A research was conducted to investigate the haematological effects of ethanolic leaf extracts of Senna occidentalis on Swiss albino mice infected with Plasmodium berghei. The extracts were found to increase the level of some haematological parameters such as Red Blood Cells, White Blood Cells and Haemoglobin. The effect is concentration dependent, increases with increase in concentration. Thus, the anti-plasmodial efficacy of the leaf extract of S. occidentalis on P. berghei is confirmed. It is therefore recommended that, 400 mg/kg leaf ethanolic extracts of S. occidentalis could be use in the treatment of malarial fever.

Key Words: Albino mice, Ethanolic Extracts, Concentrations, Plasmodium berghei, Senna occidentalis.

INTRODUCTION
Malaria remains an important public health concern in countries where transmission occurs regularly, as well as in areas where transmission has been largely controlled or eliminated (WHO, 2001). Malaria is responsible for one to three million deaths and 300 to 500 million infections annually (Krishna et al., 2009; Verma et al., 2011). Children under five and expecting mothers are particularly susceptible to malaria. The disease claims the life of a child every two minutes (WHO, 2018). The African region continues to bear 90% of malaria cases and 91% of malaria deaths worldwide and Nigeria, the continent’s most populous country, accounted for 27% of malaria cases and 24% of malaria deaths globally in 2016 (WHO, 2018). Plasmodium falciparum the most widespread etiological agent for human malaria has become increasingly resistant to standard antimalarials (such as chloroquine and antifolates) and recently to the Artemisinin combination therapies (ACTs) (Sha’a et al., 2011).

Plants are important source of drugs; especially in traditional medicine (Bako et al., 2005). It is a common practice in Nigeria and other parts of the world to use plant in the form of crude extracts, decoction, infusion or tincture to treat common infection and chronic conditions. According to WHO, over 70% of the world populations rely on medicinal plants for primary health care and there are reports from various researchers on natural substances of plant origin which are biologically active, with desirable antimicrobial and antioxidant properties (Hamid et al., 2010). Medicinal plants have been the focus for the search of new antimalaria drugs in various parts of the world (Schuster, 2001) and the present global situation indicates a recent resurgence in the severity of malaria, due to the resistance of malaria parasites to antimalarial drugs (Peter and Anatoll, 1988). Hence, there is the need to intensify research in the development of new, cheap and effective antimalarial drugs from medicinal plants.
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*Senna occidentalis* is one of the most widely used herbal plants among people of tropical and sub-tropical regions of the world (Veronique and Gabriel, 2013). It is used for various therapeutic purposes in traditional medicine (Yadav et al., 2010, Silva et al., 2011). It has been used to treat malaria since ancient time and surely if it was not effective, malaria would have devastated Africa (Idowu et al., 2010). It is a most common known practice in Nigeria that, the first line of treatment for 60% of children with high fever resulting from malaria is the use of herbal medicine at home (WHO, 2001). The rapid spread of resistance Plasmodium parasite, makes it necessary to search for more effective herbal antimalarial compounds. *Plasmodium berghei* is one of the species of intracellular protozoan parasites from the genus Plasmodium which is responsible for causing malaria. It provides a well established experimental model of malaria infection (Margarida et al., 2006) because of its critical mimicry with the species that infect human (Van der Heyde et al., 2006). The aim of the study is to determine the effect of leaf ethanolic extracts of *Senna occidentalis* on haematological parameters of Swiss albino mice infected with *Plasmodium berghei* with the view of testing the efficacy and safety of the extracts in the treatment and control of Plasmodium parasites.

**MATERIALS AND METHODS**

**Collection and Identification of Samples**

Fresh leaves of *Senna occidentalis* were obtained from the field at Kura town, Kano state in a clean sterile polythene bag and carried to the herbarium section of the Department of Botany, Ahmadu Bello University Zaria for authentification. Voucher number of 900078 was assigned to the specimen.

