Thymoquinone suppressed Cyclosporine A-induced Nephrotoxicity in rats via antioxidant activation and inhibition of inflammatory and apoptotic signaling pathway

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ABSTRACT

Cyclosporine A (CsA) is a strong immunosuppressive drug, but its use is frequently accompanied by severe renal toxicity. The potential reno-protective effect of Thymoquinone (TQ) against CsA-induced nephrotoxicity in rats was assessed. Thirty adult white male albino rats were divided into three equal groups. Group I: (Normal control), received no drugs, Group II: (CsA treated), rats received oral dose of Cyclosporine A (25 mg/kg b. wt./day) for 21 days. Group III: (TQ protected + CsA), received Thymoquinone (10 mg/kg b. wt./day) orally 7 days before and during 21 days of CsA treatment. The obtained results showed a significant increase in the concentration of urea and creatinine in serum and L-MDA level in kidney tissue with marked decrease in renal catalase activity and GSH concentration in CsA treated rats. Moreover, a significant down-regulation in Bel-2 and up-regulation of NF-κB, PAI-1, Caspase-3 and p53 gene expressions levels were observed in kidney tissues of CsA treated rats. Also, various histopathological alterations were detected in kidneys of CsA treated rats. Meanwhile, TQ potentially improved renal function and oxidative alterations related to CsA near its normal ranges. Interestingly, histopathological findings supported that TQ markedly attenuates harmful effects that CsA induced and protected kidney. Our research could conclude that, TQ has an ameliorating role as potent antioxidant and anti-apoptotic agent via inhibition of inflammatory (NF-κB, PAI-1) and apoptotic (Caspase-3, p53) signaling pathway in modulation of CsA-induced nephrotoxicity.

1. INTRODUCTION

Cyclosporins are immunosuppressant drugs isolated from Tolypocladium inflatum (Cragg and Newman, 2013). Cyclosporine A nephrotoxicity is related to its chronic use in numerous autoimmune diseases and in organ transplantation especially kidney transplantation (Raesisi et al., 2016). Oxidative stress has an important role in Cyclosporine A nephrotoxicity. Cyclosporine A-induced nephrotoxicity results from the endoplasmic reticulum stress activation, the increased production of the mitochondrial reactive oxygen species (ROS) which modifies the redox balance enhancing lipid peroxidation ends with nephrotoxicity (Wu et al., 2018). Thymoquinone, the main bioactive constituent isolated from black seed or Nigella sativa volatile oil with many pharmacological properties including anti-inflammatory, antioxidant, immunomodulatory and antineoplastic effects (Elisherbiny et al., 2017). Thymoquinone is characterized by high lipophilicity giving it easy accessibility to subcellular compartments and owing to its benzoquinone structure.

It has strong antioxidant potential with high ability to scavenge free radicals (Darakhshan et al., 2015). This study aimed to evaluate the possible protective effect of Thymoquinone against the nephrotoxicity induced by Cyclosporine A in adult white male albino rats through investigation of renal function, oxidative stress and antioxidant biomarkers, pro-inflammatory mediators, apoptotic and anti-apoptotic gene expression markers in kidney tissues in addition to evaluation of histopathological alterations of kidney.

2. MATERIAL AND METHODS

2.1. Experimental animals:
Thirty-white male albino rats, 4-5 weeks old with average body weight 140-160 g were used in this study. Rats were fed on constant ration and fresh, clean drinking water was supplied ad-libitum. All rats were acclimatized for two weeks prior to the beginning of study. The experimental protocols were approved by the Animal Care and Use committee at Benha University and are in accordance with the National Institute of Health Guide for the Care and Use of Laboratory Animals.

