Effect of methanol, n-hexane and aqueous extract of *Irvingia gabonensis* leaf on castor oil-induced diarrhoea in albino rats

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**ABSTRACT**

Bush mango leaf (*Irvingia gabonensis*) is commonly used locally to treat diarrhoea. The present study evaluated the anti-diarrhoea effect of this plant extract on albino rats induced with castor oil. Fresh tender leaf of this plant was collected, air-dried, powdered and percolated in n-hexane, methanol and aqueous solvents. Diarrhoea was induced with castor oil on albino rats using standard procedures. The extract (100, 250 and 500 mg/kg) in a dose dependent manner, significantly protected the rat against diarrhoea induced experimentally with castor oil in terms of nature and number of stool droppings within 3 hrs of administration of extract, there was no significant difference when compared with the control drug (Loperamide) (P > 0.05). The study justifies the ethno pharmacological uses of this medicinal plant for treatment of diarrhoeal disease.

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**Keywords:** Diarrhoea, castor oil, *Irvingia gabonensis*.

**INTRODUCTION**

Diarrhoea can be defined as the increased frequency of bowel movements, accompanied by a loose consistency of stools. Worldwide distribution of diarrhoea accounts for more than 5-8 million deaths each year in infants and small children less than 5 years of age. World health organization (WHO) estimate that in 1998 (Jaiarj et al., 2000), about 7.1 million deaths occurred due to diarrhoea (Lin et al., 2002). On a worldwide basis, 750 million cases are reported in children below 5 years in Asia, Latin America and Africa resulting in 4-5 million deaths (Abdullahi et al., 2001).

However, the incidence of diarrhoeal diseases varies greatly with the seasons and a child’s age. The youngest children are most vulnerable. Incidence is highest in the first two years of life and declines as a child grows older. Mortality from diarrhoea has declined over the past two decades from an estimated 5 million deaths among children under 5 years to 1.5 million deaths in 2004 (WHO, 2004).
In an effort to tackle the problems of diarrhoea, the world Health Organization (WHO) has established diarrhoea disease control programme (DDC) which includes studies of traditional medicinal practices together with the evaluation of health education and prevention approaches (3,4). In most parts of the developing countries, particularly Africa, the use of herbal remedies in management of diarrhoea is a common practice. *Irvingia gabonensis* is a tree plant popularly known as wild or Africa mango. The plant occurs freely in many parts of Africa. It belongs to the family Irvingiaceae, It is identified by various names such as “bush mango”, “dika nut” tree, “Ugiri” in Igbo, “Goron” or “Biri” in Hausa and “Apon” in Yoruba. The variability of the phytochemical compounds of each plant may explain why traditional medicine uses the combination of both plants to increase the antioxidant activities. (Koevi et al., 2015).

They have been proved to contain pharmacologically active ingredients which have over the years been exploited in traditional medicine for the treatment of various human and animal diseases (Adamu et al., 2006).

Recently, Raji et al. (2001) reported anti-diarrhoeal and anti-ulcer properties of *I. gabonensis*. Okorondu et al. (2013) observed that the inhibitory action of the plant extracts could be attributed to the presence of the phytochemical constituents in the plant extracts such as alkaloid, flavonoid and saponin. Castor oil is a ricinoleate (Mckeon et al., 1999) and its diarrhoea inducing property is known to be due to its active metabolite ricinoleic acid which increases peristaltic activity and alters permeability of the intestinal mucosa membrane to electrolytes. The objective of the study was to determine the therapeutic effect of the plant extract in rats induced with castor oil.

**MATERIALS AND METHODS**

**Plant collection**

*Irvingia gabonensis* leaf was collected from Lilu town in Ihiala L.G.A of Anambra State, Nigeria. The plant was identified and authenticated in the Department of botany, Nnamdi Azikiwe University, Awka Nigeria where the sample was deposited. The leaves spread out and dried on a clean surface under a shade at room temperature to exclude direct sunlight in order to prevent the active constituents of the leaves from being degraded due to photochemical reactions. It was air dried for about eight days after which, it was observed to be dried. The dried leaves were gathered, and crushed with grinder. The powder was weighed using an electric weighing balance by Kern ALS 220 – 4. The powder was then stored in an air tight bag at room temperature and used for further extraction.

**Preparation of plant extract**

The ground leaf was prepared in three ways to get the extracts.

**Aqueous extract (Maceration Method)**

Maceration method was used for aqueous extraction and powdered leaf of *Irvingia gabonensis* was used. 150 g of the plant was weighed and put in 375 ml of distilled water and allowed to stand for 48 hrs, agitation or shake for 45 mins. The extract was filtered using British standard mesh filter and first muslin cloth and concentrated by using air drying under constant air current and water bath at 50 °C. The extract was then transferred into a clean container and stored in the refrigerator until required for use.

