Biochemical and Haematological Studies on the Ethanol Leaf Extract of Spondias mombin Linn

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Abstract

Spondias mombin Linn (Anacardiaceae) is an edible plant that is ethnomedicinally used for induction of labour, expulsion of afterbirth, and stabilization of women after childbirth. The effects of ethanol extract of the leaf of S. mombin on hepatic, renal and haematologic function indices of albino rabbits were studied using standard methods. Acute toxicity studies of the extract showed the lethal dose (LD₅₀) to be indeterminate, while the effective (abortifacient) dose (ED₅₀) was 753.96 ±0.10 mg/kg body weight. Intraperitoneal administration of 750 mg/kg body weight of the extract significantly (p<0.05) reduced serum alanine aminotransferase and aspartate aminotransferase activities, as well as the serum concentrations of total bilirubin, urea and creatinine, but increased the serum concentrations of total protein, albumin and haemoglobin, as well as the values of white blood cell count, platelet count, and the calculated red cell indices. The results of the study showed that the extract does not have detrimental toxicological effects on the studied organs/tissue function indices. However, it’s observed effect on platelet count calls for further studies.

Keywords: Hog plum; Liver; Kidney; Haematologic indices; Electrolytes

Introduction

Spondias mombin is a common flowering plant that is commonly called hog plum. Locally in Nigeria, it is known as akika in Yoruba, ijikara in Igbo and tsadar masar in Hausa [1]. The plant leaves are common animal forage. The tender leaves and ripe fruits are also edible. The bark leaves and fruit juice have been widely used for varied folk medical purposes [2]. Preliminary researches reported the plant to have a wide range of antibacterial, antiviral and antifungal properties [3,4]. Reports on the plant have been interesting, yet controversial. It was reported to have abortifacient and uterine muscle contraction effects [5,6]. On the other hand, Kramer et al. [7] recommended its use for pregnant woman but only after five months of pregnancy. They noted that the observed cytotoxic effects of the plant may have some benefits in protecting the foetus from pathogens. Furthermore, they claimed that its high level of cytotoxicity, is indicative of analgesic properties. Thus they concluded that the use of the plant to ease pain during childbirth supports this evidence [7]. In the same vein, it was earlier observed [8] that the infusion of S. mombin leaves is variously used, without any reported collateral effects due to its activity. However, apart from the abortifacient effect noted above, Raji et al. [9] showed that the aqueous leaf extract of the plant has a dose-dependent anti-fertility action, but with full recovery achieved within four weeks after cessation of treatment with the extract.

Given the varied ethnomedicinal uses of the plant, the alleged non-collateral effects when ingested and the reported possible anti-fertility and abortifacient effects, the present study was designed to assess the safety of S. mombin leaves using the hepatic, renal and haematologic function indices of albino rabbits.

Materials and Methods

Plant material

Leaves of Spondias mombin were obtained from the bushes in Obinze and Iheagwa villages in Owerri West Local Government Area of Imo State. The leaves were duly authenticated at the Department of Crop Science, Federal University of Technology, Owerri. Voucher specimen was deposited in the departmental herbarium.

Preparation of plant material

The leaves of S. mombin were plucked from the stem stalks, rinsed in clean water and air-dried at room temperature. The dried leaves were ground into fine powder with a mechanical grinder (Heman, Japan). About 1.4 kg of the powdered leaf was soaked in 3.5L of 75 % ethanol in a 5 L beaker and covered with aluminum foil. The mixture was stirred about 3 hours for proper mixing and allowed to stand for 24 hours. The resulting decoction was filtered and the filtrate was subjected to a slow but complete solvent evaporation using a regulatory hot plate (Techmel, USA) at a temperature of 40-60°C. The extract was packaged in an air-tight container, labeled and stored below 4°C in a freezer until required [10].

Acute toxicity Tests

Lethal dose (LD₅₀) determination: The lethality of the S. mombin leaf extract was estimated using 36 healthy female albino mice divided into 6 groups with 6 mice each group. Each group received a dose of extract (ranging from 500-3000 mg/kg in 1ml of normal saline) intraperitoneally. The number of deaths in each group within 24 hours was recorded. The LD₅₀ was then estimated from the graph of percentage mortality (converted to probit) against log-dose of the extract using the probit method of Miller and Tainter [11].