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2.2. Chemicals and antioxidant agent:
1. Cyclosporine (CsA): CsA presents in the form of soft gelatin capsules containing 50 mg cyclosporine under traditional name (Sandimmune® Neoral®) obtained from (Novartis Pharma AG, Basel, Switzerland). CsA was freshly dissolved in propylene glycol. Nephrotoxicity was induced in rats after oral treatment with CsA (25mg/kg b.wt/day) for 21 successive days according to Chia et al. (2012).
2. Thymoquinone (2-isopropyl-5-methyl-1,4-benzoquinone), was purchased from Sigma-Aldrich Chemical Co. (St. Louis, MO, USA). The calculated dose of TQ was 10 mg/kg b. wt/day according to Awad et al. (2011). Thymoquinone was dissolved in absolute ethanol then diluted to the appropriate concentration by physiological saline according to Nagi and Mansour (2000).

2.3. Experimental design:
Rats were randomly divided into three groups, 10 rats each, kept in individual cages and classified as follow:
Group I: (Normal control group): Rats fed only with ordinary diet without any treatment during the experiment period.
Group II: (Cyclosporine A nephrotoxic group): Rats received oral administration of CsA (25 mg/kg b. wt/day) for 21 days.
Group III: (Thymoquinone protected group): Rats orally received Thymoquinone (10 mg/kg b. wt./day), 7 days prior to CsA administration and concurrently during CsA treatment for 21 days later.

2.4. Sampling:
2.4.1. Blood samples
Blood samples were taken via veins puncture of the medial canthus of the eye, allowed to clot, then centrifuged for 15 minutes at 3,000 rpm. Sera were separated in dry sterile tubes by automatic pipette, and then stored at -20 °C in a freezer until use for determination of urea and creatinine concentration.

2.4.2. Tissue samples:
After blood collection rats were sacrificed by decapitation according to Animal Ethics Committees and abdomen was opened, then kidneys were collected.

2.4.3. For biochemical analysis:
Briefly, one gram of kidney tissues were cut and minced into small pieces, homogenized with a glass homogenizer in 9 volume of ice-cold 0.05 mM potassium phosphate buffer (pH7.4) to make 10% homogenates, then centrifuged for 15 minutes at 6000 RPM at 4 °C. The supernatant used directly for determination of catalase (CAT) activity, reduced glutathione (GSH) and L-malondialdehyde (L-MDA) concentrations.

2.4.4. For molecular analysis:
About 0.5 g of kidney tissue was put in Eppendorf tubes, kept immediately in liquid nitrogen and stored at -80 °C till RNA extraction for determination of caspase-3, tumor suppressor protein p53, B cell lymphoma-2 (Bcl-2), nuclear factor kappa B (NF-κB) and Plasmonen activator inhibitor-1 (PAI-1) gene expressions.

2.4.5. For histopathological examination:
Small tissue specimens were collected from the kidneys and immediately fixed in 10% neutral buffered formalin solution then subjected to histopathological examination according to the technique described by Bancroft and Gamble (2008).

2.5. Analysis:
2.5.1. Biochemical analysis:
Serum urea and creatinine were measured according to methods described by Kaplan and Kohn (1992) and Jaffe, (1986), respectively. Also, kidney tissue catalase activity, GSH and L-MDA concentrations were measured according to methods described by Weydert and Cullen (2010), Moron et al. (1979) and Lahouel et al. (2004), respectively.

2.5.2. Molecular analysis:
The mRNA expression contents of Caspase-3, p53, Bcl-2, NF-κB and PAI-1 were determined using real time quantitative polymerase chain reaction analysis (real-time qPCR) in rat kidney. β-actin was used as load control. Total RNA was isolated from the kidney using High Kit for isolation of pure RNA (Thermo Scientific, Fermentas, #K0731) RNA Extraction kit according to manufacturer’s instructions. With each cDNA, sample was reverse transcribed using RevertAid TM First Strand CDNA synthesis kit (#EP0451, Thermo Scientific, Fermentas, USA). Then, real-time quantitative PCR amplification was performed on Faststart Universal SYBR Green Master (Roche, GÉR). The target gene was normalized with β-actin by the 2^-ΔΔCt method (Livak and Schmittgen, 2001). Primers sequence for real time PCR is presented in table (1)

Table 1 Forward and reverse primers sequence for real time PCR.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Caspase-3</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
</tr>
<tr>
<td>p53</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
</tr>
<tr>
<td>Bcl-2</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
</tr>
<tr>
<td>NF-κB</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
</tr>
<tr>
<td>PAI-1</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
</tr>
<tr>
<td>β-actin</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
<td>ΔΔCt+1/1AGGACAGGAAGCAATAGAGA</td>
</tr>
</tbody>
</table>

2.6. Statistical analysis:
Results were expressed as mean ± SE (Standard Error) using SPSS software (Version 13.0, 2009). Data were analyzed using one-way ANOVA followed by Duncan’s test. Values were statistically taken into account significantly at p < 0.05.

