Full Length Article

A controlled study to investigate anti-diarrhoeal effect of the stem-bark fractions of *Terminalia avicennioides* in laboratory animal models

Mohammed M. Suleiman\(^{a, *}\), Balkisu B. Oyelowo\(^a\), Ahmed Abubakar\(^b\), Mohammed Mamman\(^a\), Kamar-deen T. Bello\(^c\)

\(^a\) Department of Veterinary Pharmacology and Toxicology, Faculty of Veterinary Medicine, Ahmadu Bello University, Zaria, Nigeria
\(^b\) Department of Pharmacognosy and Drug Development, Faculty of Pharmaceutical Science, Ahmadu Bello University, Zaria, Nigeria
\(^c\) National Animal Production Research Institute, Ahmadu Bello University, Zaria, Shika, Nigeria

**A B S T R A C T**

Due to the shortcomings associated with modern synthetic anti-diarrhoeal drugs, it is important to find newer, safer and cheaper anti-diarrhoeal agents from natural sources. The study was conducted to evaluate the anti-diarrhoeal activity of the fractions of the stem-bark of *Terminalia avicennioides* in laboratory animal models. The effect of different concentrations (1.0 × 10^{-3}, 2.0 × 10^{-3}, 4.0 × 10^{-3} and 8.0 × 10^{-3} mg/mL) of the aqueous methanol (AMF), ethyl acetate (EAF) and hexane (HXF) fractions of *T. avicennioides* were tested against spontaneous and acetylcholine-induced contractions of rabbit jejunum and that of histamine-induced contraction of guinea pig ileum. Similarly, the effects of the AMF on gastro-intestinal transit time, castor oil-induced diarrhoea and castor oil-induced enteropooling were evaluated. The AMF, EAF and HXF at concentrations of 1.0 × 10^{-3}, 2.0 × 10^{-3}, 4.0 × 10^{-3} and 8.0 × 10^{-3} mg/mL attenuated the contractile effects of both the spontaneous and acetylcholine-induced contractions of rabbit jejunum and that of histamine-induced contraction of guinea pig ileum in a concentration-dependent manner. The AMF at doses of 200, 300 and 500 mg/kg produced significant (p < 0.05) reductions in gastrointestinal transit time of charcoal and incidence of castor oil-induced diarrhoea in mice relative to the untreated control. Similarly, at doses of 300 and 500 mg/kg, AMF significantly (p < 0.05) reduced the weight and volume of intestinal fluid in the treated mice when compared to the untreated animals. The results of this study showed that the stem-bark of *T. avicennioides* possesses spasmyloytic effect and could be a potential anti-diarrhoeal agent. However, detailed pharmacological trials are required to justify the clinical use of the plant for treating diarrhoea.

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1. Introduction

Diarrhoea is characterised by increased frequency of bowel, wet stool and abdominal pains [1]. It is usually a symptom of an infection in the intestinal tract, which can be caused by a variety of bacterial, viral, parasitic organisms and other non-infectious causes [2]. The World Health Organisation (WHO) estimated that 3–5 billion cases of diarrhoea occur each year (1 billion in children less than 5 years of age) [3]. Similarly, Ahmed et al. [4] reported that diarrhoea is the foremost fatal outcome among children in Nigeria under the age of five. Despite the availability of different approaches for diarrhoeal management, vast majority of the people in developing countries rely on herbal drugs for the management of diarrhoea [5]. The use of modern drugs for treating diarrhoea are usually associated with unwanted side effects (e.g. dry mouth and urinary retention often observed with the use of antimuscarinic drugs as atropine and headache, and nausea with calcium channel blockers). The synthetic opioid drugs (diphenoxylate and loperamide) cause severe constipation and may significantly slow gastrointestinal transit and increase the absorption of bacterial toxins in infectious diarrhoea [6]. Due to these shortcomings associated with modern anti-diarrhoeal drugs, it is important to find newer, safer and cheaper anti-diarrhoeal agents from natural sources [7].

*Terminalia avicennioides* (Combretaceae) is found in the savannah region of West Africa [8]. In Nigeria, the plant is locally called “bause”, “Idi”, “kpace”, “kpayi” and “Edo” in Hausa, Yoruba, Nupe, Gwari and Igbo languages, respectively [9,10]. Different parts of the plant have been used traditionally to manage conditions such as...
as gastric ulcer, gastro-intestinal disorders (diarrhoea), bloody sputum, cough, and gastro-intestinal helminth parasites [11–13]. Evaluation of the antidiarrhoeal effects of this plant is an attempt not only to validate the traditional claim but that is also hoped to provide an alternative source for an effective treatment against diarrhoea.