**Organic solvent extraction by maceration**

This was carried out at Pharmacognosy Department, Faculty of Pharmaceutical Sciences, Agulu. 150 g of the plant sample was transferred into 1000 ml volumetric flask, and then 375 ml of solvent (methanol and n-hexane) were added. This was covered and allowed for 48 hrs with continuous shaking, filtered and transferred to rotary evaporator for concentration. The extract was then transferred into a clean container and stored in the refrigerator until required for use.

**Extraction by Soxhlet method**

This method was carried out by continuously extracting a sample with a non-polar organic solvent for about 4-6 hrs.
**Experimental animal and diet**

In the study, thirty-three (33) of about 8 weeks old healthy rats (*Rattus norvegicus*) weighing 92 – 138 g were used. The rats were kept in standard metal cages obtained from the faculty. Controlled conditions of temperature (25 ± 20 °C), relative humidity (50 ± 15%) and normal photoperiod (12 – 12 hrs light – dark cycle) were maintained. Free access to standard pellet and water was ensured. Acclimatization of the rats to the metal cages (4 – 6 rats/cage) was done for seven days and maintained according to the NIH guideline for care and use of laboratory Animals (Saha et al., 2001).

Drug: Loperamide (2 mg) was obtained from Pharmacy Department, Nnamdi Azikiwe University Hospital, Nnewi, Anambra State, Nigeria.

**Anti-diarrhoea assay**

The plant extract with three different solvents (n-hexane, aqueous and methanol) were used for the study. Eleven (11) cages allotted three albino rats each, out of the eleven cages, two were used as negative control and positive control, and nine cages were used as test groups.

The animals were fasted for 12 hrs and weighed. 2 ml of castor oil were given intragastrically to each rat in each group to induced diarrhoea (Yegnanarayan and Ghoshti 1982), while papers were dropped underneath the cages. Faecal materials were checked for the nature of stool passage; 30 min after administration of castor oil, diarrhoea was established, 2 ml of water was given to negative control and 2 ml of anti-diarrhoea (Loperamide) was given to positive control and volume equivalent of plant extract (100, 250 and 500 mg/kg) were given to test groups. Faecal materials were checked for consistency/frequency every 30 min for the characteristic droppings. The presence/absence of fluid material in the stool that stained the absorbent paper placed beneath the cages and nature of stool passed during 3 hrs period was recorded to determine the effect of plant extract on castor oil induced diarrhoea. Their absence was recorded as a good protection from diarrhoea; also reduction in diarrhoea (watery Stool) is a protection.

**Statistical analysis**

The statistical method used was 2- way ANOVA and Bonferroni post-test. This was done at P < 0.05 level of significance.

**RESULTS**

Tables 1 – 3 show the results of effect of methanol, n-hexane and aqueous extracts of *Irvingia gabonensis* leaf on castor oil induced diarrhoea. After administration of castor oil, the rats were defecating watery stools (ws), then after administration of extracts, the watery stooling stopped. The methanol and n-hexane extracts of *Irvingia gabonensis* show that in high doses (500 mg/kg), the number of stool droppings within 3 hrs of administration of extract was one (1) while the number of no stool dropping (No defecation) within 3 hrs of administration of extract was five (5), when compared with the anti-diarrhoea drug (loperamide) (P < 0.05) that had the same result. Then, the aqueous extract of *Irvingia gabonensis* shows that in high dose (500 mg/kg), the number of stool droppings within 3 hrs of administration of extract was two (2). In other ways, the methanol and n-hexane extracts of *Irvingia gabonensis* were more effective than the aqueous extract of *Irvingia gabonensis* at high doses (500 mg/kg). whereas, the aqueous and n-hexane extracts had the same number of stool dropping (2) and the number of no stool dropping (4) within 3 hrs at low (100 mg/kg) and middle (250 mg/kg) doses (Tables 2 and 3).

The Methanol extract of *Irvingia gabonensis* had the same result at middle dose (250 mg/kg) with anti-diarrhoea drug (loperamide) (Table 1).

The extracts significantly protected rats against castor oil- induced diarrhoea when compared with the control as evident in the nature and prolongation of onset. Loperamide (2 mg/kg) also significantly protected rats against castor oil-induces diarrhoea. The extracts effect was dose dependent (100, 250 and 500 mg/kg).
Table 1: Effect of methanol extract of *Irvingia gabonensis* on castor oil-induced diarrhoea.

<table>
<thead>
<tr>
<th>Nature of stool, 30 min after administration of castor oil</th>
<th>Extract doses (mg/kg)</th>
<th>Weight (g)</th>
<th>Volume Equivalent to dose administered (ml)</th>
<th>Nature of stool at time interval (30 minutes)</th>
<th>Number of stool dropping within 3 hrs of adm. of extract</th>
<th>Number of no stool dropping within 3 hrs</th>
</tr>
</thead>
<tbody>
<tr>
<td>Watery stool</td>
<td>100</td>
<td>125</td>
<td>0.13</td>
<td>SWS ND ND ND ND NS</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Watery stool</td>
<td>250</td>
<td>119</td>
<td>0.29</td>
<td>ND ND ND ND ND NS</td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Watery stool</td>
<td>500</td>
<td>118</td>
<td>0.59</td>
<td>ND ND ND ND ND NS</td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Watery stool</td>
<td>Loperamide</td>
<td>121</td>
<td>0.41</td>
<td>ND ND ND ND ND NS</td>
<td>1</td>
<td>5</td>
</tr>
</tbody>
</table>

WS = watery stool; SWS = semi-watery stool; HS = Hard stool; NS = Normal stool; ND = No defecation.

