Effect of Thymoquinone and Omega-3 on Intestinal Ischemia/Reperfusion Induced Hepatic Dysfunction in Rats


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Abstract: Intestinal Ischemia-Reperfusion (I/R) is a complex phenomenon causing local and remote tissue destruction and multiple-organ dysfunction. The present study investigates the effect of thymoquinone and omega-3 on intestinal I/R-induced hepatic dysfunction. Sixty four Wistar albino rats were randomly allocated into four experimental groups: sham control, intestinal I/R control, thymoquinone (10 mg/kg) and omega-3 (300 mg/kg) pre-treated groups respectively. Intestinal I/R model were established by clamping the superior mesenteric artery for 30 min followed by 60 min reperfusion. Serum levels of aspartate Amino Transferase (AST), Alanine Aminotransferase (ALT) and Alkaline Phosphatase (ALP) were measured. Hepatic tissue contents of Malondialdehyde (MDA), reduced Glutathione (GSH), Myeloperoxidase (MPO) and Tumor Necrosis Factor alpha (TNF-α) and Superoxide Dismutase (SOD) activity were measured. Apoptosis in hepatic tissue cells was determined by immunochemical analysis of caspase-3. Hepatic histopathological examination was carried out. Intestinal I/R elevated serum AST, ALT, ALP levels. In-addition, hepatic tissue SOD activity and GSH content were decreased, while MDA, MPO and TNF-α contents were increased. Pre-treatment with thymoquinone or omega-3 corrected the histopathological changes and serum AST, ALT, ALP levels as well as hepatic tissue activity of SOD and GSH contents. In-addition, hepatic tissue contents of MDA and MPO were decreased. Furthermore, immunohistochemical examination showed remarkable activation of caspase-3 activity in hepatic tissue after intestinal I/R, which were corrected by thymoquinone or omega-3 pre-treatment. The protective potential of thymoquinone and omega-3 on hepatic dysfunction could be attributed to the known antioxidant, antiapoptotic and anti-inflammatory effects of test drugs.

Keywords: Antiapoptotic, anti-inflammatory, antioxidant, intestinal ischemia/reperfusion, MPO, omega-3, thymoquinone

INTRODUCTION

Ischemia and Reperfusion (I/R) is a pathological condition characterized by an initial restriction of blood supply to an organ followed by the subsequent restoration of perfusion and concomitant re-oxygenation (Yellon and Hausenloy, 2007). The absence of oxygen and nutrients from blood during the ischemic period and reperfusion results in inflammation and oxidative damage through the induction of oxidative stress. Surprisingly, restoration of blood flow and re-oxygenation is frequently associated with tissue injury and a profound inflammatory response called reperfusion injury (Takizawa et al., 2011).

Intestinal I/R leads to systemic inflammation and multiple organ failure in clinical and laboratory settings (Santora et al., 2010). Multiple organ dysfunction syndrome is mediated by complex mechanisms in which interactions between activated leukocytes and endothelial cells play a central role (Olanders et al., 2002).

I/R injury also including production of Reactive Oxygen Species (ROS), inflammatory cell infiltration and cytokine production (Paterno and Longo, 2008). Increased oxidative stress and inhibition of cellular antioxidant defence mechanisms are associated with apoptotic cell death (Hayes and McLellan, 1999). Malondialdehyde (MDA) is one of the final products of lipid peroxidation. MDA can be found in both tissue and blood and its concentration is directly proportional to the cell damage caused by free radicals (Halliwell and Gutteridge, 1990). Antioxidant enzymes, including reduced Glutathione (GSH) and Superoxide Dismutase (SOD), protect tissues from reperfusion injury by destroying ROS (Deshmukh et al., 1997).

One of the main orchestrators of I/R injury is the proinflammatory cytokine Tumor Necrosis Factor (TNF-α) (Pascher and Klupp, 2005). TNF-α promotes leukocyte infiltration, induces cell death and activates an inflammatory cascade that results in further production of cytokines and chemokines (Ciesielski and Modzelewski, 1995; Esposito et al., 2007).
Pharmacologic modulation of TNF-α production is a promising strategy in the prevention of I/R injury (Camara-Lemarroy et al., 2010). I/R activate various programs of cell death, which can be categorized as necrosis, apoptosis or autophagy-associated cell death.

