Eucalyptus Globulus protects against diclofenac sodium induced hepatorenal and testicular toxicity in male rats

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Abstract

The current study was conducted to investigate the protective properties of Eucalyptus globulus leaves methanolic extract (EGLME) against diclofenac sodium (DS) induced hepatorenal and testicular toxicity in male rats. A total of 40 rats were equally divided into 4 groups, Control, Diclofenac sodium (DS), EGLME and DS + EGLME groups, respectively. DS and EGLME were administered orally at dose rate 2.5 and 100 mg/kg BW, 4 times/week for 8 weeks, respectively. Administration of DS distorted hepatorenal functions manifested by alteration of serum levels of ALT, AST, total protein and albumin, creatinine and urea with changes of histological architectures. DS caused reproductive toxicity represented by minimized sperm count, individual sperm motility and viability; depleted concentration of reduced glutathione (GSH) in testicular tissue; and decreased testosterone level with alteration in testicular histological features. In contrast, co-treatment of DS intoxicated rats with EGLME protected rats against the adverse effects of DS revealing enhancing properties of EGLME on rats' liver, kidney and testes. In conclusion, we demonstrated that EGLME had a potent protecting property against DS induced hepatic, renal and testicular toxicity in male rats, with special concern to testicular tissue via modulation of GSH as an oxidant marker.

Taxonomy: (classification by EVISE): Diclofenac sodium toxicity (hepatorenal and testicular toxicity), co-treatment with natural herbal extract, blood biochemical assays, tissue anti-oxidants assay, histopathology and reproductive indices analyses.

1. Introduction

Diclofenac sodium is considered the most commonly used anti-inflammatory drug of non-steroidal nature (NSAID) and analgesic substance.1,2 It is used for medical complications like rheumatoid arthritis, inflammation, trauma, degenerative joint disease, dysmenorrhea and surgical pains.3 Diclofenac acts on decreasing the prostaglandin synthesis through inhibition of cyclooxygenase enzymes and diminishing the apoptosis.4,5 Diclofenac induces hepatotoxicity and renal toxicity as a side effect.4,5 It is known that anti-inflammatory drugs badly affect reproduction.2,3 Administration of diclofenac caused liver and kidney toxicity, besides degenerative alteration in testis, epididymis and accessory sex glands.2 Moreover, diclofenac metabolism involves the production of reactive oxygen species leading to oxidative stress and genomic DNA fragmentation.2 Furthermore, previous work has suggested that diclofenac is cytotoxic to liver via cytochrome P-450 (CYP)-mediated metabolism.2 There is clear evidence to implicate drug-induced oxidative stress as a mechanism of toxicity in numerous tissues. ROS have effects on key cellular targets, namely, DNA, lipid,
and protein macromolecules. ROS may damage these critical cellular components at the molecular level, with consequent effects of ROS on cell survival mediated by kinase cascades. These factors may have a key role in initiating cell death in response to oxidative insult.10

Eucalyptus, which includes more than 700 species is the native plant of Australia. Eucalyptus globulus (E. globulus), were transferred to Europe and North Africa where they are well cultivated to the Mediterranean shores. Eucalyptus is mostly acclimated for cosmetics and paper industrial purposes and for traditional medicine. Recent trends in medicine depends on certain species of Eucalyptus as well.12,13 Recently, Eucalyptus is extensively cultivated for many purposes including antiseptic, chemotherapeutic, antioxidant, antimicrobial, respiratory and gastrointestinal disorders, healing, acaricidal, repellent, herbicidal and nematicidal.14–16 In this study we aimed to ameliorate and protect against the side effects of treatment with diclofenac sodium (DS) using novel extracts from natural herbal plants. To our knowledge, the protective effect of E. globulus against diclofenac induced reproductive toxicity in rats has not yet been studied. Therefore, the present study was undertaken to investigate the possible mechanism through which, E. globulus leaves methanolic extract (EGLME) deserve its protective potentials against DS induced hepatic, renal and reproductive toxicity in rats.