Table 2: Effect of aqueous extract of *Irvingia gabonensis* on castor oil-induced diarrhoea.

<table>
<thead>
<tr>
<th>Nature of stool, 30 min after administration of castor oil</th>
<th>Extract doses (mg/kg)</th>
<th>Weight (g)</th>
<th>Volume Equivalent to dose administered (ml)</th>
<th>Nature of stool at time interval (30 minutes)</th>
<th>Number of stool dropping within 3 hrs of adm. of extract</th>
<th>Number of no stool dropping within 3 hrs</th>
</tr>
</thead>
<tbody>
<tr>
<td>Watery stool</td>
<td>100</td>
<td>118</td>
<td>0.11</td>
<td>ND ND ND ND ND NS</td>
<td>2</td>
<td>4</td>
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<tr>
<td>Watery stool</td>
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<td>129</td>
<td>0.32</td>
<td>NS ND ND ND ND NS</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Watery stool</td>
<td>500</td>
<td>119</td>
<td>0.59</td>
<td>NS ND ND ND ND NS</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Watery stool</td>
<td>Loperamide</td>
<td>121</td>
<td>0.41</td>
<td>ND ND ND ND ND NS</td>
<td>1</td>
<td>5</td>
</tr>
</tbody>
</table>

WS = watery stool; SWS = semi-watery stool; HS = Hard stool; NS = Normal stool; ND = No defecation.

Table 3: Effect of n-hexane extract of *Irvingia gabonensis* on castor oil-induced diarrhoea.

<table>
<thead>
<tr>
<th>Nature of stool, 30 min after administration of castor oil</th>
<th>Extract doses (mg/kg)</th>
<th>Weight (g)</th>
<th>Volume Equivalent to dose administered (ml)</th>
<th>Nature of stool at time interval (30 minutes)</th>
<th>Number of stool dropping within 3 hrs of adm. of extract</th>
<th>Number of no stool dropping within 3 hrs</th>
</tr>
</thead>
<tbody>
<tr>
<td>Watery stool</td>
<td>100</td>
<td>138</td>
<td>0.14</td>
<td>ND ND ND ND ND NS</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Watery stool</td>
<td>250</td>
<td>113</td>
<td>0.28</td>
<td>NS ND ND ND ND NS</td>
<td>2</td>
<td>4</td>
</tr>
<tr>
<td>Watery stool</td>
<td>500</td>
<td>113</td>
<td>0.59</td>
<td>ND ND ND ND ND HS</td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Watery stool</td>
<td>Loperamide</td>
<td>121</td>
<td>0.41</td>
<td>ND ND ND ND ND HS</td>
<td>1</td>
<td>5</td>
</tr>
</tbody>
</table>

WS = watery stool; SWS = semi-watery stool; HS = Hard stool; NS = Normal stool; ND = No defecation.
DISCUSSION

The results of the present study suggested that the methanol, n-hexane and aqueous extracts of *Irvingia gabonensis* had significant anti-diarrhoea effects. Castor oil is ricinoleate (McKeon et al., 1999) and its diarrhoea inducing property is known to be due to its active metabolite ricinoleic acid which increases peristaltic activity and alters permeability of the intestinal mucosa membrane to electrolyte, particularly Na⁺, Cl⁻ (Adzu et al., 2003) and water (Capasso et al., 1994).

Castor oil is also reported to induce diarrhoea by increasing the volume of intestinal content by prevention of the reabsorption of water.

The secretory diarrhoea is associated with an activation of Cl⁻ channels, causing Cl⁻ efflux from the cell, the efflux of Cl⁻ results in massive secretion of water into the intestinal lumen and profuse watery diarrhoea (Longanga et al., 2000).

Diarrhoea was established 30 mins after administration of castor oil to the rats at volume equivalent to their weights. The results revealed that the extract was able to stop the diarrhoea in the rats within 3 hrs of therapy, showing that they exhibited anti-diarrhoeal effects. Effect of the extracts on castor oil induced diarrhoea in rat also showed protection ability of the extract was high. Abdulrahman et al. (2004) reported the same. The above findings are therefore, suggestive of non-specific mechanism of action of anti-diarrhoeal activity of the extract. Moreso Unaeze et al. (2017) also suggested that the plant had anti diarrhoea effect.

Conclusion

In conclusion, the present study has shown that extract of *Irvingia gabonensis* is a potent anti-diarrhoeic agent which can be used for the effective management of diarrhoea.

COMPETING INTERESTS

The authors declare that there are no conflicts of interest.