2. Materials and methods

2.1. Plant material

The stem-bark of *T. avicennioides* was obtained from the wild, around Zaria, Nigeria. Samples of the flowers, leaves and seeds of the plant were sent to the Herbarium, Department of Biological Sciences, Ahmadu Bello University, Zaria, for identification. A voucher specimen with number 900239 was deposited at the Herbarium for reference purposes.

2.2. Experimental animals

Five adult New Zealand white rabbits weighing between 2.0 and 2.5 kg and 5 adult guinea pigs of 300 and 400 g weight were purchased from a local market in Zaria. One hundred adult Swiss albino mice of both sexes weighing between 22 and 23 g were obtained from the Department of Pharmacology and Therapeutics, Faculty of Pharmaceutical Sciences, Ahmadu Bello University, Zaria. The animals were acclimatized to laboratory conditions for two weeks and were fed on commercial rodent diet. In addition, water was provided *ad libitum*. All animal experiments were done according to the ethical guidelines on laboratory animal use and care policy, which is in line with Ahmadu Bello University Research Policy (revised in 2010).

2.3. Equipment and laboratory materials used

Locally made cages; Wooden mortar and pestle; Whatman filter paper size 1; Conical flask; Macerating bottle; Measuring cylinder, Test tubes (Pyrex, France); Syringes (1 mL and 5 mL); Weighing balance (Lab tech. BL 20001 and Mettler P162, USA); Microdynamometer (Ugo Basile, Italy); Water bath (HH-S Digital thermostatic water bath, China); Dissecting kit (Gold Cross Dissecting Set, Malaysia); Plastic ruler and Stop watch.

2.4. Drugs and chemicals

Acetylcholine (Ach) and histamine (H) were purchased from Sigma-Aldrich Inc., 3050 Spruce Street, St. Louis, USA. Castor oil (Bell, Sons and Co Ltd, Southport PR9 9 AL, England); Loperamide (Imodium® – Janssen Pharmaceutical, Pakistan); Medicinal charcoal (Ultracarbon® tablets–Merck KGaA, Darmstadt, Germany); Carboxymethylcellulose.

2.5. Plant extraction and partitioning

The stem-bark of *T. avicennioides* was air-dried at room temperature to a constant weight. The dried plant part was made into powdered form using wooden mortar and pestle. One thousand five hundred grams of the powdered stem-bark of the plant were extracted by maceration in a macerating bottle using 4.5 litres of methanol (98%) as solvent at room temperature. The process was repeated twice and the extracts were pooled together. The liquid extract was concentrated in vacuo using rotary evaporator at 40 °C. The crude methanol extract was dissolved in water and serially partitioned with n-hexane and ethyl acetate in a separating funnel. Similarly, the fractions obtained were concentrated in vacuo at 40 °C. All the fractions were weighed, labeled and stored in an air tight container at 4 °C until required.

2.6. Phytochemical screening

The fractions were screened to detect the presence of alkaloids, anthraquinones, carbohydrates, cardiac glycosides, flavonoids, saponins, steroids/triterpenes and tannins using standard test methods [14].

2.7. In vitro studies

2.7.1. Effect of fractions of *T. Avicennioides* on isolated rabbit jejunum

The rabbits were deprived of food but not water for 18 h before the study. Each rabbit was sacrificed by cervical dislocation and exsanguinated. Their abdomens were cut open and segments of the jejunum (2 cm) were cut and dissected from the adhering mesentery. Each tissue was suspended in a 25 mL organ bath containing Tyrode's solution, aerated with air and allowed to stabilize for 30 min to acclimatize. The effect of different concentrations of acetylcholine (Ach) (1.0 × 10⁻⁶ mg/mL–8.0 × 10⁻⁶ mg/mL) and the fractions of *T. avicennioides* (1.0 × 10⁻³ mg/mL–8.0 × 10⁻³ mg/mL) were tested on both spontaneous and Ach-induced contractions of the isolated tissue. The contact time for each tested fraction on the tissue was 30 s, which was followed by washing the tissue three times with Tyrode's solution. The tissue was allowed to rest for a period of 15 min before the next addition of drug or extract. Changes in tension produced by the test agent were recorded with a microdynamometer (sensitivity of 3.0 mV and speed of 24 mm/min coupled to an isotonic transducer [12].

2.7.2. Effect of fractions of *T. Avicennioides* on isolated guinea pig ileum

Similarly, the method described above for the effect of stem-bark extracts of *T. avicennioides* on isolated rabbit jejunum was used. The effects of histamine (1.0 × 10⁻³ mg/mL–8.0 × 10⁻³ mg/mL) and fractions of *T. avicennioides* (1.0 × 10⁻³ mg/mL–8.0 × 10⁻³ mg/mL) against histamine-induced contraction of the tissue were also tested.