2. Materials and methods

2.1. Animals

A total of 40 male albino rats (Rattus norvegicus), weighing 100–120 g were used in this study. The animals were obtained from Abu Rawash, Giza Governorate, Egypt. Rats were kept in plastic cages with humidity (65%), temperature (20 ± 2 °C) and 12 h dark/ light cycles and fed normal chow diet. They were acclimatized 10 days before starting the experiment. The experiments were conducted in accordance with ethical guidelines of the Animal Care and Use Committee of University of Sadat City, which followed the Guide for the Care and Use of Laboratory Animals 8th edition. Washington (DC): National Academies Press (US); 2011.

2.2. Chemicals and reagents

Diclofenac sodium (Declophen®, tablets) was purchased from Pharco Pharmaceuticals, Egypt. Diagnostic kits for assaying serum activities of alanine aminotransferase (ALT), aspartate aminotransferase (AST), urea, creatinine and reduced glutathione (GSH) in testis tissue were purchased from Bio-diagnostic company, Egypt. Diagnostic kits for assaying serum levels of total protein (Diamond diagnostics, Egypt), albumin (SPECTRUM, Egypt).

2.3. Plant extract preparation

Fresh green leaves of E. globulus, 2 kg, were obtained from road trees at Sadat City, Menoufia, Egypt (location: 30.462629, 30.782271). E. globulus leaves were identified and authenticated in Biochemistry department, Faculty of Veterinary Medicine, University of Sadat City. Methanolic extract of E. globulus leaves were prepared as follows, in brief, fresh leaves of E. globulus were rinsed thoroughly with distilled water to remove dust and debris and shade dried at room temperature then ground into powder. The powder was immersed in 80% methanol for 48 h at room temperature (22 °C) with gentle shaking. The contents were filtered through filter paper (Whatman size No. 1), then the filtrate was dried at room temperature by air current to obtain a semisolid crude extract weighing (320 g) with 16% as a crude percent. This extract was stored in airtight container at 4 °C until used.

2.4. Gas chromatograph technique for characterization of bioactive constituents of Eucalyptus globulus

The potential bioactive constituents of Eucalyptus globulus methanolic leaf extract was characterized by using a Hewlett-Packard HP 5890-Series II gas chromatograph equipped with a Hewlett-Packard HP 5971 series mass detector (MS). Briefly, a representative sample (0.2 g) of Eucalyptus Globulus methanol extract was extracted with 10 mL of methylene chloride. A 1 μL of the sample was injected onto VF-Xms Agilent capillary column of (30 m × 0.25 mm ID × 0.25 μm film thickness). The initial oven temperature of the GC was 40 °C for 4 min and the temperature then programmed at a rate of 5 °C/min to 270 °C. The injector temperature and detector temperature were 270 and 250 °C, respectively, and the carrier gas was He of 99.99% purity. The m/z (ratio of mass to charge) values, which represent the fragment ions of the compounds were recorded for each compound.

2.5. Experimental design

To evaluate protective potentials of E. globulus leaves methanolic extract (EGLME) against diclofenac sodium (DS) induced toxicity. Animals were assigned into 4 groups (n = 10/group) as the following, control group, were received distilled water, DS group, only diclofenac sodium intoxicated, rats were given orally diclofenac sodium, 2.5 mg/kg BW, 4 times per week for 8 weeks dissolved in distilled water. E. globulus leaves methanolic extract (EGLME) group, rats were given orally EGLME, 100 mg/kg BW, 4 times per week for 8 weeks. DS + EGLME group, rats were diclofenac sodium intoxicated as DS group and co-treated with EGLME as in the EGLME group.

2.6. Blood and tissues sampling

Blood samples were collected at the end of the 8th week of the experiment. Rats were anaesthetized with Diethyl ether. Blood samples were collected from orbital venous plexus. Sera samples were separated and stored at −20 °C until used for investigation of biochemical parameters. Furthermore, rats were sacrificed, and livers, kidneys and testes were removed, washed with normal physiological saline solution and divided into 2 parts. The first part was stored at −80 °C and used for assaying reduced glutathione (GSH) (antioxidant defense system biomarker) in testicular tissue. The second part was kept in 10% neutral formalin for histopathological examination.

