

Phytochemical Constituents and Antidiarrhoeal Activities of Ethanolic Extract of *Maesobotrya barteri* (Bush Cherry) Leaves in Albino Rats

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Abstract

Maesobotrya barteri, commonly known as bush cherry is used in orthodox medicine for the treatment of different ailments including diarrhoea. Hence, this study was designed to investigate the phytochemical compositions, acute toxicity (LD50) and the effect of ethanolic extract of *Maesobotrya barteri* leaves as an antidiarrhoeal agent. Castor oil was used to induce diarrhoea in albino rats. 500mg/kg, 1000mg/kg and 2000mg/kg extracts of *M. barteri* were used to protect the rats against castor oil induced diarrhoea. The quantitative phytochemical screening of *M. barteri* revealed the presence of flavonoids, Saponin, Phenol, alkaloid, oxalate and phytate at the concentrations of 101.06ug/g, 21.32ug/g, 7.42ug/g, 1.75ug/g, 1.50ug/g and 0.57ug/g respectively. No experimental animal used to determine the LD50 up to 5000mg/kg extract died. The result of the anti-diarrhoeal property of *M. barteri* after pretreatment with the extract showed a considerable dose – dependent decrease in the production of diarrhoeal faeces, reduced the rate of defecation and the onset of diarrhoea in albino rats. The inhibition of diarrhoea by the extract as shown in the result may be attributed to the high concentration of flavonoid present in the plant.

Keywords: *Maesobotrya barteri* leaves; Antidiarrhoea; Phytochemicals

Introduction

Plant materials are globally known to be sources of high concentrations of various groups of drugs. Hence, lots of researches are ongoing by scientists to unveil the potency of most plants since it is believed that all plants possess hidden virtues that will help combat diseases of nature if discovered. Owing to the outbreak of various diseases and microbial resistance to most conventional

drugs, proper identification and evaluation of the usefulness of any plant involves the investigation of proximate analysis and phytochemical constituents of such plants [1]. Despite great advances in modern medicine, many species of plants have been claimed to possess antibiotic activities in orthodox medical practice [2].

Maesobotrya barteri (bush cherry) belongs to *Euphorbiaceae* family. It is a broad leafed plant of about 10m long, found mainly in the Southern and Eastern parts of Nigeria, Sierra lone and West Africa. In Nigeria, the Benin people locally called it "Oruru" the Efik/ Ibibios, called it Nyanyatet and the Yorubas called it "Odun or Obomodu". It bears fruits from April to June [3]. The fruits are about 1cm long, ovoid and distinctly pointed, bearing succulent black-purple berries that are edible and stain the tongue [4]. The plant is highly cherished by the people of south eastern Nigeria, not just for its succulent edible fruits but also for its acclaimed medicinal effect on most ailments [5]. For a very long time, various parts of *M. barteri* have been in use in the local communities for the treatment of diarrhoea, dysentery, urethral discharge, Jaundice, cough, measles and others. Also, in the eastern and southern Nigeria, the stem is preferentially used as chewing stick and as an important raw material for juice making [6]. The fruits of *M. barteri* enhance waste removal and the fibres help lower cholesterol absorption by preventing plaque formation [7]. However, this plant remains under exploited globally as a result of lack of useful information on its potential as sources of substances that are vital to humans with respect to nutrition and medicines.

Diarrhoea is referred to as a condition of passing out at least three loose or liquid faeces per day [8]. It is characterized by an increased fluidity and hypermotility of the intestine as a result of an imbalance between the absorptive and secretory mechanisms in the gastrointestinal tract. This leads to loss of excess fluid and electrolytes in stools [8-10].