2.8. Acute toxicity test

The median lethal dose (LD₅₀) as an indication of the acute toxicity of the extract was determined by the method described by Lorke [15]. The test was carried out in two phases. All the animals were fasted for 12 h prior to oral administration of the AMF. In phase one, nine apparently healthy Swiss albino mice were randomly divided into three groups of three mice. Mice in groups one, two and three received AMF orally at 10, 100 and 1000 mg/kg, respectively. The mice were observed over a period of 48 h for signs of toxicity and mortality. In the second phase, three mice were randomly assigned into three groups of one mouse each. Animals in groups one, two and three were treated with AMF orally at 1600, 2900 and 5000 mg/kg, respectively. Similarly, the animals were observed for 48 h for any signs of toxicity or mortality. The obtained results were recorded accordingly.

2.9. In vivo studies

2.9.1. Effect of the aqueous methanol fraction on gastro-intestinal transit time in mice

The method described by Vogel and Vogel [16] was used. Twenty-five mice were deprived of feed for 12 h and randomly allotted into five groups of 5 mice each. Animals in groups one and two were administered with distilled water (5 mL/kg) and loperamide (5 mg/kg) and served as untreated and treated control groups, respectively. Similarly, mice in groups three, four and five
were given AMF at 200, 300 and 500 mg/kg, respectively. All treatments were given orally. A solution of charcoal meal was prepared by adding ground charcoal tablets (250 mg) to a boiled solution of 2% carboxymethylcellulose in a 20 mL beaker. Distilled water was added gradually to give a slurry aqueous suspension. The suspension was allowed to cool before administering it to the animals. One hour post-treatment, 0.3 mL of the charcoal meal was administered orally to each mouse. One hour after the charcoal meal was given, all the mice were sacrificed by cervical dislocation, their abdomens were cut open, and their intestines were carefully removed from the cardia to the anus. The intestines were immediately immersed in formalin to arrest peristalsis. Thereafter, the intestines were washed in clean tap water. The distance transverse by the meal through the intestine of each mouse was shown by the charcoal meal front, which was then measured using a graduated cylinder and expressed as peristaltic index (PI). The peristaltic index and percentage of inhibition were calculated using the following:

\[
\text{Peristaltic Index} = \frac{\text{Distance travelled by charcoal meal}}{\text{Total length of small intestine}} \times 100
\]

Percentage of inhibition = \(\frac{D_{c} - D_{t}}{D_{c}} \times 100\)

where 

\(D_{c}\): Mean distance travelled by the charcoal in the control group

\(D_{t}\): Mean distance travelled by the charcoal in the test group.

2.9.2. Effect of the aqueous methanol fraction on castor oil-induced diarrhoea in mice

The method described by Vogel and Vogel [16] was used in this study. Twenty-five mice were allocated at random into five groups of five mice each and were deprived of feed for 12 h. Animals in group one (untreated control) and two (treated control) were given distilled water (5 mL/kg) and loperamide (5 mg/kg), respectively. Mice in groups three, four and five were given AMF at 200, 300 and 500 mg/kg, respectively. All treatments were given through the oral route. One hour after drug treatment, all mice were orally administered castor oil (2 mL/kg). One hour after castor oil administration, the mice were sacrificed by cervical dislocation and the small intestine of each mouse was removed after tying the ends with threads and weighed. The intestinal content of each mouse was collected in a graduated cylinder and the volume measured. Thereafter, it was weighed again and the difference between the initial and final weights of the intestines was recorded. The percentage reduction of intestinal secretion (volume and weight) was calculated relative to the negative control using the formula:

Percentage of inhibition = \(\frac{\text{control} - \text{Test}}{\text{Control}} \times 100\)

3. Results

A yield of 77.37 g (5.16%) of the crude methanol extract was obtained when 1.5 kg of the dried powder of the stem-bark of \(T. avicennioides\) was completely extracted with methanol. Similarly, the aqueous methanol extract after partitioning with ethyl acetate and hexane gave yields of 31.89 g (41.22%), 1.76 g (2.27%) and 0.4 g (0.52%) of AMF, EAF and HXF, respectively. Qualitative phytochemical tests on the extracts of the stem-bark of \(T. avicennioides\) revealed the presence of saponins and steroidal/terpenoids in AMF, EAF and HXF. In addition, AMF and EAF contain carbohydrate, tannins and cardiac glycosides. Flavonoids and alkaloids were also detected in the AMF (Table 1).

The AMF, EAF and HXF inhibited the spontaneous contractions of the isolated rabbit jejunum in a concentration-dependent fashion (Figs. 1–3). In addition, the AMF and EAF produced a concentration–dependent inhibition of acetylcholine (Ach)-induced contraction of isolated rabbit jejunum (Figs. 4 and 5).