2.7. Preparation of testicular tissue homogenate for measuring of reduced glutathione (GSH)

Before dissection, tissues were perfused with phosphate buffer saline solution (PBS) pH 7.4 containing 0.16 mg/ml heparin to remove red blood cells and clots. Testicular tissues (222 mg) were homogenized by using 2 ml cold buffer (50 mM potassium phosphate, pH 7.5 and 1 mM EDTA for GSH). Tissue homogenate was centrifuged at 4000 rpm for 15 min at 4 °C. The supernatant was aspirated and stored at −80 °C till analysis.

2.8. Biochemical and hormonal assay

We have measured serum activities of ALT and AST,17 serum total protein concentration,18 serum albumin concentration,19 serum urea concentration,20 and serum creatinine level.21 Serum total testosterone level was measured using a competitive immunoassay
using direct chemiluminescent technology (The ADVIA Centaur®
CP Immunoassay System, Siemens Healthcare Diagnostics, USA).
GSH content of testicular tissue was determined according to the
procedure described by.22

2.9. Epididymal sperm preparation

The rats’ dissected cauda epididymis was kept in 1 ml of pre-
warmed phosphate buffer saline. Gentle agitation along with
tearing of the tissue was applied to make spermatozoa leak into the
PBS.23 Each sample was briefly incubated at 37 °C for 20 min for
further sperm analysis.

2.10. Evaluation of sperm count and motility

The motile sperm percents were estimated under (40 x ) using
phase-contrast microscope with high stage. Sperm motility (%) was
categorized as proportion of progressive (rapid and slow) and non-
progressive spermatozoa. Total number of motile sperm was
calculated as; motility (%) = number of motile sperm/total number
of sperms x100. Whereas total sperm count in a drop of the
resulting sperm suspension was determined using a Neubauer
chamber as described by.23

2.11. Evaluation of sperm viability

The viability was evaluated using eosin stain. Briefly, mixing
10 µl of sperm sample with 10 µl of dye (0.5% eosin stain) on a
microscope slide and covered with a coverslip. A total of 200 sperm
cells were counted shortly after addition of dye.23 Evaluation of live
(unstained) and dead (red stained) spermatozoa eae carried out
using light microscopy under X40.

2.12. Histopathological examination

Following necropsy, tissue specimens from liver, kidneys and
testes were collected and rapidly fixed in 10% neutral buffered
formalin solution. The fixed specimens were trimmed, washed,
dehydrated in ascending grades of ethyl alcohol, cleared in methyl
benzoate and processed through the conventional paraffin
embedding technique. A size of 3–5 µm sections were gained from
paraffin blocks using microtome (LEICA RM 2135) then routinely
stained by haematoxylin and eosin (H & E) stain according to
Ref.26. Prepared slides were checked using light microscopy and
photographed using a digital Leica photomicroscope (LEICA, DMLB,
Germany).

2.13. Semiquantitative histopathological evaluation

The histopathological lesions in liver, kidney and testes tissues
were examined in three rats from each group and five randomly
selected sections were examined from each rat. Semiquantitative
evaluation was performed in examined fields (n = 15) according to
the percentage, degree and extent of tissue damage and were
scored according to Michael, 2008 as follows: (−): normal
appearance (absence of pathological lesion 0%), (+): mild (<25% of
sections), (++): moderate (25–50% of sections), (+++): severe
(51–75% of sections), and (++++): very severe (>75% of sections).

2.14. Statistical analysis

The significant differences between groups were analyzed using
(GraphPad Prism 5, GraphPad Software, Inc., San Diego, CA, USA)
with a one-way analysis of variance (ANOVA) followed by Tukey’s
multiple comparison test for pairwise comparison of data from the
multiple groups. Results were statistically significant when
P < 0.05.