Apoptosis induced by I/R involves an orchestrated caspases-signaling cascade that induces a self-contained program of cell death, characterized by the shrinkage of the cell and its nucleus (Hotchkiss et al., 2009).

Thymoquinone (TQ), a component of Nigella sativa is known to have broad anti-inflammatory activities and attenuates allergic inflammation. TQ attenuates the pro-inflammatory response mainly by modulating nuclear transactivation of NF-kappa B and TNF-α production (El Gazzar et al., 2007).

Omega-3 fatty acid is one of the major constituents of fish oil. It has been shown that omega-3 has beneficial effects in multiple disease states that involve an inflammatory process. The effects of omega-3 were reported to be through modulation of inflammatory mediators such as TNF-α as well as inhibition of apoptotic marker; caspase-3 (El-Ansary et al., 2011a). The present study is constructed in order to explore the effects of TQ or omega-3 on liver dysfunction induced by intestinal I/R in rats.

MATERIALS AND METHODS

Animals: Adult male Wistar albino rats, weighing 150-180 g. Animals purchased from the National Institute of Ophthalmology, cairo, Egypt. Animals were kept in the animal center under appropriate conditions of temperature, humidity and light. The study was carried out according to the approval of ethics committee for animal experimentation at faculty of pharmacy, Cairo University. Animals were fed standard pellet chow (El-Nasr Chemical Co., Cairo, Egypt) and were allowed water ad libitum.

Drugs: Thymoquinone (Sigma-Aldrich Co, USA) and Omega-3 (Metagenics, INC., Norway), were used in this study.

Experimental design: Rats were randomly allocated into 4 groups (16 rats each) as follows:

Group 1: Rats exposed to sham operation and served as sham control.

Group 2: Rats received distilled water orally for 14 days and then exposed to intestinal I/R operation and served as I/R control.

Group 3: Rats received thymoquinone (10 mg/kg) orally (Fouda et al., 2008; Padhye et al., 2008) for 14 days and then exposed to intestinal I/R operation.

Group 4: Rats received omega-3 (300 mg/kg) orally (Kavakli et al., 2007) for 14 days and then exposed to intestinal I/R operation and served as reference standard.

Induction of intestinal I/R: I/R rat model was established according to the method described by (Cheng et al., 2013). Clamping superior mesenteric artery for 30 min followed by reperfusion for 60 min. At the end of the experiment, blood samples were collected via the retro-orbital plexus under light anesthesia (di-ethyl ether), serum was separated, for biochemical parameters. Animals were sacrificed by cervical dislocation, liver was rapidly isolated and washed with ice cold saline. Part of the liver was homogenized in phosphate buffer saline. The other part of the liver were embedded in 10% formalin for histopathological examination and immunohistochemical staining.

Biochemical assay: The serum levels of Aspartate Aminotransferase (AST), Alanine Aminotransferase (ALT) and Alkaline Phosphatase (ALP) were measured using kits supplied by Biodiagnostic, Egypt. MDA and GSH content as well as SOD activity in hepatic tissue were determined spectrophotometrically using commercial kits supplied by Biodiagnostic, Egypt.

Hepatic tissue content of TNF-α and Myeloperoxidase (MPO) were measured by ELISA (AssayPro and Hycult Biotechnology, USA respectively).

Histopathological examination: Liver specimens were fixed in 10% formalin and embedded in paraffin. Tissue sections (4 μm) were stained with Hematoxylin and Eosin (H&E). Immunohistochemistry for caspase-3 was performed in sections prepared from formalin-fixed, paraffin-embedded tissue using the avidin-biotin immunodetection complex method according to the manufacture’s instruction (Labvision, USA). Interpretation of results was done semiquantitatively by evaluating both intensity and distribution of positive cells. Hepatocyte cells showing cytoplasmic staining for caspase-3, The intensity of caspase-3 immunostaining was assessed as follows: none = 0, mild = 1, moderate = 2 and strong = 3. The immunohistochemical Histological score (H-score) was then calculated by multiplying the intensity by the percentage of liver cells showing positive staining for caspase-3, creating a range of possible scores of 0-300.