Effective Abortion dose (ED₅₀) determination: Thirty-six pregnant mice at about the 15th day (third trimester) of pregnancy were used for the study. The animals weighing between 15 and 30 g were divided into six groups with six animals in each group. They

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were treated with increasing doses of the ethanol extract ranging from 250 to 1500 mg/kg, administered intraperitoneally as a single dose in 1ml of normal saline. All the animals were allowed access to feed (Guinea Feed Nigeria Ltd) and water ad libitum, and observed for treatment related abnormalities. These included foetal abortion and presence of vaginal bleeding for a period of 24 hours. They were further observed for general behavior for extra 7 days. The number of animals that aborted within the first 24 hours was noted for each group and subsequently ED₅₀ was calculated by the graphical probit method of Miller and Tainter [11]. The ED₅₀ was determined as a guide for dosage level to use for the biochemical and haematological effects study.

Animal grouping for Biochemical and Haematological Studies:
Thirty-five female albino rabbits (1.47 ± 0.17 kg) were obtained locally from Owerri, Imo State. The animals were housed in stainless steel cages under standard laboratory condition of 12 hours light/dark cycle. They were allowed access to feed (Guinea Feed Nigeria Ltd) and water ad libitum and allowed to acclimatize for 7 days before grouping and treatment.

The thirty-five rabbits were randomly divided into 7 groups with 5 rabbits in each group as follows:

- Group 1: the baseline control that were not administered any drug and were sacrificed on the zero day.
- Group 2: administered normal saline per kg and served as control for the 7 days treatment.
- Group 3: administered 0.14 IU/kg of standard oxytocin drug (Pitocin®, USA) for 7 days.
- Group 4: administered 750 mg/kg of S. mombin extract for 7 days.
- Group 5: administered normal saline per kg and served as control for the 14 days treatment.
- Group 6: administered 1ml (0.14 IU/kg body weight) of standard oxytocin drug (Pitocin®, USA) for 14 days.
- Group 7: administered 750 mg/kg of S. mombin extract for 14 days.

The animals were administered 1 ml of their respective drugs/extracts reconstituted in normal saline, intraperitoneally, once daily for their stipulated period.

Sample collection
At the end of each experimental period, 5 fasting animals were randomly selected from each group, anaesthetized with chloroform, and about 20 ml of blood collected by cardiac puncture from each animal. The blood sample was dispensed, 5ml each, into an EDTA bottle, a lithium heparin bottle and a non-anticoagulant containing (plain) bottle. The sample in the EDTA bottle was mixed thoroughly and used for hematological studies. The blood sample in the heparin container was centrifuged (Teco, USA) at 4000 rpm for 5 min to obtain the plasma which was stored frozen until used for the determination of extracellular cations concentrations, while the packed red cells were washed thrice with normal saline, drained dry and lysed with 1ml of distilled water and stored frozen until used for the determination of intracellular cations concentrations. The sample in the plain container was allowed to stand for 30 min to clot, and then centrifuged at 4000 rpm for 5 min. The serum was carefully separated into another sterile plain bottle and stored frozen until used for the analysis for liver and kidney function indices [11].

Liver function indices
The serum activities of alanine aminotransferase (ALT; EC 2.6.1.1), aspartate aminotransferase (AST; EC 2.6.1.2) and alkaline phosphatase (ALP; EC 3.1.3.1), as well as the serum concentrations of total bilirubin, conjugated bilirubin, total protein and albumin were determined [12] using standard kits (Human Laboratories, Germany). The unconjugated bilirubin concentration was calculated as the difference between total and conjugated bilirubin concentrations.