Diarrhoea is one of the most vulnerable and life threatening diseases worldwide, reaching epidemic proportions. It has long been known as one of the most significant health challenges in developing [11]. It is the next leading cause of infant mortality after pneumonia globally, accounting for about 5-8 million deaths throughout the world annually, mostly in children under the age of 5 [12]. According to Venkat [13], diarrhoea claimed 1,793,000 lives in European countries in the year 2001. Similarly, the World Health Organization (WHO) has estimated that four to five billion cases of diarrhoea occur annually with one billion in children under 5 years of age and 5 million deaths occur in diarrhoea each year with 50% in children under 5 years of age [14]. Diarrhea is mainly caused by infectious agents, plant and animal toxins, poorly absorbable matter and inflammatory problems of the gastrointestinal tract [6]. Additionally, Nwachoko and Jack [15] reported that diarrhoea is also

connected with some terminal ailments like AIDS and Ebola viral infection and this usually contributes to the immediate death of those patients due to excessive fluid loss resulting from electrolyte imbalance. However, most pharmaceutical antidiarrhoeal drugs are not readily available particularly to those in the rural areas. Therefore, to combat the problem of diarrhoea, the World Health Organization has initiated a diarrhoeal disease control programme that encourages study of herbal medicines for treatment and prevention of diarrhoea with plants, since patients who cannot afford treatment with conventional medicines usually resort to the use of herbs [16].

Materials and Methods

Collection and Identification of Plant Sample

The leaves of *Maesobotrya barteri* were collected in a farmland behind Cauty Grammar School, in Etche Local Government Area of Rivers state, Nigeria. The plant was taxonomically identified and authenticated by Dr. Okeke Chiemezie of the Department of Plant Science and Biotechnology, University of Port Harcourt, Rivers State, Nigeria. A voucher specimen with the herbarium number: UPH/V/1250 was deposited in the herbarium of the aforementioned department.

Experimental Animal Acquisition and Maintenance

Male/female albino rats weighing 100-150g were used for the experiments. The animals were obtained and housed in the Animal House of the Department of Biochemistry, University of Port Harcourt, Nigeria. The albino rats were kept for one week for acclimatization at room temperature with proper ventilation and under a naturally illuminated environment of 12hours day and night cycle. They were given standard diet (Top feeds grower's mash) and had access to clean water *ad libitum*. Animal care protocols were followed in accordance with the National Institute of Health Guide for the care and use of laboratory animals (NIH, 1996).

Preparation of Ethanolic Extract of *M. barteri*

The leaves of *Maesobotrya barteri* were washed in a running tap water to remove sand and other particles and then air-dried at room temperature in an emptied well ventilated room for two weeks. The dried leaves were ground into coarse powder using an electric grinder. 1000g of the sample was weighed into a macerating jar and macerated with 5000ml of 99% ethanol for 72 hours with constant shaking. It was sieved using a muslin cloth

and after which filtered with No.1 Whatman filter paper. The filtrate was placed in a rotary evaporator to recover the solvent and thereafter, concentrated to dryness in an evaporating dish placed on a water bath at 45°C. The weight of the concentrates was taken and the percentage yield calculated, the extract was kept at 4°C in a refrigerator prior to usage.

Extraction of Phytochemicals

One gram (1g) of sample was measured and placed in a test tube, ethanol (15ml) and potassium hydroxide (10ml of 50%) was added. The test tube was kept in a water bath to react at 60°C for 60 minutes. Thereafter, the sample was placed in a filtering funnel and washed thoroughly using 20ml of ethanol, 10ml of hot water and 3ml of hexane. The extract was then washed three times using 10ml of aqueous ethanol, the solution was dried using anhydrous sodium sulfate and the solvent was evaporated. The product was then solubilized in 1000ul of pyridine of which 200ul was measured into a vial for analysis.

Quantification by GC-FID

The analysis of the sample was performed on a BUCK M910 Gas Chromatography equipped with a flame ionization detector. A RESTEK 15 Meter MXT-1 column (15um x 250um x 0.15um) was employed. The temperature was 280°C with split less injection of 2ul of sample and a velocity of 30cms⁻¹. The oven which operated initially at 200°C was then heated to 330°C at a rate of 3°C min⁻¹ and the detector was set at a temperature of 320°C. The phytochemicals were then evaluated as the ratio of the area and mass of internal standard to the area of the identified phytochemicals. The amount of the different phytochemicals was expressed in ug/g.