Statistical analysis: Results are expressed as means±Standard Error of Mean (S.E.M.). Comparisons between different groups were carried out by one-way Analysis of Variance (ANOVA) followed by the Tukey-Kramer test. The level of significance was set at p<0.05. Graphpad software instat (version 2) was used to carry out statistical analysis.
RESULTS

Effect of TQ or omega-3 on ALT, AST and ALP:
Normal serum level of AST, ALT and ALP were 8.785, 18.75 U/mL and 53.142 IU/L, respectively. I/R resulted in significant increase to 84.125, 69.013 U/mL and 126.511 IU/L, respectively. Pre-treatment of TQ resulted in significant decrease to 17.750, 22.375 U/mL and 60.233 IU/L respectively, while treatment of omega-3 resulted in significant decrease to 28.375, 23.250 U/mL and 66.250 IU/L, respectively (Table 1).

Effect of TQ or omega-3 on hepatic MDA, GSH and SOD:
Normal hepatic tissue content of MDA, GSH and SOD activity were 109.330 nmol/g, 457.221 mg/g.tissue and 93.147 U/mg.tissue, respectively. I/R resulted in significant increase of MDA content to 171.101 nmol/g.tissue and significant decrease of GSH content and SOD activity to 285.110 mg/g.tissue and 37.753 U/mg.tissue respectively. Pre-treatment of TQ or omega-3 significantly decreased MDA content to 114.207 and 111.233 nmol/g.tissue, significantly increased GSH content to 438.366 and 447.250 mg/g.tissue and increased SOD activity to 79.851 and 87.897 U/mg.tissue, respectively (Table 2).

Table 1: Effect of TQ or omega-3 on the serum level of AST, ALT and ALP in intestinal I/R rats

<table>
<thead>
<tr>
<th>Groups</th>
<th>AST unit/mL</th>
<th>ALT unit/mL</th>
<th>ALP IU/L</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham control</td>
<td>8.785±1.086</td>
<td>18.75±1.750</td>
<td>53.142±2.539</td>
</tr>
<tr>
<td>I/R control</td>
<td>84.125±7.691</td>
<td>69.013±5.868</td>
<td>126.511±6.582</td>
</tr>
<tr>
<td>Thymoquinone</td>
<td>17.750±2.104</td>
<td>22.375±2.203</td>
<td>60.233±3.454</td>
</tr>
<tr>
<td>Omega-3</td>
<td>28.375±2.620</td>
<td>23.250±2.624</td>
<td>66.250±3.612</td>
</tr>
</tbody>
</table>

Thymoquinone (10 mg/kg; P.O.) or omega-3 (300 mg/kg; P.O.) was administered for 14 days before I/R; Data are presented as the mean±S.E.M.; a: Significantly different from sham control at p<0.05; b: Significantly different from I/R control at p<0.05; Statistical analysis was carried out by one way Analysis of Variance (ANOVA) followed by Tukey-kramer test

Table 2: Effect of TQ or omega-3 on hepatic MDA and GSH contents and SOD activity in intestinal I/R in rats

<table>
<thead>
<tr>
<th>Groups</th>
<th>MDA nmol/g.tissue</th>
<th>GSH mg/g.tissue</th>
<th>SOD U/mg.tissue</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham control</td>
<td>109.330±8.750</td>
<td>457.221±19.711</td>
<td>93.147±8.250</td>
</tr>
<tr>
<td>I/R control</td>
<td>171.101±15.751</td>
<td>285.110±23.696</td>
<td>37.753±3.110</td>
</tr>
<tr>
<td>Thymoquinone</td>
<td>114.207±8.368</td>
<td>438.366±33.854</td>
<td>79.851±6.749</td>
</tr>
<tr>
<td>Omega-3</td>
<td>111.233±10.258</td>
<td>447.250±38.542</td>
<td>87.897±8.102</td>
</tr>
</tbody>
</table>

Thymoquinone (10 mg/kg; P.O.) or omega-3 (300 mg/kg; P.O.) was administered for 14 days before I/R; Data are presented as the mean±S.E.M.; a: Significantly different from sham control at p<0.05; b: Significantly different from I/R control at p<0.05; Statistical analysis was carried out by one way Analysis of Variance (ANOVA) followed by Tukey-kramer test

Fig. 1: Effect of TQ or omega-3 on hepatic content of TNF-α in intestinal I/R in rats
Thymoquinone (10 mg/kg; P.O.) as well as omega-3 (300 mg/kg; P.O.) was administered for 14 days before I/R; a: significantly different from normal control at p<0.05; b: Significantly different from I/R control at p<0.05; Statistical analysis was carried out by one way Analysis of Variance (ANOVA) followed by Tukey-kramer test for multiple comparisons
Effect of TQ or omega-3 on hepatic TNF-α: The normal value of hepatic tissue content of TNF-α was 72.380 pg/mg.tissue I/R resulted in significant increase to 211.649 pg/mg.tissue. Pre-treatment with TQ or omega-3 before I/R resulted in significant decrease to (64.626%) and (44.867%) respectively of I/R control value (Fig. 1).