The AMF and EAF inhibited histamine (H)-induced contraction of isolated guinea pig ileum in a concentration-dependent fashion (Figs. 6 and 7).

Oral administration of the extract at a dose of up to 5000 mg/kg did not produce mortality or any toxic effect in the tested animals. The median lethal dose \((LD_{50})\) was therefore considered to be >5000 mg/kg in mice. The extract at doses of 300 and 500 mg/kg significantly \((p < 0.05)\) reduced both the length transversed by the charcoal meal and the peristaltic index when compared with the untreated control (distilled water) group. Loperamide (5 mg/kg), which is a standard antidiarrhoeal drug, showed a greater inhibitory effect (51.61%) than the extract, as shown in Table 2.

The extract at doses of 200, 300 and 500 mg/kg significantly \((p < 0.05)\) afforded protection against diarrhoea in mice induced by castor oil in a dose-dependent manner when compared with the distilled water (untreated) control group (Fig. 8).

The extract at 200, 300 and 500 mg/kg, significantly \((p < 0.05)\) inhibited castor oil-induced enteropooling in mice. At the same doses, it also inhibited the weight of intestinal fluid content of treated mice by 21.44%, 32.53% and 43.62%, respectively. The potent anti-diarrhoeal drug; loperamide (5 mg/kg), also significantly \((p < 0.05)\) inhibited the weight of intestinal fluid content (54.71%) relative to the untreated control. Similarly, the volume of the diarrhoeal index for each group was calculated by multiplying the number of mice in each grade by the number of grade divided by the number of mice in each group.
Table 1
Phytochemical constituents investigated in the extracts of the stem-bark of *Terminalia avicennioides*.

| Phytochemical constituent | Phytochemical test | Inference | AMF | EAF | HXF |
|---------------------------|-------------------|-----------|-----|-----|-----|
| Alkaloids                 |               |           | +   |     |     |
| Anthraquinones            |               |           |     |     |     |
| Carbohydrates             | Fehlings        |           |     | +   |     |
| Cardiac glycosides        | Kelle-Kiliani   |           |     |     |     |
| Flavonoids                | Shinoda         |           |     |     |     |
| Saponins                  | Frothing        |           |     | +   |     |
| Steroids/triterpenes      | Liberman Buchard|           |     |     | +   |
| Tannins                   | Ferric chloride |           |     |     |     |
|                           | Lead acetate    |           |     |     |     |

Key: + = present; − = not present; AMF = Aqueous methanol fraction; EAF = Ethyl acetate fraction; HXF = Hexane fraction.

Fig. 1. Effects of different concentrations (1.0 × 10⁻³ mg/mL–8.0 × 10⁻³ mg/mL) of aqueous methanol fraction (AMF) of the stem-bark of *Terminalia avicennioides* on the spontaneous contraction of isolated rabbit jejunum.

Fig. 2. Effects of different concentrations (1.0 × 10⁻³ mg/mL–8.0 × 10⁻³ mg/mL) of the ethyl acetate fraction (EAF) of the stem-bark of *Terminalia avicennioides* on spontaneous contraction of isolated rabbit jejunum.
Fig. 3. Effects of different concentrations (1.0 × 10^{-2} mg/mL–8.0 × 10^{-2} mg/mL) of the hexane fraction (HXF) of the stem-bark of *Terminalia avicennioides* on spontaneous contraction of isolated rabbit jejunum.

Fig. 4. Effects of different concentrations (1.0 × 10^{-3} mg/mL–8.0 × 10^{-3} mg/mL) of the aqueous methanol fraction (AMF) of the stem-bark of *Terminalia avicennioides* on acetylcholine (Ach)-induced contraction of rabbit jejunum.

Fig. 5. Effects of different concentrations (1.0 × 10^{-3} mg/mL–8.0 × 10^{-3} mg/mL) of the ethyl acetate fraction (EAF) of the stem-bark of *Terminalia avicennioides* on acetylcholine (Ach)-induced contraction of isolated rabbit jejunum.
intestinal fluid in mice was significantly (p < 0.05) reduced at all the tested doses of the extract, as shown in Table 3.

### Table 3

| Treatment (mg/kg) | Mean length of small intestine ± SEM (cm) | Mean distance of charcoal meal ± SEM (cm) | Peristaltic index | % of inhibition |
|------------------|-------------------------------------------|-------------------------------------------|-------------------|----------------|
| Distilled Water  (5 mL/kg) | 46.12 ± 1.80                             | 37.2 ± 2.43                               | 80.34 ± 2.36      | –              |
| Loperamide (5)   | 43.90 ± 1.11                              | 17.12 ± 0.97                              | 38.88 ± 1.37      | 51.61          |
| AMF (200)        | 47.84 ± 1.44                              | 35.66 ± 1.30                              | 74.49 ± 0.91      | 7.28           |
| AMF (300)        | 45.10 ± 2.26                              | 24.90 ± 2.20                              | 54.81 ± 2.23      | 31.78          |
| AMF (500)        | 49.0 ± 1.64                               | 22.44 ± 0.75                              | 45.86 ± 1.21      | 42.92          |

Means on the same column with different superscript letters are significantly (p < 0.05) different.