**Extraction of Plant Ethanolic Extracts**

The extraction methodology followed the protocols of Sasidharan et al. (2011). The collected samples were thoroughly washed and air-dried under shade at room temperature for 2 weeks. The dried leaves were then ground into fine powder with mortar and pestle, and were stored in dry containers until needed. The ethanolic extracts of *S. occidentalis* leaves were prepared by soaking 100 g of each powder in 150 ml of 95% ethanol and shaken in orbital shaker at 120 rpm. The preparations were left to stand for another 24 hours and then filtered through a gauze and then Whatman No 1 filter paper. The filtrates were concentrated to dryness at 40°C under reduced pressure on a rotary evaporator and were stored in a refrigerator at −4°C until the need arise. Different concentrations of 100, 200 and 400mg/kg of the ethanolic extracts were prepared from the leaves of *S. occidentalis*.

**Phytochemical Analysis**

The phytochemical screening of the ethanolic extract of *S. occidentalis* was carried out according to the methods described by Sofowara (1993), Mukherjee (2006) and Adegoke et al. (2010) to determine the presence of active constituents in the plant leaves. Ethanolic extract of *S. occidentalis* was subjected to qualitative test for the presence of bioactive components that include Molisch’s test for detection of Carbohydrates, Meyer’s test for detection of Alkaloids, Wagner’s test for detection of Alkaloids, Lead subacetate test for detection of Tannins, Keller- Killiani’s test for detection of Cardiac Glycosides, Frothing/Foaming test for detection of Saponins, Libermann-Burchard’s test for detection of Steroids and Triterpenoids, Copper Acetate’s test for detection of Diterpenoids, Alkaline test for the detection of Flavonoids, Xanthoproteic test for the detection of Protein and Borntrager’s test for the detection of Anthraquinones.

**Experimental Animals**

Ethical clearance was obtained from Ahmadu Bello University Committee on Animal Use and Care (ABUCAUC). The protocol employed met the guidelines of the Good Laboratory Practice (GLP) regulations of World Health Organization. Apparently healthy white Swiss albino mice of both sexes weighing between 140 and 260 g were used for the work and were obtained from the Faculty of Veterinary Medicine, Ahmadu Bello University, Zaria. The mice were housed in metal cages and kept in a well ventilated room with 12 hours dark/light cycle. They were fed with standard feed pellets and tap water ad libitum. The animals were acclimatized for 2 weeks before proceeding with the experiment.

**Inoculation**

*P. berghei* was obtained from the Department of Parasitology and Entomology, Faculty of Veterinary Medicine, Ahmadu Bello University, Zaria. Parasites were harvested from the blood of a donor rat at peak parasitemia (10⁹ parasites/ml) and was diluted with normal saline and then used for infection of experimental animals. The inoculums consisted of *P. berghei* parasitized erythrocytes. This was prepared by determining both the percentage parasitemia and the erythrocytes count of the donor mouse and diluting the blood with normal saline in proportions indicated by both determinations as described by Akuodor et al. (2010).
Each mouse was inoculated on day 0, intraperitoneally with 0.2ml of infected blood containing approximately 1x10^7 P. berghei parasitized red blood cells. In addition, the newly inoculated animals were monitored daily to determine their degree of infestation with the parasite.

A 4-day suppressive test was conducted according to the method described by Mbah et al. (2012). After four hours of inoculation, the infected mice were randomly divided into 5 groups of 3 mice per cage and were treated for four consecutive days. Group 1 received 0.2 mL of normal saline (drug free control). Group 2, 3, 4 received 100, 200 and 400mg/kg of the ethanol leaf extract respectively while group 5 received 10mg/kg of chloroquine phosphate. All doses were administered orally using oral cannula according to their weight. On the fifth day, thin film of blood was collected from the tail of each mouse and thick films fixed with methanol, stained with 4% Giemsa at pH 7.2 for 30 min were prepared. Parasitemia was determined by counting the parasitised red blood cells out of 1000 red blood cell in 10 random macroscopic fields. Slides were viewed using light microscopy with oil immersion (X1000 magnification). The percentage of infected RBCs was determined by enumerating the number of infected RBCs in relation to the number of uninfected RBCs as described by Akuodor et al. (2010). A minimum of 500 RBCs total was counted.