Effect of TQ or omega-3 on hepatic MPO: The normal value of hepatic tissue content of MPO was 33.200 U/mg.tissue. I/R resulted in significant increase to 87.250 U/mg.tissue. Pre-treatment with TQ or omega-3 before I/R resulted in significant decrease to (68.401%) and (52.149%), respectively of I/R control value (Fig. 2).

Fig. 2: Effect of TQ or omega-3 on hepatic content of MPO in intestinal I/R in rats
Thymoquinone (10 mg/kg; P.O.) as well as omega-3 (300 mg/kg; P.O.) was administered for 14 days before I/R; a: Significantly different from normal control at p<0.05; b: Significantly different from I/R control at p<0.05; Statistical analysis was carried out by one way Analysis of Variance (ANOVA) followed by Tukey-kramer test for multiple comparisons

Fig. 3: Hepatic histological findings of sham control group showed normal hepatocytes (A) those from I/R control group (B) showed strong degeneration and congestion in the hepatocytes with dilatation in the portal vein. Those treated with thymoquinone (C) showed minor degeneration in hepatocytes and dilatation in the portal vein (pv); Those treated with omega-3 (D) showed normal hepatocytes and dilatation in the central vein
Effect of TQ or omega-3 on hepatic histopathological finding: Rats subjected to I/R showed strong degeneration and congestion in the hepatocytes with dilatation in the portal vein. Rats treated with TQ showed minor degeneration in hepatocytes and dilatation in the portal vein. Rats treated with omega-3 showed normal hepatocytes and dilatation in the central vein (Fig. 3).

Effect of TQ or omega-3 on hepatic immunohistochemical staining of caspase-3: Immunohistochemical staining of caspase-3 in sham control group was moderately positive in 30% of hepatocytes (H-Score = 60). While it was strongly positive in 40% of hepatocytes (H-Score = 120) in rats subjected to I/R. Rats treated with TQ and omega-3 showed moderate staining of caspase-3 in 40% of hepatocytes (H-Score = 80; Fig. 4).

DISCUSSION

Intestinal I/R is considered to be a grave and triggering event in development of local and distant organ dysfunction (Zhao et al., 2010), which involves liver diseases (Sasaki and Joh, 2007).

In the present study; intestinal I/R-induced hepatic failure confirmed biochemically by increasing in the hepatic enzymes AST, ALT and ALP as a measure of hepatic function. The elevation of liver enzymes is accepted as the most sensitive indexes of acute hepatic injury (Zhang et al., 2006). These results are in agreement with Li et al. (2008) and Sheth et al. (2011). Who reported that I/R significantly increased the transaminase levels as compared with the sham operated group.

In support of our result histopathological examination showed strong degeneration and congestion in the hepatocytes in intestinal I/R rats which confirm the destructive effect of I/R on liver. The injury most likely due to the sudden reduction of the portal blood flow after the occlusion of the mesenteric artery followed by reperfusion, which increases the inflammation and oxidative damage in the liver. The significant reduction reported in the present study in the serum levels of AST, ALT and ALP of the thymoquinone and omega-3 treated groups confirm thecytoprotective effects of these drugs. Moreover thymoquinone showed a marked improve in the hepatocytes structure as well as omega-3 showed a normal histopathological result. This may be due to the direct protective effect of the drugs on the hepatocytes in addition to their anti-oxidant effect. TQ effect was in agreement with Daba and Abdel-Rahman (1998) who reported that TQ significantly protected isolated hepatocytes from tertbutylhydroperoxide induced toxicity and restores the levels of ALT and ALP. Omega-3 effect was in agreement with Meganathan et al. (2011) who reported that Omega-3-Fatty acids lowering AST, ALT and ALP levels, beside its anti-inflammatory effect due to decrease the production of pro-inflammatory cytokines.