4. Discussion

This study was carried out to evaluate and validate the traditional claims on the antidiarrhoeal effect of the stem-bark of *Terminalia avicennioides*, which is a plant used widely among the Fulanis (major cattle rearing people of West Africa) for treating diarrhoea in both animals and humans.

The in vitro effect of the different fractions (AMF, EAF and HXF) of the plant obtained from the crude methanol extract was tested against contractions of isolated rabbit jejunum (spontaneous and acetylcholine-induced) and guinea pig ileum (histamine-induced) in an attempt to find the fraction with the most promising effect against diarrhoea.

Diarrhoea is usually the frequent passage of liquid stools. It involves increased gastrointestinal motility and secretion, a decrease in absorption of fluid, and loss of electrolytes and water [18]. In some types of diarrhoea, the secretory component...
predominates, while other types of diarrhoea are characterized by hypermotility [19].

The results of the phytochemical screening of the stem-bark detected metabolites that concur with that earlier reported by Mann and Yusuf [20]. Garba et al. [21] reported the presence of alkaloids, saponins, cardiac glycosides, steroids, tannins and phe-nols in the roots of *T. avicennioides.*

The fractions of the crude methanol extract (AMF, EAF and HXF) of *T. avicennioides* concentration-dependently inhibited both the spontaneous and acetylcholine-induced contractions of isolated rabbit jejunum. Similarly, the fractions attenuated the contraction of guinea pig ileum caused by histamine in a concentrationdependent manner. Acetylcholine and histamine are agonists that cause intestinal smooth muscle contractions by activating the muscarinic M3 and histamine H1 receptors, respectively. Acetylcholine induces smooth muscle contraction via inositol phosphate (IP3) pathway, which mediates Ca2+ release from sarcoplasmic reticulum [22]. The role of calcium ions (Ca2+) in producing smooth muscle tension has been well established [23]. Ca2+ serves as a second messenger in smooth muscle cells. An increase in cytoplasmic Ca2+ concentration and binding to calmodulin, and the activation of myosin light chain kinase, is the primary stimulus for contraction. The Ca2+ used in the activation of the contractile apparatus enters the Cytoplasmic compartment during periods of membrane depolarisation, mechanical distortion, or stimulation by agonists. Release of Ca2+ from intracellular stores is a second means of increasing calcium ion concentration [24,25].

In the mammalian tissues, histamine is distributed widely. The enterochromaffin-like cells (ECF) that are found mainly in the fundic mucosa of the stomach produce histamine by decarboxylation of the amino acid, histidine. Histidine is converted to histamine by a specific enzyme, histidine decarboxylase. Histamine is known to contract several types of smooth muscle, including those of the bronchi, gut and large blood vessels [26]. Histamine-induced contraction of the smooth muscle of guinea pig ileum is mediated by H1 receptors found throughout the gastrointestinal tract. [27]. Moreover, the contractile effect of histamine on both vascular smooth muscle and endothelial cells cause increase in vascular permeability [28].

In a similar study, the extract of the roots of *T. avicennioides* was shown to attenuate acetylcholine-induced contraction of isolated rabbit jejunum [12]. Saponins have been shown to inhibit the in vitro release of histamine [29]. Tannins, flavonoids, saponins, sterols/triterpenes, present in some medicinal plants, were demonstrated to possess spasmylic effect [30]. It was shown also that flavonoids inhibit intestinal secretion of autacoids and prostan-gldins [31] and inhibit intestinal contractions induced by spasmo-gens [32]. Flavonoids were shown to have the ability to attenuate the contraction of guinea pig ileum induced by some spasmodgens such as acetylcholine [33]; they also have the ability to inhibit small intestinal transit [34]. In this study, phytochemicals such as alkaloids, flavonoids, saponins, tannins and terpenoids detected in the fractions of *T. avicennioides* are reported to be responsible for the observed spasmylic effect of the plant [35,36]. It is of interest to note that the AMF produced more spasmylic effect on both the rabbit and guinea pig ileum during the *in vitro* tests. This finding clearly suggested that flavonoids which were detected only in the AMF largely are responsible for the spasmylic effect of the plant.

