The Effect of Steroidal Extract of Nerium oleander on the Liver Tissue Structure of Albino Mice

Sahar A.A. Malik Al-Saadi, Karim H. Al-Derawi, Shiama Abd Alrasool
College of Science, Department of Biology, University of Basrah, Iraq
E.mail: Saharmalik2010@gmail.com

Abstract: Nerium oleander is an evergreen shrub of Apocynaceae family cultivated worldwide as an ornamental plant. All parts of the plant are toxic and contain a mixture of very toxic cardiac glycosides of cardenolides. Albino mice were divided into four groups. Group I was normal control group; and others represented three concentrations of steroid component of Nerium oleander (0.003, 0.006 and 0.012 mg/kg). Liver marker enzymes were assayed in serum. Levels of marker enzymes such as alanine aminotransferase (ALT) and aspartate aminotransferase (AST) were increased significantly. Sections of livers were observed under microscope for the histopathological changes such as destruction of the liver architecture with increased inflammation was observed compared with control.

Keywords: Nerium oleander, Leaves, Steroid extract, Toxicity, Mice.

I. Introduction
Nerium oleander (common oleander) is an evergreen shrub belongs to the family of Apocynaceae. It originates from the Mediterranean region and widely cultivated as an ornamental plant in many parts of the world, particularly in warm temperate and subtropical regions. This plant has long been known to be poisonous to animals and human beings. All parts of the plant are toxic and dry leaves are about as toxic as green ones (Kingsbury, 1964; Hardin and Arena, 1974; Zibbu and Batra 2010).

There are large numbers of secondary metabolic compounds found in plants such as alkaloids, saponins, flavonoids and anthraquinones phytosterol. Parts of the plant can be ingested intentionally in suicidal attempts or accidentally. Following the ingestion of oleander, the clinical picture usually starts with gastrointestinal signs: vomiting, nausea, abdominal pain and diarrhea. Later cardiovascular and neurological symptoms can occur. (Goetz, 2005) reported that Oleander contains a mixture of cardenolides, the most important of which are oleanadrin and oleandrinigenin (Zibbu and Batra 2010). Oleandrin constitutes approximately 0.08% of the total content of cardenolides present in N. oleander (Marchioni and Distefano, 1989).

Nerium oleander as important plant used the leaves and the flowers are cardiotonic, diaphoretic, diuretic, anticanancer, antifungal antibacterial (Abe and Yamauchi, 1992; Wang et al. 2000). This is a very poisonous plant, containing a powerful cardiac toxin and should only be used with extreme caution. The root is powerfully resolvent. Oil prepared from the root bark is used in the treatment of leprosy and skin diseases of a scaly nature. Plants have an extensive root system and are often used to stabilize soil in warmer areas. Seeds are Poisonous, abortifacient and alternative. They used as purgative in dropsy and rheumatism. The whole plant is said to have anticanancer properties (Abe and Yamauchi, 1992; Wang et al. 2000; Zibbu and Batra 2010). For this purpose, we extraction of toxic components of N. oleanderspecialized steroids extraction.

The aim of this research is to investigate and examine the hepatic histologist structure alterations and biochemical parameters that may be caused by administration of Nerium oleander leaf extract to albino mice.

II. Material and Methods

Leaf extract
Leaf samples of Nerium oleander were collected from University of Basrah from the garden, where the leaves are in fully green color. The leaves were thoroughly washed with water, dried and then ground with a blender. The powder (25) gm was extracted with methanol 90%. The extract was with by using sexhlet aperture for 24 h, the extract filtered and then dried.

The animals
A total number of 80 adult male albino mice were used in this research. The mice were in good health, approximately having the same weight (30 ± 5 gm) and 75 days in age. They were collected from Medical college Basrah University. They were housed in groups of 3 animals per cage at a temperature of 18-25°C under a 12 h dark–light cycle. The animals were left for one week before start of treatments. The animals were grouped into four groups each one is ten mice, the first one is control which was treated orally with distilled water, the second group 2,3,4: was treated orally using the stomach tube with N. oleander extract at dose of
0.003, 0.006, 0.012 gm/kg respectively. At the end of each treatment the animals were anaesthetized with diethyl ether, fixed the liver were taken for sectioning.

**Biochemical determination**

The biochemical parameters (serum enzymes): alanine aminotransferase (ALT) and aspartate aminotransferase (AST) and Alkaline phosphatase (ATP) were assayed spectrophotometrically using a commercially available assay kits according to the manufacturer’s protocol.

**Preparation of tissues**

Tissue samples were fixed in equivalent formaldehyde solution (10%), then washed with tap water. The samples were then dehydrated by being passed through series of ethanol from 30% up to 100%. Xylene was then applied for 30-40 minutes to clear the samples from ethanol. The tissues were subjected to paraffin infiltration using melted paraffin wax inside the oven at 60°C for one hour and the tissue samples were transferred from the oven inside these moulds which were then cooled to room temperature. The tissue sections were stained by using hematoxylin for one minute then in alcoholic eosin 1%.