In the present study, hepatic tissue content of TNF-α was significantly elevated in the intestinal I/R control group as compared to the sham-operated rats in agreement with Yang et al. (2001). Among the various cytokines, TNF-α has emerged as a key factor in assorted liver diseases (Hsu et al., 2010).
Reperfusion of the ischemic intestine results in intestinal mucosal barrier impairment, leading to endotoxemia that triggers a deleterious cascade including the production of ROS and further activates the intestinal macrophages and Kupffer Cells (KCs) (Towfigh et al., 2000). Activation of KCs, upregulates the production of inflammatory mediators including interleukin-1b, interleukin-6, tumor necrosis factor-alpha and platelet activating factor (Giakoustidis et al., 2006).

In the current study, thymoquinone and omega-3 markedly decreased hepatic tissue content of TNF-α. This is in accordance with Tekeoglu et al. (2006) who reported the anti-inflammatory effects of TQ on experimentally-induced arthritis in rats is associated with decreased levels of TNF-α. Salem (2005) attributed this to its antioxidant and anti-inflammatory properties. Omega-3 effect was in agreement with Skuladottir et al. (2007) who refer the anti-inflammatory effect of omega-3 due to the significant reduction in the levels of IL-6 and TNF-α.

The obtained data revealed that intestinal I/R resulted in a marked decrease in hepatic tissue content of GSH and the activity of SOD along with increased tissue content of LPO and MPO. Zhao et al. (2010) reported that the activity of SOD in the liver tissue decreased after intestinal I/R and stated that hepatic tissue content of GSH significantly decreased in the intestinal I/R group. A growing body of evidence indicates that oxidative stress plays an important role in the pathogenesis of many clinical conditions (Cross et al., 1987). These changes were significantly ameliorated by TQ and omega-3 concurrent treatment. Omega-3 administration could cause its beneficial effects by its known anti-inflammatory properties through displacement of arachidonic acid from the cellular membrane, shifting of prostaglandin E2 and leukotriene B4 production (Beyazit et al., 2010).

A very valuable and more selective tool to identify apoptotic cells in solid tissues may be the use of antibodies that specifically detect the cleaved (active) subunits of caspases-3, because activation of these enzymes constitutes a key molecular event during the process of apoptosis. In fact, the utility of this approach has been proven in previous studies by other groups (Tanaka et al., 2000; Gamonal et al., 2001). The present data showed that caspase-3, in rat liver was strongly activated in 40% of hepatocytes after intestinal I/R, compared to moderate activation in 30% of hepatocytes in sham control rats. This result are in agreement with Giakoustidis et al. (2006) and El Gazzar et al. (2007), who supposed that activated caspase-3 was widely expressed in the I/R group, although a very limited amount was detected in the sham operation animals. This activation of caspase-3 may explain the hepatic injury caused after the intestinal I/R. Thymoquinone administration in this study reduced apoptosis through decreasing the activation of caspase-3. This is in agreement with Fouda et al. (2008) who assumed that apoptosis and proliferative reactions are reduced by thymoquinone. On the other hand Xuan et al. (2010) believed that TQ is known to confer protection against tumor growth due to stimulation of tumor cell apoptosis. Administration of omega-3 in the present study also reduced caspase-3. This is in agreement with El-Ansary et al. (2011b). This reduction in caspase-3 may be attributed to the powerful antioxidant effect of omega-3.

As a quinone, TQ can be reduced by a variety of reductases to yield semiquinone (one reduction) or thymohydroquinone (two reductions). While the latter molecule is reported to have antioxidant effects (El-Najjar et al., 2010). TQ has potent superoxide anion scavenging abilities and inhibits iron-dependent microsomal lipid peroxidation (Badary et al., 2003). While the effect of omega-3 may be due to their anti-inflammatory effect (Beltz et al., 2007). The mechanisms include that omega-3 fatty acids decrease the production of pro-inflammatory cytokines, such as TNF-α (Song et al., 2009).

**CONCLUSION**

In conclusion, TQ and omega 3 have potential antifibrotic effect beside its anti-inflammatory and antioxidant activity via the protection of hepatic dysfunction induced by intestinal ischemia reperfusion.

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