3. RESULTS
Data were presented in table (2) showed that, serum urea and creatinine levels were significantly elevated in CsA treated rats. However, in TQ protected group urea and creatinine values were significantly reduced compared with CsA nephrotoxic group.

Table 2 Protective effect of Thymoquinone administration on serum urea and creatinine concentrations in Cyclosporine A-induced nephrotoxicity in rats.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>Urea (mg/dl)</th>
<th>Creatinine (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group I: Normal control</td>
<td>35.44 ± 0.13</td>
<td>0.92 ± 0.01</td>
</tr>
<tr>
<td>Group II: Cyclosporine A</td>
<td>59.07 ± 0.23</td>
<td>1.58 ± 0.06</td>
</tr>
<tr>
<td>Group III: Thymoquinone + Cyclosporine A</td>
<td>38.70 ± 0.36</td>
<td>1.02 ± 0.01</td>
</tr>
</tbody>
</table>

Data are presented as (Mean ± S.E): S.E = Standard error. Mean values with different superscript letters in the same column are significantly different at (P ≤0.05).

The obtained results in table (3) showed a significant increase in L-MDA level while GSH content and CAT activity were markedly decreased in kidney tissue of CsA.
treated rats compared with normal control group. However, TQ protected group showed significant decrease in L-MDA level with marked increase in GSH level and CAT activity comparing with CsA nephrotoxic group.

Table 3 Protective effect of Thymoquinone administration on kidney tissue L-MDA, GSH concentrations and CAT activity in Cyclosporine A-induced nephrotoxicity in rats.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>L-MDA (μmol/g. tissue)</th>
<th>GSH (μmol/g. tissue)</th>
<th>CAT (U/g. tissue)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal control</td>
<td>3.34 ± 0.09</td>
<td>34.73 ± 0.32</td>
<td>9.75 ±0.15</td>
</tr>
<tr>
<td>Group II: Cyclosporine A</td>
<td>17.03 ± 0.35</td>
<td>10.77 ±0.28</td>
<td>2.41 ±0.30</td>
</tr>
<tr>
<td>Group III: Thymoquinone + Cyclosporine A</td>
<td>8.45 ± 0.23</td>
<td>24.10 ±0.55</td>
<td>8.37 ±0.15</td>
</tr>
</tbody>
</table>

Data are presented as (Mean ± SEM). SEM = Standard error mean. Mean values with different superscript letters in the same column are significantly different at (P≤0.05).

The presented data in table (4) showed a significant up-regulation in kidney tissue NF-κB and PAI-1 gene expressions in CsA treated rats comparing with normal control group. However, TQ protection significantly down-regulate these expressions compared with CsA nephrotoxic group.

Table 4 Protective effect of Thymoquinone administration on kidney tissue NF-κB and PAI-1 gene expression levels in Cyclosporine A-induced nephrotoxicity in rats.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>Nuclear factor kappa B (NF-κB)</th>
<th>Plasminogen activator inhibitor-1 (PAI-1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group I: Normal control</td>
<td>1.00 ± 0.07</td>
<td>1.00 ± 0.09</td>
</tr>
<tr>
<td>Group II: Cyclosporine A</td>
<td>4.35 ± 0.21</td>
<td>3.23 ± 0.33</td>
</tr>
<tr>
<td>Group III: Thymoquinone + Cyclosporine A</td>
<td>2.01 ± 0.09</td>
<td>1.00 ± 0.07</td>
</tr>
</tbody>
</table>

Data are presented as (Mean ± SEM). SEM = Standard error mean. Mean values with different superscript letters in the same column are significantly different at (P≤0.05).