**Statistical analysis**

The data are expressed as mean ± SD. Data were analyzed by analysis of variance (ANOVA) followed by multiple comparisons using SPSS test to compare all groups against control. Results were considered statistically significant at $p < 0.05$.

### III Results and Discussion

#### Biochemical parameters

Biochemical parameters studied were found to increase in the extract treated mice at three doses as compared to the control mice. Values of AST (Aspartate aminotransferase), ALT (Alanin aminotransferase) and ALP (Alkaline phosphatase) significantly increased ($p < 0.05$) by 126.50, 95.67 and 33.17 % respectively in the mice treated with doses of 0.012 mg/kg body weight/day as compared to the controls. Such increases were insignificant with the 0.006 mg/kg body weight/day in ALT and AST stroidal extract treatment (Table 1).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>ALT</th>
<th>AST</th>
<th>ALP</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>85.17±1.17°</td>
<td>24.50±2.06°</td>
<td>24.83±1.12</td>
</tr>
<tr>
<td>0.003</td>
<td>74.83±2.12°</td>
<td>113.00±2.32°</td>
<td>19.83±1.11 °</td>
</tr>
<tr>
<td>0.006</td>
<td>88.00±2.40°</td>
<td>88.17±22.14°</td>
<td>20.00±1.33°</td>
</tr>
<tr>
<td>0.012</td>
<td>95.67±2.15 °</td>
<td>126.50±2.17°</td>
<td>33.17±2.15 °</td>
</tr>
<tr>
<td>R.L.S.D.</td>
<td>4.42</td>
<td>9.92</td>
<td>3.83</td>
</tr>
</tbody>
</table>

*Values are mean±SEM.

#### Histopathology of the liver

Histopathological examinations of liver sections under light microscope revealed that there were several changes in the mice treated with all doses of the steroidal extract as compared with the control group. These changes include: congestion, inflammations around central vein in the 0.003 mg/kg bodyweight/day treated mice after two weeks, as well as showed inflammations around portal triads, Kupffer cells appear after four weeks (Fig 1); cellular infiltrations and severe cytoplasm vacuolations (hydropic degenerations) of hepatocytes around central veins in which their nuclei were stained pale and pushed to the periphery in the 0.012 mg/kg body weight/day steroidal extract treated mice, found necrosis and degenerations liver cells in addition inflammation of cells after four weeks. In addition, the mice treated with 0.012 mg/kg body weight/day showed congestion and degenerations liver cells and necrosis after four weeks from treatment. One of the mechanisms of plant extracts is the process of inhibiting the synthesis of purine and Alburuyemidn (Henghe Tian et al., 2007)
The Effect of Steroidal Extract of Nerium Oleander on the Liver Tissue Structure of Albino Mice

Figure -1: Photomicrographs of H and E stained liver sections from mice treated with Steroidal extracts of *Nerium oleander* at control mice (A) 0.003 mg/kg body weight/day showed congestion central vein and degeneration of endothelial cells after two weeks (B), 0.003 mg/kg body weight/day showed inflammations around portal triads, Kupffer cells appear after four weeks (C), 0.003 mg/kg body weight/day. Expanded central veins and The nuclei became Thickness of nuclei (D), aggregate of inflammation thicker, cells and sever hydropic degenerations (HD)(E), In 0.006mg/kg body weight/day observed hydropic degenerations and haemorrhage (F), while in 0.012mg/kg weight/day found necrosis and degenerations liver cells in addition inflammation of cells after four weeks(G). In mice treated with 0.012 mg/kg body weight/day showed congestion and degenerations liver cells and necrosis after four weeks from treatment(H). CV in A, B, C, D, E, F G & H = Central vein (Magnifications, all 450 X).

Pyknotic nuclei

III. Results and Discussion

observed in the study, the steroidal extract compared with treatment control group showed an increase in the levels of ALT, AST and increase alkaline phosphatase levels in 0.012 mg/kg body weight/day usually increases remarkably in disease and lesser extent in hepatocellular diseases. The decrease observed in the increased levels of the enzymes AST, ALT, Alkaline phosphatase may be attributed to the protection offered by the administration of the steroidal extract of Nerium oleander. Many researcher reports the chemical component found in plant extract contain phytochemicals can interrel with metabolism process and indicates the role of certain flavonoids, triterpenoids and steroids in hepatoprotection against hepatotoxins. The presence of those compounds in Nerium oleander may be responsible for the protective effect on liver damage in mice.

In conclusion, our results show that many effects of *N. oleander* on liver and enzymes the hepatoprotective effects of may be due to its antioxidant and free radical scavenging properties.
The Effect of Steroidal Extract of Nerium Oleander on the Liver Tissue Structure of Albino Mice

References