Kidney function and electrolyte indices
Serum urea and creatinine concentrations were determined by Jaffe’s reaction and urease enzymatic method respectively [13], using standard kits (Human Laboratories, Germany). Serum bicarbonate (as total CO₂) concentration was determined by the use of a CO₂ gas electrode (ASTRA CO₂ apparatus, Beckman Instruments, USA), while chloride concentration was determined by the titrimetric method of Schales and Schales [13]. Serum (extracellular) and erythrocytic (intracellular) sodium, potassium, calcium and magnesium concentrations were determined by use of an atomic absorption spectrophotometer (Alpha 4, Chem. Tech. Analytical, England).

Haematological indices
The haematological indices involving packed cell volume (PCV), haemoglobin (Hb) concentration, white blood cell (WBC) count, platelet count and the red blood cell indices including mean corpuscular volume (MCV), mean corpuscular haemoglobin (MCH) and mean corpuscular haemoglobin concentration (MCHC) were determined using an automated haematology analyzer machine (Mindray BC 2300, USA) which performs blood cell count by Direct Current detection method, haemoglobin analysis by non-cyanide haemoglobin analysis method and calculates the red blood cell constants automatically from red blood cell count, Hb and PCV.

Statistical analysis
Data generated were analyzed by the use of one-way ANOVA and Bonferroni’s multiple comparison tests, with the aid of a computer-based statistical package (Graphpad Prism 5.3). Inferences were made at 95% confidence level.

Results

Yield of ethanol extract of dry leaves of S. mombin
One kilogram weight of dry leaves of S. mombin yielded 27.71g (1.98%) of crude ethanol extract.

Acute toxicity studies
The results of the acute toxicity studies showed the LD₅₀ to be indeterminate because no animal died within the period of study, while the abortion ED₅₀ in pregnant mice was 753.96 ± 0.10 mg/kg body weight.

Effect of spondias mombin leaf extract on liver function indices
Administration of S. mombin extract and oxytocin drug significantly (P<0.05) reduced serum ALT and AST activities at both the 7th and 14th days of treatment (Table 1). Serum ALP activity was, on the other hand, significantly (P<0.05) increased by the extract, only after 14 days of administration from a baseline value of 64.08 ± 3.60 IU/l to 103.09 ± 11.05 IU/l. The extract administration non-
significantly (p>0.05) reduced serum total and unconjugated bilirubin concentrations. Conversely, the extract significantly (p<0.05) increased serum total protein and albumin concentrations in comparison with their respective baseline values.

Effect of *Spondias mombin* leaf extract on kidney function and electrolyte indices