The median lethal dose (LD50) as an indication of acute toxicity of AMF, which showed the most promising effect during the in vitro trial was calculated to be ≥5000 mg/kg. According to Lorke [15], any substance with LD50 of ≥5000 mg/kg should be considered relatively safe. Determination of median lethal dose of a chemical is usually the first step taken to ascertain the safety of that chemical with the sole intent of using it as a drug. Acute toxicity test also allows further toxicological and pharmacological testing of candidate drugs to justify their use in clinical situations.

Diarrhoea is usually considered a result of altered motility and fluid accumulation within the intestinal tract. Many antidiarrhoeal agents act by reducing gastro-intestinal motility and/or the secretions [37]. The extract significantly reduced intestinal transit time in mice fed charcoal meal. Drugs that display such properties are known to be good candidates as antidiarrheal agents [38]. Loperamide (an opioid derivative), has been shown to slow intestinal motility by its action on mu (μ) receptors on neurons in the submu-cosal neural plexus of the intestinal wall and by its antimuscarinic activity in the gastrointestinal tract [39]. Antimuscarinic drug possess their antidiarrheal action by inhibiting both the gastrointestinal secretions and motility [18]. The antidiarrheal activity of some plant extracts had been ascribed to their antagonistic action on muscarinic receptors found in the gastro-intestinal tract [40]. Studies on the functional role of tannins against diarrhoea, revealed that they inhibit gastro-intestinal movement by reducing the intracellular Ca2+ inward current or by activation of the calcium pumping system [41] as well as forming protein tannates, which make the intestinal mucosa more resistant and hence, reduce peristaltic movement [42]. The extract also inhibited the intestinal propulsion as shown by its inhibitory action against charcoal meal motility. Agents with such activity were known to have antidiarrheal effect [43].

![Fig. 8. Effect of the aqueous methanol fraction (AMF) of the stem-bark of *Terminalia avicennioides* on castor oil-induced diarrhoea in mice. Untreated control group was dosed with distilled water (DW) while treated control group was dosed with loperamide (LP). Data are expressed as mean ± SEM. *p < 0.05; **p < 0.01 compared to the untreated control group; (Kruskal-Wallis one-way ANOVA test).](image)

| Treatment (mg/kg) | Mean weight of small intestine (g) | % inhibition | Mean volume of small intestinal contents (ml) | % inhibition |
|------------------|-----------------------------------|-------------|---------------------------------------------|-------------|
| Distilled water (5 ml/kg) | 1.123 ± 0.07 | – | 0.888 ± 0.03 | – |
| Loperamide (5) | 0.486 ± 0.01 | 54.71 | 0.374 ± 0.02 | 61.80 |
| AMF (200) | 0.846 ± 0.02 | 21.44 | 0.734 ± 0.02 | 17.98 |
| AMF (300) | 0.730 ± 0.01 | 32.53 | 0.566 ± 0.03 | 35.96 |
| AMF (500) | 0.630 ± 0.01 | 43.62 | 0.520 ± 0.02 | 43.82 |

Means on the same column with different superscript letters are significantly (p < 0.05) different.
Similarly, the study showed that the extract inhibit castor oil-induced diarrhoea in mice in a dose-dependent manner. Castor oil, an irritant or stimulant laxative, is hydrolysed in the upper small intestine to ricinoleic acid, a local irritant that irritates the mucosa of the gastrointestinal tract resulting in increase in small intestine to mucosa of the gastrointestinal tract resulting in increase in volume of intestinal fluid in mice when compared to animals in the untreated control group. This effect can be attributed to either a decrease in mucosal secretion or increase in mucosal absorption. Nwafor et al. [49] showed that the suppression of intestinal fluid accumulation by plant extracts produced antidiarrhoeal activity via inhibition of the gastrointestinal function. In this study, the extract significantly reduced both the weight and volume of intestinal content which may have promoted reabsorption of fluids due to decrease propulsion in the intestinal of mice. This may possibly explain the mechanism of the extract anti-enteropooling action. Longanga et al. [29] screened a number of medicinal plants and showed that the antidiarrhoeal activity of those plants were due to tannins, alkaloids, saponins, flavonoids, sterols, triterpenes and reducing sugars contained in them. In a similar study, the antidiarrhoeal activities of some medicinal plants were found to be due to tannins, alkaloids, saponins and flavonoids contained in them [50]. Hence, the antidiarrhoeal activity exhibited by the aqueous methanol fraction of the stem-bark of *T. avicennioides* could be due to presence of one or more of these metabolites.

5. Conclusions

The present study showed that the fractions (AMF, EAF and HXF) of the crude methanol extract of *T. avicennioides* possess spasmolytic effect in an *in vitro* study. In addition, the AMF possesses antidiarrhoeal effects which could be due to the metabolites present in the fraction. This study therefore, supports the traditional use of the plant as an antidiarrhoeal agent. Further studies are ongoing to isolate, characterize and elucidate the mechanism(s) of compounds with antidiarrhoeal action in the plant.