\[
\text{Percentage of Infected RBCs} = \frac{\text{Number of infected RBCs}}{\text{Total Number of RBCs Counted}} \times 100
\]

Average chemo suppression was calculated as

\[
\% \text{ Suppression} = \frac{\text{Average parasitemia in Control} - \text{Average parasitemia in Treated}}{\text{Average parasitemia in Control}} \times 100
\]

**Acute Toxicity Test**

The safety of the extract to the mice was determined by estimating the lethal concentration (LC50) of the ethanolic leaf extract of *S. occidentalis* using the method as described by Lorke (1983) with modification. Mice of both sexes were starved for 8 hours prior to the toxicity test. The study was carried out. The mice were administered 100, 200 and 400mg/kg of the leaf extract. The acute toxicity LC50 was calculated as geometric mean of the concentration that resulted in 100% lethality and that which caused no lethality at all. Toxicity signs such as death, changes in physical appearance or behavioral changes were observed for over a period 72 hours after administration of the plant extract.

Packed cell volume (PCV) was measured to predict the effectiveness of the test extract in preventing haemolysis resulting from increased parasitaemia. Heparinized capillary tubes were used for collection of blood from tail of each mouse. Blood sample collected was suspended in EDTA to prevent blood coagulation. The capillary tubes were filled with blood up to three-third of their volume and sealed at the dry end with sealing clay. The tubes were then placed in a micro-haematocrit centrifuge (Gelma Awhksley, England) with the sealed end outwards and centrifuged for 5 min at 12,000 rpm. The tubes were then taken out of the centrifuge and PCV was determined using a standard micro-haematocrit reader using the protocols of Dacies and Lewis (1975). Haemoglobin level was determined by the method of Crosby et al. (1954) as modified by Pla and Fritz (1971). Estimation of white and red blood cell counts were computed using the method described by Dacies and Lewis (1975).

The formula for computing the PCV follows the following relationship:

\[
\text{PCV} = \frac{\text{Volume of Erythrocytes in a Given Volume of Blood}}{\text{Total Blood Volume}}
\]

**Results**

The result for the phytochemical screening of the leaf extracts of *S. occidentalis* is presented in Table 1. The result revealed the presence of ten active constituents in the form of alkaloids, carbohydrates, cardiac glycosides, diterpenoids, flavonoids, proteins, saponins, steroids, tannins and triterpenoids.
Table 1: Phytochemical Constituents of the Leaves of *S. occidentalis*

<table>
<thead>
<tr>
<th>S/N</th>
<th>Phytochemical Compound</th>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Alkaloids</td>
<td>Wagner’s test</td>
<td>+</td>
</tr>
<tr>
<td>2</td>
<td>Anthraquinones</td>
<td>Bornfrager’s</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>Carbohydrate</td>
<td>Molisch</td>
<td>+</td>
</tr>
<tr>
<td>4</td>
<td>Cardiac glycosides</td>
<td>Keller-Killani</td>
<td>+</td>
</tr>
<tr>
<td>5</td>
<td>Diterpenoids</td>
<td>Copper acetate</td>
<td>+</td>
</tr>
<tr>
<td>6</td>
<td>Flavonoids</td>
<td>Alkaline reagent test</td>
<td>+</td>
</tr>
<tr>
<td>7</td>
<td>Protein</td>
<td>Xanthoproteic</td>
<td>+</td>
</tr>
<tr>
<td>8</td>
<td>Saponins</td>
<td>Frothing test</td>
<td>+</td>
</tr>
<tr>
<td>9</td>
<td>Steroids</td>
<td>Libermann-Buchard’s test</td>
<td>+</td>
</tr>
<tr>
<td>10</td>
<td>Tannins</td>
<td>Lead subacetate test</td>
<td>+</td>
</tr>
<tr>
<td>11</td>
<td>Triterpenoids</td>
<td>Libermann-Buchard’s test</td>
<td>+</td>
</tr>
</tbody>
</table>

Key: (+) = Present  (-) = Absent

The result for acute toxicity test revealed no mortality observed in mice after oral administration of the ethanolic leaf extracts and the LC₅₀ value of 1000 mg/kg was estimated which signifies that the oral LC₅₀ was greater than 1000mg/kg. There were no observed clinical signs of toxicity in the treated mice during a 4-day observation period except mild drowsiness and mild weakness in the treated mice.