**Table 1:** Effect of ethanol extract of *S. mombin* leaves on liver function indices.

<table>
<thead>
<tr>
<th>Indices</th>
<th>Baseline</th>
<th>Control</th>
<th>Oxytocin</th>
<th>Extract</th>
<th>Day 0</th>
<th>Day 7</th>
<th>Day 14</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alanine Aminotransferase (IU/l)</td>
<td>26.41 ± 2.52a</td>
<td>26.70 ± 2.21a</td>
<td>15.15 ± 2.92a</td>
<td>17.89 ± 1.08a</td>
<td>28.82 ± 2.20a</td>
<td>18.09 ± 1.01a</td>
<td>19.81 ± 1.75a</td>
</tr>
<tr>
<td>Aspartate Aminotransferase (IU/l)</td>
<td>43.27 ± 1.56b</td>
<td>43.42 ± 0.48b</td>
<td>17.43 ± 1.32b</td>
<td>13.64 ± 1.38b</td>
<td>42.24 ± 2.17b</td>
<td>14.19 ± 0.82b</td>
<td>16.47 ± 0.87b</td>
</tr>
<tr>
<td>Alkaline Phosphatase (IU/l)</td>
<td>64.08 ± 3.60b</td>
<td>67.72 ± 3.87b</td>
<td>75.48 ± 2.52b</td>
<td>84.83 ± 7.49b</td>
<td>65.16 ± 2.19b</td>
<td>88.40 ± 4.45b</td>
<td>103.09 ± 11.05b</td>
</tr>
<tr>
<td>Total bilirubin (µmol/l)</td>
<td>5.51 ± 1.51c</td>
<td>5.54 ± 1.53c</td>
<td>5.36 ± 0.71c</td>
<td>4.90 ± 0.20c</td>
<td>5.40 ± 0.12c</td>
<td>6.94 ± 0.68c</td>
<td>4.61 ± 0.86c</td>
</tr>
<tr>
<td>Conjugated bilirubin (µmol/l)</td>
<td>1.96 ± 0.22c</td>
<td>2.06 ± 0.27c</td>
<td>2.45 ± 0.31c</td>
<td>2.22 ± 0.34c</td>
<td>1.95 ± 0.15c</td>
<td>2.25 ± 0.09c</td>
<td>1.98 ± 0.22c</td>
</tr>
<tr>
<td>Unconjugated bilirubin (µmol/l)</td>
<td>3.53 ± 0.86c</td>
<td>3.48 ± 1.46c</td>
<td>3.91 ± 0.66c</td>
<td>2.69 ± 0.40c</td>
<td>3.45 ± 0.06c</td>
<td>4.69 ± 0.85c</td>
<td>2.63 ± 0.80c</td>
</tr>
<tr>
<td>Total protein (g/l)</td>
<td>61.33 ± 1.89b</td>
<td>62.14 ± 4.23b</td>
<td>52.56 ± 3.22b</td>
<td>66.43 ± 5.86b</td>
<td>67.51 ± 3.50b</td>
<td>57.06 ± 3.41b</td>
<td>72.49 ± 3.00b</td>
</tr>
<tr>
<td>Albumin (g/l)</td>
<td>30.23 ± 4.12b</td>
<td>31.99 ± 4.70b</td>
<td>30.39 ± 3.56b</td>
<td>47.89 ± 1.79b</td>
<td>30.64 ± 1.96b</td>
<td>35.59 ± 3.85b</td>
<td>53.50 ± 2.96b</td>
</tr>
</tbody>
</table>

Values are mean ± standard deviation (number of animals per group = 5); values with different superscript(s) in a row are significantly different (p<0.05).

Effect of *Spondias mombin* leaf extract on haematological indices

Administration of both *S. mombin* extract and oxytocin did not significantly (p>0.05) alter packed cell volume, mean corpuscular haemoglobin and mean corpuscular haemoglobin concentration values of the treated animals in comparison with those of the controls. However, treatment with the extract for up to 14 days significantly (p<0.05) increased mean corpuscular haemoglobin concentration (Figures 2, 3). Both the extract and oxytocin administrations significantly (p<0.05) increased the white blood cells and platelet counts of the treated animals (Figure 4).

Discussion

Acute toxicity effect of *Spondias mombin* leaf extract

The results of the acute toxicity study showed an indeterminate LD₅₀ even at 3 g/kg extract administration. The animals, especially those on higher doses of the extract showed reduced agility and were occasionally clustered together at one corner of their cages. This is in corroboration with earlier findings [14] that, apart from occasional clustering of the rats, acute toxicity test with up to 2 g/kg of ethanol leaf extract of *S. mombin* showed no lethal effect.

The abortion ED₅₀ in the pregnant mice was found to be 753.96 ± 0.10 mg/kg. The result is in consonance with an earlier report of 750 ± 20 mg/kg [5].

Effect of *Spondias mombin* leaf extract on liver function indices

The reductions in alanine and aspartate aminotransferases (ALT and AST) activities observed following administration of *S. mombin* leaf extract may be attributed to reduced rate of synthesis of the liver enzyme as a consequence of extract exposure. Administration of the extract for up to 14 days did not further depreciate the enzymes activities in comparison with activities noted at the 7th day, indicating an adaptation by the liver cells to the assault from the plant extract.