The results presented in table (5) revealed significant up-regulation in kidney tissue Caspase-3, p53 and down-regulation in Bcl-2 gene expressions in CsA treated rats comparing with normal control group. Meanwhile, TQ protection caused significant down-regulation in kidney tissue Caspase-3, p53 and up-regulation in Bcl-2 gene expressions when compared with CsA nephrotoxic group.

Table 5 Protective effect of Thymoquinone administration on kidney tissue apoptotic (Caspase-3, p53) and anti-apoptotic (Bcl-2) gene expression levels in Cyclosporine A-induced nephrotoxicity in rats.

<table>
<thead>
<tr>
<th>Animal Groups</th>
<th>Caspase-3</th>
<th>Tumor suppressor protein p53</th>
<th>Bcl-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1.00</td>
<td>0.06</td>
<td>1.00</td>
</tr>
<tr>
<td>Group II: Cyclosporine A</td>
<td>6.28</td>
<td>0.15</td>
<td>0.01</td>
</tr>
<tr>
<td>Group III: Thymoquinone + Cyclosporine A</td>
<td>2.01</td>
<td>0.09</td>
<td>0.01</td>
</tr>
</tbody>
</table>

Data are presented as (Mean ± SEM). SEM = Standard error mean. Mean values with different superscript letters in the same column are significantly different at (P≤0.05).

Histopathological examination:
Microscopical examination of kidney tissues obtained from normal control rats (Group I) revealed normal histological structure with normal glomeruli and proximal and distal convoluted tubules. Microscopical examination of renal tissues obtained from CsA treated rats with oral dose of (25 mg/kg b.wt/day) for successive 21 days revealed various histopathological alterations including congestion of inter-tubular blood capillaries and renal blood vessels (Figure 1a) with perivascular mononuclear leukocytic cellular infiltrations mainly lymphocytes and macrophages in renal cortex (Figure 1b). Vacuolation of the endothelial cell lining the glomerular tuft (Figure 1c) as well as hyper-segmentation of the glomerular tuft with adhesions between glomerular tufts and Bowman’s capsule as well as thickening in the wall of Bowman’s capsules were recorded (Figure 1d). Necrosis and disintegration of the glomerular tuft with shrinkage of the glomerular tuft was seen in few cases. The lining epithelium in renal cortex exhibited extensive degenerative changes such as vacuolar and hydropic degeneration characterized by swollen pale vacuolated cytoplasm in association. Necrosis of the lining epithelium of some renal tubules were also observed (Figure 1e). Meanwhile, the microscopical of the kidneys obtained from rats in (Group III), the protected group with thymoquinone; revealed moderate improvement in the renal tissue histology as congestion of inter-tubular capillaries and renal blood vessels with few perivascular mononuclear leukocytic cellular infiltration was detected in few cases. Proliferation of the endothelial cell lining the glomerular tuft resulting in its adhesions with Bowman’s capsule and absence of Bowman’s space in association with mild degenerative changes in the lining epithelium of some proximal and distal convoluted tubules as cloudy swelling was also observed in renal cortex (Figure 1f). Moreover, few numbers of renal tubules were ecstatic and lined by attenuated epithelium (Figure 1g).

4. DISCUSSION

Nephrotoxicity could be the major problem in transplant medicine as a side effect caused by Cyclosporine A (CsA) as an immunosuppressive drug. Oxidative stress has a crucial role nephrotoxicity induced by Cyclosporine A (Raesi et al., 2016). Apoptosis increased oxidative stress, and mitochondrial dysfunction were reported in CsA nephrotoxicity (Hausenloy et al., 2012). Renal failure was the end stage results from nephrotoxicity associated with long term administration of Cyclosporine A (Caires et al., 2018).