Determination of Acute Toxicity

The acute toxicity LD₅₀ was determined in albino rats, following Lorke [17] method with little modification (use of rats instead of mice). Twelve (12) albino rats weighing 100-125g were divided into 4 groups of 3 rats per group. The ethanolic extract of *Maesobotrya barteri* was given orally in doses of 1000, 2000, 3000 and 5000mg/kg body weight respectively to the animals. Each group received single dose of the extract and the general signs and symptoms of toxicity such as ingestion of food and water, mortality, alterations in physical appearance and behavioural changes were noticed for a period of 48 hours and then for a period of 7 days. The acute toxicity LD₅₀ was estimated as geometric mean of the dose which

resulted in 100% lethality and that which caused no lethality at all.

Experimental Procedure on Castor Oil induced Diarrhoea in Albino Rats and Faecal Count

Castor oil-induced diarrhoea was determined by the method employed by Awouters, et al. [18]. Twenty albino rats of both sexes, weighing between 100-150g were fasted for 18 hours and distributed into five groups of four animals each. Group1 which served as the control were given distilled water (1ml) via the oral route, Group2 received the standard drug, Loperamide hydrochloride (5mg/kg body weight), while Group3, 4 and 5 were administered with the ethanolic extract of *M. barteri* at doses of 500, 1000 and 2000mg/kg body weight respectively. One hour after drug pretreatment, all the animals were given 1ml of castor oil orally. The animals were then placed in metabolic cages on a clean and dried floor for faecal count. The positions of the metabolic cages were changed after each hour. The rats were monitored for seven hours after castor oil administration. The time taken for onset of diarrhoea was recorded, the mean faecal droppings evaluated and expressed as percentage (%) inhibition of diarrhea [19].

$$\text{Percentage inhibition of wet faeces} = \frac{T_0 - T_1}{T_0} \times 100$$

Where

T₀ = number of wet faeces in control group

T₁ = number of wet faeces in test group

Results

The result of quantitative phytochemical analysis of *M. barteri* is as shown in Table 1. The table shows the concentration of flavonoids to be highest followed by concentration of saponin while phytate had the least concentration of 0.57ug/ml. There was no death noticed in the rats upon oral administration of ethanol extract of *M. barteri* even at 5000mg/kg body weight. This signifies that the LD₅₀ of the plant extract was higher than 5000mg/kg. Also, apart from slight sedation and weakness noticed at doses of 3000mg/kg and 5000mg/kg, the plant extract did not show any significant signs of toxicity in albino rats during the observation period.

The effect of *M. barteri* on castor oil induced diarrhoea in albino rats is as shown in Table 2. The result showed 60 to 120 minutes delay before the onset of diarrhoea in

animals with the extract treated groups and percentage inhibition of diarrhoea was highest in Group 2 while the

least percentage inhibition was noticed in Group 3.



Figure 1: *M. barteri* plant in its natural habitat.

Component	Subclass	Concentration (ug/g)
Flavonoid	Kaemferol	45.49
Flavonoid	Rutin	27.85
Flavonoid	Catechin	25.18
Flavonoid	Epicatechin	2.54
Saponin	-	21.32
Phenol	-	7.42
Alkaloid	Ribalidine	1.75
Oxalate	-	1.5
Phylate	-	0.57

Table 1: Quantitative Phytochemical Constituents of leaves of *M. barteri*.

Group	Treatment	OD (MIN)	MWF	I (%)
1	Control	60	3.57±1.02b	-
2	Loperamide 5mg/kg	240	0.43±0.30	87.96
3	Extract 500mg/kg	60	2.14±0.74	40.06
4	Extract 1000mg/kg	60	2.00±0.69	43.98
5	Extract 2000mg/kg	120	0.71±0.36b	80.11

Table 2: Effect of ethanolic extract of *Maesobotrya barteri* leaves on faecal count of castor oil-induced diarrhoea in albino rats.

Results reported as Means ± standard error (M ± SEM), n =7

The Superscript “b” indicates a statistical significant difference ($p < 0.05$) when the extract treated groups are compared to the control group.