However, the result for the *in vivo* assays for four-day suppressive effect is presented in Table 2. The result for the percentage suppression analysis of the leaf ethanolic extracts of *S. occidentalis* revealed significant decrease (P≤0.05) in parasitaemia at all dose levels as compared to the negative control group. The group that received 400 mg/kg of the extract exhibited maximal suppression (75.00%) of parasitaemia than the other groups. All doses of the extract significantly enhanced the survival time of the mice in a dose dependent manner as compared to the negative control group (Table 2). The percentage suppression increases with increase in concentration of the extracts.

Table 2: Effect of Leaf Ethanolic Extract of *S. occidentalis* on *P. berghei* infected mice on suppressive test

<table>
<thead>
<tr>
<th>S/N</th>
<th>Dose (mg/kg)</th>
<th>Parasitemia Level/µl of Blood</th>
<th>% Suppression</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Control I</td>
<td>4.20±0.21a</td>
<td>-</td>
</tr>
<tr>
<td>2</td>
<td>Control II</td>
<td>4.10±0.40a</td>
<td>2.00</td>
</tr>
<tr>
<td>3</td>
<td>100</td>
<td>2.70±0.1b</td>
<td>35.00</td>
</tr>
<tr>
<td>4</td>
<td>200</td>
<td>1.83±0.55c</td>
<td>56.00</td>
</tr>
<tr>
<td>5</td>
<td>400</td>
<td>1.03±0.03d</td>
<td>75.00</td>
</tr>
</tbody>
</table>

Key: Control I: 0.2 mL Normal Saline, Control II: 10 mg/kg of Chloroquine

More so, the result for the effect of different concentrations of the extracts on haematological characteristics of the experimental animals is shown in Table 3. The result on the Packed Cell Volume of the Swiss albino mice infected with *Plasmodium berghei* and treated with different concentrations of the extracts showed significant increase (P≤0.05) in PCV level with increase in the concentrations of the extracts. Similar results were found in the haemoglobin and Red Blood Cells levels. The levels of PCV, Hb and RBC decreased among the positive and negative controls. However, 100 mg/kg of the extracts showed increased in the levels of White blood cells while 400 mg/kg and the negative control inducing the least.

Table 3: Effect of leaf extract on some haematological parameters

<table>
<thead>
<tr>
<th>Parameters</th>
<th>0.2ml Normal saline</th>
<th>10 mg/kg Chloroquine</th>
<th>100mg/kg</th>
<th>200mg/kg</th>
<th>400mg/kg</th>
</tr>
</thead>
<tbody>
<tr>
<td>PCV</td>
<td>31.30±0.88d</td>
<td>42.00±7.02c</td>
<td>42.00±6.56c</td>
<td>45.67±1.20b</td>
<td>56.67±2.40a</td>
</tr>
<tr>
<td>Hb</td>
<td>10.50±0.39d</td>
<td>13.97±2.33c</td>
<td>13.63±1.86c</td>
<td>15.20±0.42b</td>
<td>17.87±1.94a</td>
</tr>
<tr>
<td>RBC</td>
<td>5.70±0.38d</td>
<td>7.17±1.17c</td>
<td>7.01±0.92c</td>
<td>7.83±0.50b</td>
<td>9.33±0.48a</td>
</tr>
<tr>
<td>WBC</td>
<td>8.97±1.79c</td>
<td>7.67±1.73d</td>
<td>13.97±2.03b</td>
<td>16.93±1.22a</td>
<td>7.87±1.27d</td>
</tr>
</tbody>
</table>
DISCUSSION