On the other hand, the initial 7 days administration of the extract did not affect serum alkaline phosphatase (ALP) activity, another marker enzyme employed to assess the integrity of hepatic plasma membrane. However, further administration for up to 14 days increased significantly ALP activity. The later increase in ALP activity observed may be due to enzyme induction by the extract [15]. The increased ALP activity may be attributed to the effects of some divalent ions such as Mg²⁺ and Ca²⁺ which are established activators of ALP and are reported to be present in the leaves [16]. This inference could further be buttressed by the observed overall reduction in the concentrations of total and unconjugated bilirubin, indicating that the xenobiotic conjugative and excretive potentials of the organ were not affected by the administration of the extract. Furthermore, the extract elicited time-dependent increases in serum total protein and albumin concentrations. The liver is the sole site for synthesis of albumin, which makes up approximately 60% of serum protein concentration. The observed increases in total protein and albumin concentrations with administration of the extract indicate that the synthetic function of the hepatocytes were not impaired. The increased synthesis of albumin may also have contributed in the reduction in overall unconjugated bilirubin, since it is the sole transporter of this water insoluble by-product of haemoglobin metabolism to the liver.

![Figure 1: Effect of ethanol extract of Spondias mombin leaves on serum urea and creatinine concentrations.](image-url)
It could be inferred that the administration of *S. mombin* leaf extract does not have hepatotoxic effect. The extract is rather hepatoprotective. This inference is corroborated by earlier observations [9] that *S. mombin* extract administration does not significantly alter liver function parameters. Thus, it may be claimed that the extract may prevent hepatic cell destruction, which is usually marked by an increase in blood aminotransferases activities, increased bilirubin concentration and reduction in serum protein and albumin concentrations [17]. Similarly, administration of the oxytocin standard drug did not cause a significant (p>0.05) increase in intracellular Ca\(^{2+}\) concentration, while significantly (p<0.05) affect both chloride and bicarbonate ions concentrations. This also points to the diuretic potential of the plant in ethnomedical practice. Thus, it may be claimed that the extract may prevent hepatic cell destruction, which is usually marked by an increase in blood aminotransferases activities, increased bilirubin concentration and reduction in serum protein and albumin concentrations [17].

**Effect of Spondias mombin leaf extract on kidney function and electrolyte indices**

Administration of *S. mombin* extract significantly (p<0.05) reduced serum urea and creatinine concentrations. This substantiates the reported diuretic potential of the plant in ethnomedical practice [2]. The extract did not significantly (p>0.05) affect intracellular concentrations of Na\(^+\) and K\(^{+}\), but increased (non-significantly, p>0.05) extracellular Na\(^+\) with a concomitant time-dependent reduction in K\(^{+}\) extracellular concentration. This also points to the diuretic potential of the extract, given that the extract is a rich source of potassium [16]. On the other hand, the administration of the extract did not significantly (p>0.05) affect both chloride and bicarbonate ions concentrations.

Furthermore, the extract caused a significant (p<0.05) increase in the intracellular Ca\(^{2+}\) concentration, while significantly (p<0.05) reducing the extracellular Ca\(^{2+}\) concentration. The extract as well as oxytocin standard drug had inverse effects on intracellular and extracellular Mg\(^{2+}\) to those on Ca\(^{2+}\). These inverse effects on intra- and extracellular Mg\(^{2+}\) concentrations by both the extract and oxytocin may be a reflection of the tendency of the body to balance cations level in the intra- and extracellular fluids. Factors that modulate renal Mg excretion can have profound effects on Mg balance. Thus, the apparent hypermagnesaemia created by the administration of the extract may lead to increased urinary excretory potential and consequently increased diuresis.

Unlike the extract, oxytocin administration reduced the concentrations of intracellular K\(^{+}\), explaining its action through efflux of K\(^{+}\) and influx of Ca\(^{2+}\) into the cell. Meanwhile, the significant (p<0.05) increase in intracellular Ca\(^{2+}\) concentration elicited by both the extract and oxytocin indicates activation of the Ca\(^{2+}\)/K\(^{+}\) pump which is necessary for the maintenance of the action potential required for initiation of muscle contraction.