3. Results

3.1. Characterization of bioactive constituents of Eucalyptus
globulus

The potential bioactive constituents of Eucalyptus globulus
methanolic leaf extract was characterized by using a Hewlett-
Packard HP 5890-Series II gas chromatograph and revealed 4
compounds o-Isopropylanisole, Spathulenol, 2-Butanone, 4-(2,6,6-
trimethyl-2-cyclohexen-1-yl)- and β-Gurjuren (Table 1).

3.2. Eucalyptus globulus ameliorated diclofenac sodium induced
alteration in liver function biomarkers

Rats of the DS group showed significantly elevated (p < 0.05)
sodium activities of ALT, AST and significantly reduced serum levels
of total protein and albumin compared to those of rats in control
group. However, co-treatment of rats with EGLME and diclofenac
sodium in group DS + EGLME, significantly reduced (p < 0.05)
sodium activity of ALT, AST and significantly increased serum levels
of total protein and albumin compared to those of rats in DS group.
Administration of rats in EGLME group had no significant effects on
the mentioned parameters compared to control group (Table 2).

3.3. Eucalyptus globulus ameliorated diclofenac sodium induced
alteration in kidney function biomarkers

Rats of the DS group showed significantly elevated (p < 0.05)
sodium activities of creatinine and urea compared to those of rats in
control group. However, co-treatment of rats with EGLME and
diclofenac sodium in group DS + EGLME, significantly reduced
(p < 0.05) serum activity of creatinine and urea compared to those
of rats in DS group. Administration of rats in EGLME group had no
significant effects on the mentioned parameters compared to control
group (Table 2).

3.4. Eucalyptus globulus ameliorated diclofenac sodium induced
alteration in reproductive parameters

3.4.1. Regulation of reduced glutathione (GSH) concentration in
testicular tissue

Rats of the DS group showed significantly decreased (p < 0.05)
GSH concentration in testicular tissue compared to those of rats in
control group. However, co-treatment of rats with EGLME and
diclofenac sodium in group DS + EGLME, significantly enhanced
(p < 0.05) GSH concentration in testicular tissue compared to those
of rats in DS group. Administration of rats in EGLME group had no
significant effects on GSH concentration compared to control group
(Table 2).

3.4.2. Regulation of serum total testosterone level

Rats of the DS group showed significantly decreased (p < 0.05)
sodium total testosterone level compared to those of rats in control
group. However, co-treatment of rats with EGLME and diclofenac
sodium in group DS + EGLME, significantly enhanced (p < 0.05)
sodium total testosterone level compared to those of rats in DS group.
Administration of rats in EGLME group had no significant effects on
serum total testosterone level compared to control group
(Table 2).

3.4.3. Eucalyptus regulated sperm parameters

3.4.3.1. Total sperm count. Rats of the DS group showed
significantly decreased \((p < 0.05)\) total sperm count compared to those of rats in control group. However, co-treatment of rats with EGLME and diclofenac sodium in group DS + EGLME, significantly increased \((p < 0.05)\) total sperm count compared to those of rats in DS group. Administration of rats in EGLME group significantly increased \((p < 0.05)\) total sperm count compared to those of all groups (Table 3).

### Table 1
The major antioxidant compounds identified in Eucalyptus globulus leaf extract.

| Retention time (min) | Area % | Compound name | Structure |
|----------------------|--------|---------------|-----------|
| 25.95                | 3.05   | o-Isopropylanisole | ![](image) |
| 33.27                | 48.43  | Spathulenol     | ![](image) |
| 36.73                | 16.21  | 2-Butanone, 4-(2,6,6-trimethyl-2-cyclohexen-1-yl)- | ![](image) |
| 36.91                | 3.06   | \(\beta\)-Gurjurenene | ![](image) |

### Table 2
Effect of Diclofenac sodium and/or Eucalyptus globulus leaves methanolic extract on serum liver and kidney functions biomarkers, testicular tissue Reduced Glutathione (GSH) and serum total testosterone.

