Effect of Ethanol Extracts of *Musa paradisiaca* Fruit Pulp and Peels on Haematological Indices and Liver Enzymes of Experimental Rats

Emuesiri Goodies Moke1,*, Emuesiri Kohworho Umukoro1, Evelyn Tarela Ojugbemi2, Theresa Ezedom2, Tarela Melish Elias Daubry3, Iziege Lisa Omorodion1

1Department of Pharmacology and Therapeutics; 2Department of Medical Biochemistry; 3Department of Physiology, Faculty of Basic Medical Sciences, Delta State University, Abraka, Nigeria

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INTRODUCTION

Plants possess the ability to synthesize a variety of chemical compounds with various biological functions. The plant kingdom remains a harvest for many species of plants with key medicinal value which are yet to be discovered. Reports show that a large population (80%) of people in developing countries depend primarily on medicinal plants for their primary health care (Mahomoodally, 2013; Ekor, 2014). These plants and herbs are taken in different ways and forms (whole or in parts) for the alternative management of diseases traditionally (Benzie and Wachtel-Galor, 2011; Emudainohwo et al., 2015; Moke et al., 2019; Okafo et al., 2019).

*Musa paradisiaca* belongs to Musaceae family and is popularly known as plantain. The plant is widely distributed in the Southern part of Nigeria, West and East Africa, Malaysia, Cameroun and Southern parts of United States (Uwaoma, 2003; Nayar, 2010). *Musa paradisiaca* (Plantain) is an important staple crop that contributes to the calories and substance economic in Africa (Oyeyinka and Afolayan, 2019). Abundant medicinal activities of parts of *Musa paradisiaca* have been reported. It has been shown to possess hepatoprotective activity (Nirmala et al, 2012; Issa et al., 2018) as well as antidiabetic (Vilhena et al., 2020), antiulcer (Onasandwo et al., 2013; Ezekwesili et al., 2014; Moke et al., 2017), antimicrobial (Fagbemi et al., 2009), wound healing (Agarwal et al. 2009), and antioxidant properties (Yin et al., 2008).

The present study is aimed at evaluating the effect of parts of the ethanol fruit extracts of *Musa paradisiaca* on haematological indices and serum liver enzymes.

MATERIAL AND METHODS

Plant Collection and Preparation

*Musa paradisiaca* fruits (unripe) were purchased locally from the market, and were identified and authenticated by a taxonomist with existing specimen deposited at the herbarium of the Department of Botany, Delta State University, Abraka, Nigeria. The fruits were rinsed with water, and both the fruit pulps and fruit peels were air dried. The dried pulps and peels were grounded separately into pulverized powder using a grinding machine for ease of extraction.

The powdered materials of *Musa paradisiaca* fruits (400 g each) were separately extracted exhaustively with ethanol using Soxhlet extractor at 25 ºC. The filtrates were concentrated using Rotary evaporator at 40 ºC. The percentage yields were 8.4% (fruit pulp) and 9.73% (fruit peel). The filtrates were concentrated using Rotary evaporator at 40 ºC. The percentage yields were 8.4% (fruit pulp) and 9.73% (fruit peel). The filtrates were concentrated using Rotary evaporator at 40 ºC. The percentage yields were 8.4% (fruit pulp) and 9.73% (fruit peel).
(fruit peel). The concentrated ethanol extracts were refrigerated prior to use.

Animals
Wistar rats (150 – 180 g) were obtained from the Animals’ House facility of the Faculty. The animals were acclimatized for 7 days prior to the study, and were fed with rat feed and clean water ad libitum. Guidelines followed in the handling of animals were in accordance with the ethical standards of the Institutional Animals Ethics Committee (IAEC), as adopted by the ethical committee of the Faculty of Basic Medical Science, Delta State University, Abraka, Nigeria.

Experimental Design
The animals were divided into five (5) groups of six animals each:
- Group 1 – Normal saline (Control) 10 ml/kg
- Group 2 – *Musa paradisiaca* fruit pulp extract (MPF Pulp) 200 mg/kg
- Group 3 – *Musa paradisiaca* fruit pulp extract (MPF Pulp) 400 mg/kg
- Group 4 – *Musa paradisiaca* fruit peel extract (MPF Peel) 200 mg/kg
- Group 5 – *Musa paradisiaca* fruit peel extract (MPF Peel) 400 mg/kg

The experimental animals were administered the extracts orally daily for 14 days according to their body weights.

Sample Collection
At the end of the 14-days treatment period, the animals were anesthetized using chloroform. Blood samples were collected by cardiac puncture into labeled EDTA bottles for haematological analysis and liver function tests.

Determination of Haematological Indices
The method as described by Tietz (1976) and Baker et al. (1998) were used for determining the red blood cells (RBC) counts, haemoglobin (Hb) concentration, and haematocrit level.

Determination of Liver Function Test
Alkaline phosphatase (ALP), aspartate aminotransferase (AST), and alanine transaminase (ALT) in serum were determined according to methods described by Reitman and Frankel (1957) and Roy (1970).

Statistical Analysis
Results are presented as the mean ± standard error of the mean (SEM). Data were analysed using one-way analysis of variance (ANOVA) followed by Tukey’s post hoc test. P-values < 0.05 were taken as significant.