Key: OD = Onset of diarrhoea
MWF= Mean wet faeces after 7 hours
I = Inhibition

Discussion

M. barteri which is generally known as ‘bush cherry’ belongs to a Euphorbiacea family. It is used in the traditional system in managing diarrhoea by orthodox medical practitioners mostly in Southern and Eastern Nigeria [4,6]. The study was designed to investigate the phytochemical constituents and antidiarrhoeal property of ethanolic extract of *M. barteri* leaves.

Table 1 showed the concentration of phytochemicals of *Maesobotrya barteri* leaves in this order; Flavonoids > Saponin > Phenol > Alkaloid > Oxalate > Phytate. The Phytochemical result revealed high to moderate concentrations of kaemferol, rutin, catechin, epicatechin as subclasses of Flavonoid and ribalidine, a subclass of alkaloid. This result has provided useful information on the phytochemical compositions of *M. barteri* leaves. The result corroborates with the findings of Ogunka–Nnoka, et al. [6] who opined that due to lack of useful information on the potential of *M. barteri* as sources of substances that are vital to humans with respect to nutrition and medicine, the plant has remained under exploited globally. Additionally, Nwachoko, et al. [15] reported that

proper identification and evaluation of the usefulness of any plant involves the investigation of proximate analysis and phytochemical constituents of such plant. Research has shown that medicinal plants are the valuable bisource of drugs used frequently in traditional system of medicine and pharmaceutical intermediates for synthetic drugs [20]. Also, reports from several findings noted that the phytochemicals found in most medicinal herbs are responsible for their antimicrobial, antibiotic, anticancer, anti-helminthic and anti-sickling properties [21-24].

The acute toxicity result of the ethanolic of *M. barteri* in albino rats indicated a high LD₅₀. This shows that the plant may be regarded as safe and this high level of safety agrees with its popular consumption by the people of the Southern and Eastern Nigeria

Table 2, showed the inhibitory effect of ethanolic extract of *M. barteri* leaves on the wet faecal count of castor oil induced diarrhoea in albino rats. The 2000mg/kg extract treated group (Group 5) which had a mean wet faecal count value of 0.71±0.36 showed a significant decrease (P > 0.05) when compared with the control group (Group 1) which had 3.57±1.02 mean wet faecal count value. Animals administered with the 500mg/kg and 1000mg/kg extract (Group 3 and 4) showed a non-significant decrease (P>0.05) in their mean wet faecal count (2.14±0.74 and 2.00±0.69), when respectively compared to that of the control group. Also, the result of the percentage inhibition of castor oil induced diarrhoea as shown in Table 2, revealed that the inhibitory effect of the plant extract was dose-dependent when the group pretreated with standard drug (Group 2) which had 87.96 percentage inhibition was compare with the extract treated group (Group 3,4 and 5) with percentage inhibition of 40.06, 43.93 and 80.11 respectively. The prevention of diarrhoea and the decrease in defecation rate are the basis of pharmacological investigation of a viable antidiarrhoeal agent [25]. Castor oil causes diarrhoea due to its active metabolite ricinoleic acid, which produces irritation and inflammation in the intestinal mucosa. This lead to the liberation of prostaglandin that elicits hypersecretion and changes the electrolytes permeability in the intestinal mucosa [26,27]. Reports have shown that plants phytochemical components (Flavonoids) are capable of evoking anti-diarrhoeal effect due to their ability to form complexes with protein to inactivate microbial adhesions, decrease intestinal motility and hyperhydrolitic secretion [19,26,28-30]. Hence, the presence of flavonoids at high concentration as shown in

Table 1, may be responsible for the antidiarrhoeal properties of *M. barteri* leaves.

Conclusion

This study revealed that the plant contains phytochemicals that are of medicinal important and its ethanolic extract showed remarkable dose-dependent decrease in the production of diarrhoeal stool, reduced the frequency of defecation and delayed the onset of diarrhoea in castor oil induced diarrhoea in albino rats. The findings suggested that the extract of *M. barteri* may have antidiarrhoeal potential which may provide a scientific rationale for its used by traditional healers for the management of various diarrhoeal related conditions.

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