| Variables                  | Control       | DS            | EGLME         | DS + EGLME    |
|----------------------------|---------------|---------------|---------------|---------------|
| ALT (U/ml)                 | 16.22\(a\) ± 0.6915 | 23.54\(a\) ± 1.246 | 14.73\(b\) ± 0.4963 | 17.70\(b\) ± 0.2793 |
| AST (U/ml)                 | 90.93 ± 0.5586 | 104.0\(b\) ± 1.436 | 89.81\(b\) ± 0.7327 | 96.50\(b\) ± 1.203 |
| Albumin (g/dl)             | 3.140\(a\) ± 0.07385 | 2.129\(a\) ± 0.03621 | 2.964\(a\) ± 0.02467 | 2.875\(a\) ± 0.03846 |
| Total protein (g/dl)       | 8.268\(a\) ± 0.04716 | 6.108\(b\) ± 0.2360 | 8.228\(a\) ± 0.1588 | 7.806\(a\) ± 0.2040 |
| Creatinine (mg/dl)         | 0.3905\(a\) ± 0.02231 | 0.7937\(a\) ± 0.1208 | 0.4683\(a\) ± 0.01068 | 0.4048\(a\) ± 0.02005 |
| Urea (mg/dl)               | 46.74\(a\) ± 0.8980 | 58.11\(a\) ± 0.4042 | 46.14\(a\) ± 0.5095 | 47.51\(a\) ± 0.9046 |
| Reduced Glutathione mmol/g testicular tissue | 11.84\(a\) ± 0.06994 | 10.77\(b\) ± 0.01453 | 11.74\(a\) ± 0.1564 | 11.48\(a\) ± 0.08881 |
| Testosterone (ng/dl)       | 2.460\(a\) ± 0.2552 | 1.177\(b\) ± 0.1969 | 2.960\(a\) ± 0.09512 | 2.623\(a\) ± 0.2229 |

- Mean value ± SEM (standard error of mean). The mean difference is significant at \(P < 0.05\). The values carrying different superscript letters \((a, b, c)\) in the same row were statistically different.

### Table 3
Effect of Diclofenac sodium and/or Eucalyptus globulus leaves methanolic extract on sperm parameters (sperm count, motility and viability).

| Variables                  | Control       | DS            | EGLME         | DS + EGLME    |
|----------------------------|---------------|---------------|---------------|---------------|
| Count \((\times 10^6)\)    | 33.16\(a\) ± 2.45 | 19.33\(b\) ± 1.11 | 45.8\(a\) ± 0.91 | 39\(a\) ± 2.52 |
| Rapid motility (%) \(\text{Grade a}\) | 25\(a\) ± 1.66 | 10.91\(b\) ± 0.62 | 38.57\(b\) ± 2.62 | 21.25\(a\) ± 1.85 |
| Slow motility (%) \(\text{Grade b}\) | 35\(a\) ± 0.96 | 18.5\(a\) ± 2.07 | 37\(b\) ± 1.83 | 33.33\(a\) ± 1.66 |
| Non progressive motility (%) \(\text{Grade c}\) | 22\(a\) ± 0.79 | 22.85\(b\) ± 1.13 | 17.5 \(b\) ± 1.72 | 25.83 \(a\) ± 1.13 |
| Immotile sperm (%) \(\text{Grade d}\) | 18\(a\) ± 0.7 | 47.7\(a\) ± 2.98 | 6.93\(b\) ± 1.09 | 19.59\(a\) ± 0.7 |
| Total motility (%) \(\text{Grade a, b, c}\) | 82\(a\) ± 4.27 | 52.2\(b\) ± 7.69 | 93.07\(a\) ± 1.13 | 80.41\(a\) ± 1.13 |
| Viability (%)               | 80\(a\) ± 0.96 | 58.3\(b\) ± 1.88 | 86.4\(b\) ± 1.73 | 78.5\(a\) ± 1.36 |

- Mean value ± SEM (standard error of mean). The mean difference is significant at \(P < 0.05\). The values carrying different superscript letters \((a, b, c)\) in the same row were statistically different.