RESULTS AND DISCUSSION
Effect of Ethanol Fruit Extracts of *Musa Paradisiaca* on Haematological Indices of Wistar rats
Figures 1-3 depict the effect of ethanol fruit extracts (pulp and peel) of *Musa paradisiaca* on the red blood cell count, haematocrit level, and haemoglobin concentration of normal Wistar rat. MPF Pulp and MPF Peel at a dose of 200 mg/kg had a non-significant (P>0.05) increase in red blood cell count when compared to the control, however, at 400 mg/kg, there was a significant (P<0.05) increase in red cell count. High dose (400 mg/kg) of both fruit extracts significantly (P<0.05) increased the haematocrit level as compared to the control. There was a non-significant (P>0.05) increase in haemoglobin concentration in the treated rats as compared to the control.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Red Blood Cell (RBC) Count (× 10^12/L)</th>
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<tbody>
<tr>
<td>Control</td>
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<tr>
<td>MPF Pulp 200</td>
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<td>MPF Pulp 400</td>
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<td>MPF Peel 200</td>
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<td>MPF Peel 400</td>
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Figure 1. The effect of ethanol fruit extracts of *Musa paradisiaca* on red blood cell count of Wistar rats.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Haematocrit (%)</th>
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<tr>
<td>Control</td>
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<td>MPF Pulp 200</td>
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<td>MPF Pulp 400</td>
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Figure 2. The effect of ethanol fruit extracts of *Musa paradisiaca* on haematocrit level of Wistar rats.
Blood is composed of a variety of living cells that circulate through the heart and the blood vessels carrying nutrients, hormones, vitamins, antibodies, heat and oxygen to the body’s tissue. The components of blood include red blood cells, white blood cells and platelets which are suspended in plasma (Basu and Kulkami, 2014). Red cell contains haemoglobin, a protein that carries oxygen to all the tissues of the body. Haematocrit or packed cell volume is a measurement of the proportion of blood that is made up of cells. Following centrifugation, it is an estimate of the ratio of the volume of red blood cells to the total volume of blood (Mondal and Budh, 2020). Haematocrit and haemoglobin values are useful for assessing anaemia, polycythemia, and also for estimating response to treatment (Northrop-Clewes and Thurham, 2013; White, 2018; Mondal and Budh, 2020). Hematopoiesis is the process involved in the formation of blood cells (Rieger and Schroeder, 2012). This study showed the positive effect of the fruit pulp and peel of Musa paradisiaca on hematopoiesis. Musa paradisiaca was also revealed to have no toxic effect on liver enzymes.

The assessment of haemotological parameters could be used to reveal the deleterious effect of foreign compounds including plant extract on the blood constituent of animals. They can also be used to determine possible alteration in the levels of biomolecules, metabolic products, as well as histomorphology of the organs (Magalhães et al., 2008). Following the administration of the extracts, there was an increase in red cells count and haematocrit level, which was significant at a higher dose of 400 mg/kg as compared to the control group. MPF pulp had as better increase in red cell count when compared with MPF peel, whereas MPF peel showed a much effect in increasing the haemotocrit level as compared to MPF pulp. The results also revealed an increase in haemoglobin concentration (Figures 1-3). These increments indicate that M. paradisiaca contains phytochemicals that stimulate the synthesis of erythrocytes possibly by stimulating erythropoietin formation and secretion. Erythropoietin is a glycoprotein hormone which stimulates stem cells in the bone marrow to produce red blood cells (Ohlsson and Aher, 2009).

**Effect of Ethanol Fruit Extracts of Musa Paradisiaca on Serum Liver Enzymes of Wistar rats**

Figure 4 shows the effect of ethanol fruit extracts (pulp and peel) of Musa paradisiaca on the liver enzymes of normal Wistar rat. There was a non-significant (P>0.05) change in AST, ALT, and ALP level of the treated rats as compared to the control.

An assessment of the effect of ethanol fruit extracts of Musa paradisiaca on liver enzymes of Wistar rats showed a non-significant change in serum aspartate aminotransferase (AST), alanine aminotransferase (ALT) and serum alkaline phosphatase (ALP) levels of the treated rats as compared to the control (Figure 4). Serum AST, ALT, and ALP, which are cytoplasmic enzymes released into circulation after cellular damage are useful enzymes biomarkers in predicting liver damage (Ramaiah 2011; Zhao et al., 2018). ALT and AST are largely used in the assessment of liver damage by drugs or any other hepatotoxins, while ALP is a marker enzyme for the plasma membrane and endoplasmic reticulum (Giannini et al., 2005; McGill, 2016). The observed non-significant differences in the liver enzymes are an indication that both fruit pulp and peels extracts of Musa paradisiaca are non-toxic to the hepatic cells, thus, suggesting that Musa paradisiaca fruit extracts may not possess hepatotoxic effects, perhaps, a protective effect by stabilization of plasma membrane thereby preserving the structural integrity of the cell (Pari and Murugen, 2004). This is corroborated by the findings of Iweala et al (2011) which reported significantly reduced liver enzymes level with the consumption of a Musa paradisiaca-supplemented diet by Wistar rats. The hepatoprotective properties of Musa

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**Figure 3.** The effect of ethanol fruit extracts of Musa paradisiaca on haemoglobin concentration of Wistar rats.

**Figure 4.** The effect of ethanol fruit extracts of Musa paradisiaca on liver enzymes of Wistar rats.
paradisiaca against experimentally induced hepatotoxic models have also been reported (Nirmala et al., 2012)

CONCLUSION
This study evaluated the effect of ethanol fruit extracts of Musa paradisiaca on haematological indices and serum liver enzymes. Fruit pulp and peel of Musa paradisiaca improve erythrocytes count and haematocrit level, and they may not be associated with liver toxicity. Fruits of Musa paradisiaca can be used as food therapy in raising red cells synthesis in anemic conditions.

Conflict of Interest: The authors declare that there are no conflicts of interest concerning the publication of this article.

REFERENCES