**Effect of Spondias mombin leaf extract on haematological indices**

This study shows that *S. mombin* extract administration did not significantly increase haemoglobin (Hb) concentration, packed cell volume (PCV), erythrocyte sedimentation rate (ESR), and white blood cell count (WBC) levels of the treated animals. This indicates that the extract does not have a haemotoxic effect. This observation is corroborated by the earlier discussed overall reduction in total and unconjugated bilirubin concentrations, which are indicative of non red blood cell-lysing effect of the extract. However, the slight increase in WBC observed after 14 days of extract and oxytocin administrations could be attributed to the response of the immune system to the assault on the animals’ system by the drugs’ administration. The PCV was not significantly affected by the extract indicating that the amount of red blood cells present in the animals remained fairly constant. Similarly, the extract did not cause a drop in the haemoglobin concentration. Thus, the oxygen-carrying capacity of the blood of the animals was therefore not detrimentally affected by the extract administration. The erythrocyte indices, mean corpuscular haemoglobin (MCH) and mean corpuscular haemoglobin concentration (MCHC) are used to mathematically define the concentration of Hb within the cell. Extract administration produced an overall non-significant increase in MCH and MCHC, further supporting the observation that Hb concentration and hence, the oxygen-transporting potential of the blood of the extract-administered animals were not detrimentally affected. The mean corpuscular volume (MCV) is regarded as the average volume of a single red blood cell. There was a non-significant reduction in the MCV when compared with the respective control (Day 14) value.

![Figure 2](http://example.com/f2.png)  
*Figure 2: Effect of ethanol extract of *Spondias mombin* leaves on packed cell volume and haemoglobin concentrations. *Value significantly (p<0.05) reduced in comparison with the respective baseline (Day 0) value.*

### Table 2: Effect of ethanol extract of *S. mombin* leaves on serum and red blood cell electrolyte indices

<table>
<thead>
<tr>
<th>Indices</th>
<th>Baseline</th>
<th>Control</th>
<th>Oxytocin</th>
<th>Extract</th>
<th>Control</th>
<th>Oxytocin</th>
<th>Extract</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chloride (mM/l)</td>
<td>107.00 ± 7.56(^a)</td>
<td>105.89 ± 8.75(^d)</td>
<td>112.01 ± 8.92(^c)</td>
<td>114.60 ± 9.07(^c)</td>
<td>109.90 ± 11.19(^d)</td>
<td>111.21 ± 13.07(^d)</td>
<td>117.00 ± 9.72(^a)</td>
</tr>
<tr>
<td>Bicarbonate (mM/l)</td>
<td>19.65 ± 1.67(^a)</td>
<td>20.02 ± 2.51(^c)</td>
<td>17.54 ± 1.62(^a)</td>
<td>18.94 ± 1.93(^a)</td>
<td>19.50 ± 1.69(^a)</td>
<td>19.60 ± 2.32(^a)</td>
<td>18.78 ± 1.64(^a)</td>
</tr>
<tr>
<td>Extracellular Na(^+) (mM/l)</td>
<td>125.80 ± 4.82(^a)</td>
<td>126.78 ± 13.56(^a)</td>
<td>121.20 ± 5.26(^c)</td>
<td>135.60 ± 6.15(^a)</td>
<td>126.00 ± 5.43(^c)</td>
<td>125.40 ± 7.13(^c)</td>
<td>138.60 ± 4.10(^a)</td>
</tr>
<tr>
<td>Intracellular Na(^+) (mM/l)</td>
<td>5.91 ± 1.20(^a)</td>
<td>5.62 ± 0.82(^a)</td>
<td>4.81 ± 0.39(^a)</td>
<td>5.26 ± 0.91(^a)</td>
<td>6.06 ± 1.36(^a)</td>
<td>4.86 ± 0.41(^a)</td>
<td>5.56 ± 0.74(^a)</td>
</tr>
<tr>
<td>Extracellular K(^+) (mM/l)</td>
<td>8.47 ± 0.64(^a)</td>
<td>8.74 ± 0.69(^a)</td>
<td>6.10 ± 1.02(^a)</td>
<td>5.26 ± 0.40(^a)</td>
<td>6.86 ± 0.69(^a)</td>
<td>5.96 ± 0.67(^a)</td>
<td>5.48 ± 0.37(^a)</td>
</tr>
<tr>
<td>Intracellular K(^+) (mM/l)</td>
<td>26.30 ± 4.72(^ac)</td>
<td>26.20 ± 5.29(^ac)</td>
<td>18.63 ± 2.65(^c)</td>
<td>27.85 ± 5.70(^c)</td>
<td>27.04 ± 2.94(^ac)</td>
<td>18.30 ± 1.16(^c)</td>
<td>26.65 ± 4.14(^ac)</td>
</tr>
<tr>
<td>Extracellular Ca(^{2+}) (mM/l)</td>
<td>3.41 ± 0.91(^a)</td>
<td>3.31 ± 0.75(^a)</td>
<td>1.26 ± 0.28(^a)</td>
<td>1.03 ± 0.16(^b)</td>
<td>3.70 ± 0.61(^a)</td>
<td>2.03 ± 0.96(^b)</td>
<td>1.65 ± 0.54(^a)</td>
</tr>
<tr>
<td>Intracellular Ca(^{2+}) (mM/l)</td>
<td>0.95 ± 0.10(^a)</td>
<td>0.86 ± 0.12(^a)</td>
<td>4.54 ± 1.05(^a)</td>
<td>4.63 ± 1.31(^b)</td>
<td>0.95 ± 0.07(^a)</td>
<td>4.27 ± 0.69(^b)</td>
<td>3.86 ± 1.04(^a)</td>
</tr>
<tr>
<td>Extracellular Mg(^{2+}) (mM/l)</td>
<td>0.90 ± 0.25(^a)</td>
<td>0.80 ± 0.14(^a)</td>
<td>1.70 ± 0.36(^a)</td>
<td>1.60 ± 0.33(^a)</td>
<td>0.85 ± 0.13(^b)</td>
<td>0.96 ± 0.24(^b)</td>
<td>1.56 ± 0.74(^b)</td>
</tr>
<tr>
<td>Intracellular Mg(^{2+}) (mM/l)</td>
<td>5.10 ± 0.54(^a)</td>
<td>5.19 ± 0.86(^a)</td>
<td>3.90 ± 0.46(^a)</td>
<td>3.30 ± 0.09(^a)</td>
<td>5.05 ± 0.40(^a)</td>
<td>4.51 ± 0.72(^a)</td>
<td>3.19 ± 0.22(^a)</td>
</tr>
</tbody>
</table>