3.4.3.2. Sperm individual motility. Rats of the DS group showed significantly decreased \((p < 0.05)\) total sperm individual motility compared to those of rats in control group. However, co-treatment of rats with EGLME and diclofenac sodium in group DS + EGLME, significantly increased \((p < 0.05)\) total sperm individual motility compared to those of rats in DS group. Administration of rats in EGLME group significantly increased \((p < 0.05)\) total sperm individual motility compared to those of all groups (Table 3).
3.4.3.3. Sperm viability. Rats of the DS group showed significantly decreased (p < 0.05) sperm viability compared to those of rats in control group. However, co-treatment of rats with EGLME and diclofenac sodium in group DS + EGLME, significantly increased (p < 0.05) sperm viability compared to those of rats in DS group. Administration of rats in EGLME group significantly increased (p < 0.05) sperm viability compared to those of all groups (Table 3) (Fig. 1).

3.5. Eucalyptus globulus attenuated diclofenac sodium induced pathological alterations in rats’ hepatic tissues

Normal hepatocytes arranged in trabecular manner around normal central vein were observed in liver of control and E. globulus administrated rats. Administration of DS induced histopathological alterations in the liver including congestion of central and portal veins, granular degeneration of hepatocytes with granular cytoplasm, diffuse hydropic degeneration and vacuolation of hepatocytes in liver parenchyma. On the other hand, administration of EGLME with DS protected the liver from diclofenac toxicity and only single cell necrosis with pyknotic nucleus and mild congestion in hepatic sinuoids were observed (Fig. 2 and Table 4).

3.6. Eucalyptus globulus reduced diclofenac sodium induced pathological alterations in rats’ renal tissues

Normal histologic architecture of glomeruli, proximal and distal convoluted tubules were observed in kidneys of control and E. globulus administrated rats. Administration of DS resulted in shrinkage of glomerular tuft with increase of Bowman’s space, necrosis with pyknotic nuclei of glomerular cells, coagulative necrosis of some proximal convoluted tubules, damaged distal convoluted tubules with hyaline and cellular casts and interstitial edema. More normal glomeruli and renal tubules were observed in DS + EGLME group (Fig. 3 and Table 4).

3.7. Eucalyptus globulus restored diclofenac sodium induced pathological deteriorations in rats’ testicular tissues

The histo-architecture of testes of control and E. globulus administrated groups showed compactly arranged seminiferous tubules with germinal layers of all stages of spermatogenesis, spermatids, normal lumen of seminiferous tubules which contain spermatzoa, normal interstitium and interstitial Leydig cells. Testes of DS-administrated rats showed degenerated seminiferous tubules; some with desquamation of all spermatogenetic epithelium in their lumen, other seminiferous tubules with loss of primary and secondary spermatocytes and testicular degeneration with appearance of spermatic giant cell beside edematous fluid and vacuolation in the interstitial tissue. More or less normal seminiferous tubules with appearance of spermatzoa in their lumen were observed in DS + EGLME group (Fig. 4 and Table 4).

4. Discussion

Diclofenac sodium (DS) is one of the non-steroidal anti-inflammatory drugs (NSAIDs) with reversible inhibition of cyclooxygenase and causing serious clinical sequelae in human with acute NSAID overdose and these include convulsions, metabolic acidosis, coma and acute renal failure, and also were known for their side effects in rats including hepatotoxicity and renal toxicity; and seriously affect reproduction. Our study aimed to ameliorate the side effects of DS using a novel extract from natural herbal plants. The present study was undertaken to investigate the possible mechanism through which, E. globulus leaves methanolic extract (EGLME) protects against DS induced hepatic, renal and reproductive toxicity in rats.