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References

[1] Ezekwesili CN, Obiara KA, Ugwu OP. Evaluation of anti-diarrhoeal properties of crude aqueous extract of *Ocimum gratissimum* L. (Labiatae) in rats. Biochem 2000;16:122–31.

[2] Mzengu I, Inabo HI, Olonitola SO, Aminu M. Antibiotic susceptibilities of *Salmonella* species prevalent among children of 0–5 years with diarrhoea in Katsina state, Nigeria. Arch Med Biomed Res 2016;3:39–51.

[3] Heinrich M, Heneka B, Anka A, Rimpler H, Sticher O, Kostka T. Spasmolytic and antidiarrhoeal properties of the Yuteen Mayan medicinal plant *Cosmopterus tetramerus*. J Pharm Pharmacol 2005;57:1081–5.

[4] Ahmed AA, Zezi AM, Yaro AH. Antidiarrhoeal activity of the leaf extracts of *Dolichos lablab*, *Hitch and Dalz* (Fabaceae) and *Ricinus communis*, MDQ (Moraceae). Afr J Tradit Complement Altern Med 2007;4:524–8.

[5] Atta AH, Mounir SM. Antidiarrhoeal activity of some Egyptian medicinal plants extract. J Ethnopharmacol 2004;92:303–3.

[6] Patrick MG. Drugs affecting gastrointestinal function. In: Riviere JE, Papich MG, editors. Veterinary pharmacology and therapeutics. Ames, Iowa, USA: Wiley-Blackwell; 2009. p. 1247–72.

[7] Benoit N, Gessain BB, Dosso K, Gnangoun BN, Ansamet PG, Asiedu-Gyekye IJ, et al. Antibacterial and antispasmodic activities of a dichloromethane fraction of an ethanol extract of stem bark of *Pilostigma reticulatum*. J Pharm Biol Allied Sci 2015;7:128–35.

[8] Tausa A, Ajoboso OSO, Ajegbe S, Gbade M, Isahia S. Evaluation of the wound healing activity of ethanol extract of *Terminalia avicennioides* root bark on two wound models in rat. Int J Med Arom Plants 2011;1:95–100.

[9] Dalziel JM. The useful plants of west tropical Africa. Crown Agents for Overseas Governments and Administrations, London; Millbank; 1955. p. 81.

[10] Salau AK, Yakubu MT, Oladji AT. Cytotoxic activity of aqueous extracts of *Anogeissus leiocarpus* and *Terminalia avicennioides* root barks against Ehrlich ascites carcinoma cells. Int J Pharm Res 2013;83:381–5.

[11] Veirbano J, Adam JC, de la casamace (Senegal). Ann Pharm Fr 1963;21:853–70.

[12] Abdulahi AL, Agbo MO, Amos S, Gamanial KS, Wambebe C. Antidiarrhoeal activity of the aqueous extract of *Terminalia avicennioides* roots. Phytother Res 2001;15:431–4.

[13] Mann A. Survey of ethnomedicine for the treatment of tuberculosis: chemistry perspective. Minna, Niger State, Nigeria: Ayanwara Printing Works; 2007. p. 1–117.

[14] Tauxe GE, Evans WC. Pharmacognosy. Oxford: Alder Press; 1996. p. 213–2.

[15] Lorke D. A new approach to practical acute toxicity testing. Arch Tox 1983;54:275–87.

[16] Vogel HC, Vogel WH. Drug discovery and evaluation, pharmacological assays. In: Vogel HC, Vogel WH, editors. Methods in clinical pharmacology, Berlin: Springer; 1997. p. 757.

[17] Igboeli O, Onyeto CA, Okorie AN. Antidiarrhoeal activity of methanol leaf extract of *Lophira lanceolata* var *Tiegh* (Ochnaceae). Mer Res J Environ Sci Tech 2015;3:59–64.

[18] Hardman JG, Limbird LE. Drugs affecting gastrointestinal function. In: Goodman and Gilman’s the pharmacological basis of therapeutics. New York: McGraw-Hill; 1992. p. 914–31.

[19] Mann A, Yusuf I. Antibacterial activity of methanic extracts of *Terminalia avicennioides* against fish pathogenic bacteria. Am J Res Commun 2014;4:133–47.

[20] Garba S, Salihu L, Shoje M. Antidiarrhoeal activities of some medicinal plants. Med Chem 2015;52:001.

[21] Gordienko DV, Harhun MI, Kustov MV, Pucovsky V, Bolton TB. Sub-plasmalemmal (Ca**2+**)- upstroke in myocytes of the guinea-pig small intestine evoked by muscicarn stimulation: ITPR-mediated Ca**2+** release induced by voltage-gated Ca**2+** entry. Cell Calcium 2008:43:122–41.