The presence of vast array of phytochemicals in the leaves of *S. occidentalis* signifies the relative importance of this plant in the treatment of *P. berghei* infections. This finding is in agreement with that of Saidu *et al.* (2011) who reported that, the efficacy of *S. occidentalis* leaf extracts in the treatment of *Salmonella typhi* infections depends on the phytochemical constituents of the extracts. The fact that, malarial parasites depend on blood and blood system, analysis of the effects of such parasites and efficacy of the extracts on blood parameters is critical in understanding the mechanism of infection and control. It is a known fact that malaria infection leads to anaemia as reported by Balogun *et al.* (2009). More so, the growing parasite consumes and degrades the intracellular proteins which are mainly hemoglobin as stressed by Gavigan *et al.* (2001). This may account for further reduction in Hemoglobin level. These reductions were considerably reversed in the infected mice treated with the extracts. This suggests that the extract may enhance the production of red blood cells. This subsequently causes an increase in PCV and Hb levels observed in extract treated mice. However, the decrease in the RBC, WBC and Hb level exposes the animal to several potential health hazards such as anaemia, low immune system and suffocation. Decrease in WBC level in infected mice leaves the mice at great risk of having infections. The observed increase in WBC in the infected mice group treated with 100mg/kg of the leaf extract may probably be as a result of stimulation of the immune system of the mice to fight malaria parasites as reported by Yakubu *et al.* (2007). The ethanolic extract has the ability of boosting the immune-response of the infected mice. Thus, the plasmodium infections affect the intestine at the immunological level when the immune system is low as reported by Wang *et al.* (2013).

In acute toxicity test, no adverse effect or mortality was observed in the mice up to during the 72 h observation. This finding is in conformity with the previous report contained in the guideline set by Organization for Economic Co-operation and Development, OECD (2001) and Akhilia *et al.* (2007) that these plants are not toxic and safe for use. This finding is also in conformity with the earlier work of Ayoka *et al.* (2005) and Uchendu and Isek (2008) who individually reported similar findings using *Spondias mombin* extracts. Similar finding was also reported by Sini *et al.* (2010) using *Cassia occidentalis* extracts. The *S. occidentalis* extracts show significant effects on the the haematological parameters of infected mice. The extracts displayed erythropoiesis as indicated by significant increase in the PCV (42.00-56.67%) compared to the controls (31.30-42.00%). Erythropoiesis is an increase in the level of red blood cells than normal compared to the whole blood. This finding is in conformity with the previous work of Taylor (2006) who reported similar finding using *Spondias mombin* extracts and that of Yadav *et al.* (2009) who reported significant role played by *Cassia occidentalis* extracts in increasing the levels of haematological parameters. Similarly, Hamenoo (2010) reported increased in the levels of haematological parameters due to *S. mombin* extracts administration in rodents. Similarly, the haematological parameters of all the test animals treated with the leaf extracts of *C. occidentalis* increased significantly. This finding corresponds to that of Nuhu and Aliyu (2008) who reported significant increase in the level of haematological parameters of *C. occidentalis* extracts treated rats. More so, Emeka and Funmilayo (2011) reported similar finding among alloxan-induced diabetic rats treated with the extracts of *S. mombin* and *Parinari polyandra*. More so, the acute toxicity assessment of the extracts on to the test animals shows no gross physical and behavioral changes such as anorexia, diarrhea, depression, abnormal secretion and hair erection for 24 h. All the mice survived the 2-week observation period. The direct toxic effects of the crude extract on blood parameter (PCV) in healthy mice were evaluated by administering higher doses of the extract.

CONCLUSION

It was concluded that, the leaf ethanolic extracts of *Senna occidentalis* significantly reduce paracetaemia due to *P. berghei* infection and did not show any toxicity effect. Thus, the extracts within the range of 100-400 mg/kg is safe for consumption and effective against malaria. It was therefore recommended that, 400 mg/kg leaf ethanolic extracts of *Senna occidentalis* is the most effective dose in the treatment of malarial infection caused by *P. berghei*.

Acknowledgements

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