Our findings reported that DS induced alteration in liver function biomarkers as it significantly elevated ALT, AST activities; significantly reduced serum total protein and albumin levels; and these findings were in accordance with the previous study reported by who showed that liver injury resulted from NSAIDs including diclofenac sodium may be proposed to their acidic moiety or reactive metabolites that bind to host proteins inducing cellular injury, and agreed with who denoted that diclofenac form significant alterations in serum protein contents by interference in protein metabolism. Furthermore, our results were in parallel with who reported that increased levels of serum glutamic oxaloacetic transaminase (GOT) and serum glutamic pyruvic transaminase (GPT) indicate a marker of liver injury and damage

3.7.1.3. Sperm viability. Rats of the DS group showed significantly decreased (p < 0.05) sperm viability compared to those of rats in control group. However, co-treatment of rats with EGLME and diclofenac sodium in group DS + EGLME, significantly increased (p < 0.05) sperm viability compared to those of rats in DS group. Administration of rats in EGLME group significantly increased (p < 0.05) sperm viability compared to those of all groups (Table 3) (Fig. 1).

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liver cell. Moreover, diclofenac affects serum glutamic pyruvic transaminase (GPT) transaminases and change the concentration of glutamate and aspartate in the extra-cellular environment.33

Liver is the organ for drug filtration, concentration, and excretion. The capillary filtration is the main method for excretion of soluble boundless medication, as an example, diclofenac.34 Prostaglandins are to blame for the regulation of nephritic blood flow, capillary filtration, modulation of coagulase unharNESS, hollow particle transport, and water metabolism.34

The present study revealed that, DS badly affected the renal function as represented by significantly elevated serum creatinine and urea when compared with control group (Table 2). As noticed from previous reports, the kidney is extremely active in the metabolism and synthesis of prostaglandins,35 diclofenac prevents the synthesis of prostaglandin by inhibiting COX pathway in the kidney36 resulting in renal dysfunction and pathophysiological alterations.37 In addition, it is well established that male fertility can be affected and impaired by several kinds of toxicants through their direct effects on testicular tissue or disrupt the endocrine activities of male reproduction.37–42 DS significantly decreased serum total testosterone level and reproductive sperm parameters such as sperm count, sperm individual motility and viability when compared to those of rats in control group (Table 3), these findings were similar to those obtained by43 and in harmony with the current results, diclofenac sodium decreased sperm count, sperm motility and density and inhibited prostaglandins synthesis in rats.36,44 Additionally,45 reported a significant decrease in the daily sperm production in the diclofenac treated rats besides a significant decrease in sperm motility and count with a significant decrease in the live/dead sperm in the diclofenac administrated rats, and we attributed these effects to the histopathological alterations noticed in testes of rats in DS group (Fig. 4). These findings were contrary with that, administration of diclofenac affected serum cholesterol and glucose concentrations which in turn reduced androgen in treated group as cholesterol is the precursor of the androgens.46 Treatment with diclofenac sodium salts reduced testosterone levels, feed intake and body weight.47 For understanding the mechanism behind the adverse effects of DS in testes, we analyzed the antioxidant marker, reduced glutathione (GSH) and our results revealed that there was a great depletion in GSH concentration in testicular tissue and this agreed with previous studies by,47 about the relation between diclofenac toxicity and GSH tissue concentrations.

On the other hand, we reported the enhancing properties of our novel methanol extract from Eucalyptus globulus leaves. In the concurrent study, Eucalyptus globulus ameliorated diclofenac sodium induced alteration in liver and kidney function biomarkers. Interestingly, Eucalyptus globulus ameliorated diclofenac sodium induced alteration in reproductive parameters through a mechanism depending on improving the antioxidant system inside the cell resulting in significant enhancement of GSH concentration in testicular tissue (Table 2). Administration of EGLME increased the serum total testosterone level, sperm parameters including sperm