Values are mean ± standard deviation (number of animals per group = 5); values with different superscript(s) in a row are significantly different (p<0.05).
MCV after the first 7 days of extract administration, which however rose non significantly by the 14th day of treatment in comparison with those of the baseline and control values. This later increase may be due to the observed non-significant increase in Hb content of the red blood cells. These beneficial effects of the extract on the animals’ haematological profile, may be attributed to the presence in the plant of flavonoids, vitamin C and other substances with antioxidant properties [16,18], which may help in red cell membrane stabilization [19], as well as presence of Fe required for Hb synthesis. The presence of these antioxidant molecules could possibly explain why the red blood cell integrity was generally unaffected by the extract administration.

Interestingly, oxytocin standard drug administration did not have such positive haematologic potentials like the S. mombin extract. Oxytocin administration significantly reduced Hb concentration and increased WBC count. These effect variations may be because of the absence in this pure standard drug of antioxidant active molecules present in the plant’s extract.

On the other hand, both the extract and oxytocin significantly elicited a time-dependent increase in platelet count of the treated animals. Although, the reason for this increase is not presently known, it may not be unrelated to the reported use of the plant for the control of bleeding especially after childbirth [10]. The presence of appreciable amounts of vitamin C in the plant leaves may also contribute immensely to this potential of bleeding control associated with S. mombin [20].

In conclusion, administration of S. mombin leaf extract does not have significant toxic effects on liver and kidney cellular functions as well as on the haematological indices of rabbits. Rather, it elicited a dose-dependent increase in serum protein and haemoglobin concentrations, reduction in serum bilirubin concentration, enhanced efflux of K+ and influx of Ca2+, and thus a possible activation of cellular cation pumps necessary for initiation of muscle contraction.

References