[22] Bolton TB. Mechanisms of action of transmitters and other substances on smooth muscle. Physiol Rev 1979;59:606–718.

[23] Sanders KM. Invited review: mechanism of calcium handling in smooth muscles. J Appl Physiol 2001;91:1438–49.

[24] Sanders-Bush E, Mayer SE. 5-Hydroxytryptamine (serotonin): receptor agonist and antagonist. In: Hardman JG, Limbird LE, editors. The pharmacological basis of therapeutics. New York: McGraw-Hill; 2001. p. 269–90.

[25] Adams HR. Histamine, serotonin, and their antagonists. In: Riviere JE, Papich MG, editors. Veterinary pharmacology and therapeutics. Ames Iowa USA: Wiley-Blackwell; 2009. p. 411–27.

[26] Bertaccini G, Molina E, Zappia L, Zoesi J. Histamine receptors in guinea pig ileum. Naunyn-Schmeideberg’s Arch Pharmacol 1979;309:65–8.

[27] Coruzzi G, Poli E, Morini G, Bertaccini G. The histamine H3 receptor. In: Gaggini TS, Guglietta A, editors. Molecular targets for drug development: GI diseases. New Jersey, USA: Humana Press; 2000. p. 239–67.

[28] Tiwari P, Kumar B, Kaur M, Kaur G, Kaur H. Phytochemical screening and extraction: a review. Int J Pharm Sci 2011;3(1):98–106.

[29] Loganga OA, Vercruysse A, Foriers A. Contribution to the ethnobotanical, phytochemical and pharmacology studies of traditionally used medicinal plant in the treatment of dysentery and diarrhoea in Lomela area, Democratic Republic of Congo (DRC). J Ethnopharmacol 2000;71:41–23.

[30] Vimala R, Nagarajan S, Alam M, Susan T, Joy S. Antiinflammatory and antioxidant activity of *Michelia champaca* L. (White variety), *bora brachata Roxb.*, and *Rhynchosia cana* (Willd.) D.C. flower. Ind J Exp Biol 1997;35:1310–4.
[31] Capasso F, Pinto A, Mascolo N, Autore G, Franco MP. Effects of flavonoids on PGE2 and LTD4-induced contractions of guinea pig isolated ileum. Pharm Res Commun 1988;20:201–2.
[32] Hejazian SH, Bagheri SM, Dashti-R MH. Relaxant effect of *Humulus lupulus* extracts on isotonic rat’s ileum contractions. Avian J Phytomed 2013;4:53–8.
[33] Khoshnazar SM, Bahaddini A, Najafipour H. Effect of alcoholic extract of licorice (*Glycyrrhiza glabra*) rhizome on isolated duodenum motility in male rats and its interference with cholinergic, nitroglic, and adrenergic systems. Bull Environ Pharm Life Sci 2013;2:173–7.
[34] Naz A, Wadood AS. Antispasmodic activity of *Teucrium stocksianum* Bios. Pak J Pharm Sci 2011;24:171–4.
[35] Cortes AR, Delgadilo AJ, Hurtado M, Dominguez-Ramizez AM, Medina JR, Aoki K. The antispasmodic activity of *Buddleja scordiodes* and *Buddleja perfoliata* on isolated intestinal preparations. Biol Pharm Bull 2006;29:1186–90.
[36] Spruill WJ, Wade WE. Diarrhoea, constipation and irritable bowel syndrome. In: Matzke GR, Wells BG, Possey ML, editors. Pharmacotherapy: a pathophysiological approach. New York: McGraw-Hill Medical Publishing Division; 2005. p. 312–6.
[37] Adzua B, Tarfab F, Gamaniel KS. The efficacy of *Sphaeranthus senegalensis* Vaill extract against diarrhoea in rats. J Ethnopharmacol 2004;95:173–6.
[38] Waller DG, Renwick AG, Hillier K. Medical pharmacology and therapeutics. 2nd ed. London: Elsevier Saunders; 2005. p. 417–8.
[39] Rang HP, Dale MM, Ritter JM, Moore PK. The gastrointestinal tract. Pharmacology. 5th ed. Edinburgh: Churchill Livingstone; 2003. p. 376–377.
[40] Offiah VN, Chikwendu UA. Antidiarrhoeal effects of *Ocimum gratissimum* leaf extract in experimental animals. J Ethnopharmacol 1999;68:327–30.
[41] Belentrougui RG, Constantin B, Cognard C. Effects of two medicinal plants, *Psidium guajava* L. (*Myrtaceae*) and *Diospyros mespiliformis* L. (*Ebenaceae*) leaf extracts on rat skeletal muscle cells in primary culture. J Zhejiang Univ Sci 2000;7:56–63.