In the present study, cyclosporine A treated rats revealed significant increases in serum concentration of urea and creatinine. Abdel-Wahab, (2015) demonstrated that CsA administration for 21 days impaired the proper kidney function and induced the renal oxidant/antioxidant homeostasis disturbance. Urea and creatinine levels were increased, while glomerular filtration rate (GFR) was decreased on chronic CsA administration. This was related to reactive oxygen species (ROS) formation in kidney revealed by increased lipid peroxidation (LPO), decreased GSH in addition to increased inducible nitric oxide synthase (iNOS) level establishing the oxidative stress role in pathogenesis of renal damage, vascular injury, and endothelium dysfunction induced by Cyclosporine A (Hewed and Mostafa, 2016). Conversely, administration of Thymoquinone to CsA treated rats revealed significant decreases in serum levels of urea and creatinine. Similar data were recorded by Zidan et al. (2018) who revealed the dramatic fall in creatinine and BUN after thymoquinone administration. This effect was related to its antioxidant effect which reduced the nephrotoxicity, due to the free radical scavenging properties of thymoquinone against ROS involved in vasoconstriction and the impairment in glomerular filtration rate (Abdel-Wahab, 2015).
Figure 1: H&E stained sections of kidney tissue from (a-e) treated rats with Cyclosporine-A (Group II) and (f-g) from rats treated with (Thymoquinone + Cyclosporine-A) (Group III) showing (a) marked congestion of the inter-tubular blood capillary (C, x200), (b) perivascular mononuclear aggregation mainly lymphocytes and macrophages (arrow), with necrosis of glomerular tuft (asterisk, x400), (c) swelling and vacuolization of endothelial cells of glomerular tuft (arrow, x400), (d) hyper-segmentation of glomerular tuft (asterisk) with accumulation of eosinophilic proteinaceous substance on Bowman’s capsule (arrow, x400), (e) vacuolation (arrow) and entire necrosis (zigzag arrow) of the lining epithelium of renal tubules with any layer of the blood vessel (D, x400), (f) hyper-cellularity of glomerular tuft to the extent of its adhesion with the parietal layer of Bowman’s capsule with cloudy swelling of the epithelial cell lining of the renal tubules (x200), (g) cystic dilatation of some renal tubules (arrow, x200).
A significant increase in kidney tissue L-MDA concentration with marked decreases in GSH level and CAT activity were observed in CsA treated rats. The increased lipid peroxidation increases peroxo and hydroxyl radicals resulting in oxidative damage (Mustafa et al., 2018). Our results agreed with Hussein et al. (2014) who demonstrated that a significant increase in renal L-MDA level resulted by CsA treatment in rats, suggesting the role of oxygen free radicals involved in renal injury. ROS production by CsA may be due to the drug action as an uncoupler and inhibitor of the mitochondrial electron transport system and metabolism of CsA by cytochrome P450 3A (Ateyya, 2015). Endogenous antioxidant defense system decrease results from ROS production. Excessive ROS decreased the renal antioxidant capacity and so the kidney became more susceptible to oxidative damage. Significant decreases in kidney GSH content and CAT activity were reported in CsA treated rats (Abdel-Wahab, 2015). It is possible that depletion of the production of NADPH which is required for activation of catalase from its inactive form during CsA-Treatment could decrease the catalase activity (Hussein et al., 2014).

Meanwhile, protection with Thymoquinone resulted in significant decrease in the L-MDA level in kidney tissue and markedly increases in GSH content and CAT activity. Zidan et al. (2018) showed that thymoquinone treatment revealed significant increases in GSH level and CAT activity and significant decrease in L-MDA concentration. Protective effects of Thymoquinone was explained by its antioxidant property related to the free radicals scavenging ability, the protection of cellular membranes against lipid peroxide attack, maintaining the intracellular GSH levels and increasing of antioxidant genes expressions in tissues (Farag et al., 2015). Our study showed that CsA treatment induced significant up-regulation of kidney tissue NF-κB and PAI-1 gene expression levels. CsA administration reduced the renal content of GSH, CAT and SOD activities resulted in oxidative damage expressed by increase in lipid peroxidation, iNOS and NF-kB expression (Balah, 2014). The NF-kB transcription factor has a role involved in the transcription of inflammatory iNOS and other inflammatory genes in response to oxidative stress (Aktan, 2004). Plasminogen activator inhibitor-1 (PAI-1) is a pro-fibrotic cytokine considered as the major inhibitor of fibrinolysis (Ghosh and Vaughan, 2012). PAI-1 increased expression in vivo inhibits fibrinolysis and leading to the pathological fibrin deposition and tissue damage (Aso, 2007). CsA nephrotoxicity mechanism involves endothelial function impairment, vasoconstrictors and vasodilators mediators unbalance, activation of renin-angiotensin–aldosterone system and increased cell proliferation. These mediators increased transforming growth factor-β1 (TGF-β1) expression and plasminogen activator inhibitor-1 (PAI-1) which led to disruption of tissue architecture by increasing extracellular matrix synthesis and decreasing its degradation, resulting in fibrosis of tubule-interstitial and arteriolopathy (Young et al., 1995).