### Table 4
Lesions in liver, kidney and testes from all groups and its histopathological grades.

| Lesions                                    | Control | DS    | EGLME | DS + EGLME |
|--------------------------------------------|---------|-------|-------|-------------|
| Liver lesions                              |         |       |       |             |
| Congestion of central & portal vein        | –       | ++    | –     |             |
| Congestion in hepatic sinusoids            | –       | ++    | +     |             |
| Coagulative necrosis in hepatocytes        | –       | ++    | –     | +           |
| Granular degeneration of hepatocytes       | –       | ++    | –     | +           |
| Hydropic degeneration                      | –       | ++    | –     | +           |
| Kidney lesions                             |         |       |       |             |
| Shrinkage of glomerular tuft with increase of Bowman’s space | –       | ++    | –     |             |
| Necrosis with pyknotic nuclei of glomerular cells | –       | ++    | –     |             |
| Focal coagulative necrosis of renal tubules | –       | ++    | –     |             |
| Vacularation of epithelial lining renal tubules | –       | ++    | –     |             |
| Hyaline and cellular casts                 | –       | ++    | –     | +           |
| Interstitial edema                         | –       | ++    | –     | –           |
| Testes lesions                             |         |       |       |             |
| Degenerated atrophied seminiferous tubules | –       | ++    | –     |             |
| Desquamation of spermatogenic epithelium & debris in the lumen | –       | ++    | –     |             |
| Loss of primary and secondary spermatocytes | –       | ++    | –     |             |
| Testicular degeneration with appearance of spermatid giant cell | –       | ++    | –     |             |
| Congestion of interstitial blood vessels   | –       | ++    | –     | +           |
| Edematous fluid and vacuolation in the interstitial tissue | –       | ++    | –     | –           |
and secondary spermatocytes (thin arrow) and edematous showing degenerated seminiferous tubules; some with desquamation of all spermatogenic epithelium in their lumen (thick arrow), other seminiferous tubules with loss of primary iniferous tubules with desquamated and necrosed spermatogenic epithelium and testicular degeneration with appearance of spermatid giant cell (thick arrow) (X20).

From the histopathological point of view, in the current study, absence of secondary spermatocytes, spermatids and spermatoozua indicated the impairment of spermatogenesis at the late sper-matocyte stage, which could be due to reduced testosterone level caused by diclofenac. Multinucleate giant cells were observed in the seminiferous tubules possibly due to fusion of macrophages to engulf deteriorated spermatogenic cells. Parallel with current data, diclofenac induced degenerative changes during treatment directly or indirectly via modulation of testosterone metabolism. Moreover, mentioned that treatments with diclofenac 0.5 mg/kg and 1.0 mg/kg in mice impaired seminiferous tubules and form degeneration, vacuolation, apoptosis in spermatogonia, primary spermatocytes, secondary spermatocyte and spermatids as well as in Sertoli cells and Leydig cells. These changes may lead to androgen disrupting activities as well as cellular supportive nutri-tive cellular toxicity. Here, the desquamated spermatogenic cells reflected the degenerative changes in the seminiferous tubules. These degenerative changes in the testes were not associated with lymphocyte or neutrophil aggregation supporting the fact that apoptosis was involved in diclofenac induced testicular toxicity. Interestingly, E. globulus restored the normal testicular structure. Some researches had pointed that apoptosis is the way by which the damaged spermatogenic cells were removed (Jahuukainen et al., 2000; Sertoli cells play important role in spermatogenesis. Leydig cells are main source of androgen production. Both types of cells can be affected by toxicants and chemical drugs. Alteration in the function of these cells may lead to changes in hormonal balance, disturbance in the process of spermatoozua development and impaired male fertility. Sertoli cells were degenerated, dilated intercellular space among germ cells, sloughing of the germ cells and accumulation in the lumen of seminiferous tubules which disturbed testicular tissue. Moreover, Diclofenac having drastic effects on testicular structure especially spermatogenic cell lines, interstitial cells and sertoli cells. In conclusion, we demonstrated for the first time that EGLME had a potent protecting properties against DS induced hepatic, renal and reproductive testicular toxicity in male rats, with special concern to testicular tissue via modulation of cellular antioxidant markers especially the reduced glutathione, we recommend for conducting further deep studies on farm animals for validation of the EGLME as a liver, kidney and testicular natural protector.

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