method (Gokus Erol et al. 2013). Shihab et al. (1999) proved that CsA-induced nephrotoxicity is associated with apoptosis caused by upregulation of the proapoptotic factors p53 and Bax and downregulation of the antiapoptotic factor Bcl2 in a rat chronic nephrotoxicity model. In pancreatic islets of diabetic mice, vildagliptin treatment was found to inhibit apoptosis (Wu et al. 2015). Tarantino et al. (2011b) deduced that there is a certain balance between apoptosis and antiapoptosis in determining the net effect on hepatocytes survival and hypothesized that serum Bcl2 levels reflect the steady state of this antiapoptotic protein. Serum Bcl2 increase is a likely response to the apoptotic process to improve survival of hepatocytes. If the response to the metabolic injury was good (increased serum levels of Bcl2) probably the more severe form would not develop. Tarantino et al. (2011a) illustrated similar levels of cytochrome c and gamma-glutamyl transferase in patients suffering from simple fatty liver or from the more severe non-alcoholic steato-hepatitis. A similar effect of vildagliptin in reducing oxidative stress and apoptosis was reported in cardiac ischemia/reperfusion injury (Apajai et al. 2014). Also, Chinda et al. (2014) reported a similar effect of vildagliptin on oxidative stress and apoptosis and explained it by the ability of vildagliptin to attenuate cardiac mitochondrial dysfunction through reducing ROS level and mitochondrial swelling.

NF-κB is a nuclear transcription factor found in the cytoplasm of liver cells where it plays an essential role in the regulation of inflammatory signaling pathways (Luedde and Schwabe 2011). A heterodimeric complex of p65/RelA and p50 is the most common form of NF-κB in mammalian cells. In the cytoplasm, NF-κB is bound to its inhibitory subunit IκB. As a result of oxidative stress, NF-κB is activated and phosphorylated from its inhibitory subunit and translocated to the nucleus of the hepatic cell where it binds to DNA. This causes upregulation of DNA transcription of many inflammatory genes like cytokines, chemokines and receptors of advanced glycation end products (Zhu and Fung 2000). In rat hepatocyte cultures, CsA induces transcriptional activation of NF-κB (Andrés et al. 2002). Assessment of hepatic nuclear extract in the current study showed significant increase NF-κB p65 DNA binding activity in CsA-treated rats as compared with control. Vildagliptin, however, reduced hepatic NF-κB activity caused by CsA administration. Miyagawa et al. (2013) demonstrated the anti-inflammatory benefits of vildagliptin in diabetic mice. They reported slight improvement of hepatic steatosis with a significant reduction of plasma inflammatory markers (i.e., C-reactive protein, IL-6, and TNF-α levels) as compared with control and attributed these effects to activation of GLP-1R on hepatocytes caused by enhanced levels of GLP-1 by vildagliptin.

To further elucidate the possible mechanism by which vildagliptin improve CsA-induced hepatotoxicity, DPP-4 level was estimated during CsA administration. The data showed that rats treated with CsA alone exhibited significant increase in DPP-4 level compared with untreated rats. However, vildagliptin concomitant with CsA administration significantly normalized DPP-4 levels. The liver is one of the organs that highly express DPP-4 (Mentzel et al. 1996). Hepatic DPP-4 is thought to be involved in the regulation of hepatic metabolism (Itou et al. 2013). Balaban et al. (2007) reported that serum DPP-4 activity and hepatic expression of DPP-4 are correlated with hepatic steatosis and non-alcoholic fatty liver disease grading. Non-alcoholic fatty liver disease is characterized by the accumulation of fat in the liver when it exceeds 5%–10% of its mass. In addition to leading to major histopathological alterations, it may be associated with elevated liver enzymes and abnormal liver function, ranging from steatosis to steatohepatitis, fibrosis, and cirrhosis (Tarantino and Finelli 2013). Moreover, DPP-4 deficient rats show lower levels of hepatic pro-inflammatory and pro-fibrotic cytokines and reduced hepatic steatosis compared to wild type rats (Balaban et al. 2007).

Also, Gaetaniello et al. (1998) reported that inhibition of DPP-4 in human hepatoma cells suppress tyrosine kinase, leading to anti-apoptotic effects. Thus, from the present study, the preventive effect of vildagliptin was attributed to DPP-4 inhibition that leads to anti-inflammatory and antiapoptotic as evidenced by significant effect of vildagliptin on the altered activity and expression of NF-κB, Bax and Bcl2, respectively.

In conclusion, the results of the present study demonstrate that administration of vildagliptin might have the potential to protect the liver against CsA toxic effects through reducing hepatic oxidative stress, apoptosis, and inflammation.

Conflict of interest

The authors declare that there is no conflict of interest associated with this work.

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