On other hand, Thymoquinone administration significantly down-regulated NF-kB and PAI-1 gene expressions. Thymoquinone inhibits ROS which induces NF-kB and leads to inhibition of apoptosis and pro-inflammatory cytokines (El-Mahdy et al., 2014). Increasing glutathione levels assumed to be an important mechanism implicated in thymoquinone protective effect against inflammatory disorders (Majdalawiich and Fayyad, 2015). Oskouei et al., (2018) illustrated that pre- and post-administration of thymoquinone resulted in tubular renal cells and hemodynamic functional parameters improvement in and impairment of some pro-inflammatory cytokines and profibrotic gene expressions, including tumor necrosis factor alpha (TNF-α), transforming growth factor beta 1 (TGF-β1) and the type-1 inhibitor of plasminogen activator (PAI).

Cyclosporine A treated rats showed significant up-regulation in kidney tissue gene expressions of apoptotic (caspase-3, p53) and down-regulation of antiapoptotic (Bcl-2). Renal cell apoptosis and kidney dysfunction were important features of chronic CsA nephrotoxicity (Xiao et al., 2013). The significant up-regulation in kidney tissue p53 gene expression in CsA treated rats came in accordance with Moon and Kim, (2019). Activation of p53 induces a series of programs, including cell-arrest, cellular senescence and apoptosis (Hsin et al., 2006). Cell apoptosis signaling pathway is activated by the mitochondrial and endoplasmatic reticulum (ER) stress. Besides, mitochondrial injury induced mitochondria release of cytochrome C, resulting in mitochondrial transmembrane potential loss (Muthuraman et al., 2011) and activation of caspase family of proteases; caspase-3 and caspase-9 (Liu et al., 2019). The anti-apoptotic Bcl-2 protein is able to inhibition of Bax-induced apoptosis (Rossé et al., 1998). The observed down-regulation in Bcl-2 protein expression came in accordance with (Ateyya, 2015). CsA down-regulated Bcl-2 and Bcl-XL resulted in Bax translocation to the mitochondria and impaired Bcl-2 and Bax balance in renal tubular cells.

Thymoquinone showed its anti-apoptotic effect by down-regulating kidney tissue Caspase-3 and p53 and up-regulating Bcl-2 gene expression levels in protected group. Rahmani et al. (2019) indicated that thymoquinone attenuated liver injury induced by CCl4 in mice by p53 modulation. Moreover, Mahmoud et al., (2014) indicated that administration of thymoquinone induced the protective effects on the kidney by inhibiting the apoptotic cascade through the up-regulation of Bcl-2 expression, suppression of caspase-3 and caspase-9 activation and reduced the cleavage of Poly [ADP-ribose] polymerase 1 (PARP-1).

5. CONCLUSION

It could be concluded that, thymoquinone has a renoprotective effect by enhancing antioxidant defense system and attenuates CsA induced nephotoxicity and oxidative stress. Moreover, thymoquinone has a strong anti-inflammatory effect through modulation of NF-kB, PAI-1 gene expressions, and anti-apoptotic effect by inhibiting caspase-3, p53 and activating Bcl-2 signaling pathways.

6. REFERENCES


