

**Antidiarrhoeal Activity of Ethanolic Extract of
Adansonia digitata Fruit Pulp in Rats**

Maha Y. Abdelrahim, Babiker M. A. Elamin, Dalia J. Khalil and Samia M. A. El
Badwi.

J Phy Pharm Adv 2013, 3(6): 172-178

DOI: 10.5455/jppa.20130624013026



Antidiarrhoeal Activity of Ethanolic Extract of *Adansonia digitata* Fruit Pulp in Rats

*¹Maha Y. ¹Abdelrahim, ¹Babiker M. A. ²Elamin, Dalia J. ¹Khalil and ¹Samia M. A. E. ¹Badwi.

¹ Department of Pharmacology and Toxicology Faculty of Veterinary Medicine University of Khartoum- Sudan.

².Department of Medicine Pharmacology and Toxicology collage of Veterinary Medicine University of Bahri.

Abstract

The antidiarrhoeal activity of ethanolic extract of *Adansonia digitata* fruit pulp was evaluated in rats. Studies were investigated on castor oil –induced diarrhoea, castor oil induced fluid accumulation and electrolyte secretion. *Adansonia digitata* was orally administered to rats at dose rate of 250 and 500mg/kg and Loperamide was used at 3mg/kg as reference antidiarrhoeal drug. The extract produced a dose dependant and significantly protected rats against castor oil induced diarrhoea. The frequency of defecation as well as weight of the feces was significantly ($p>0.05$) reduced. A preliminary phytochemical screening of the ethanolic extract of *A. digitata* revealed the presence of, flavonoids, saponins, tannins triterpenes, alkaloids and glycosides.

Keywords: *Adansonia digitata*, antidiarrhoeal activity, castor oil.

* Corresponding Author: Department of Pharmacology and Toxicology Faculty of Veterinary Medicine University of Khartoum- Sudan.

Received on: 05 Jun 2013.

Revised on: 24 Jun 2013.

Accepted on: 27 June 2013.

Online Published on: Jun 2013.

Introduction

African baobab is a very long-living tree, in the Sudan it was known locally as Tabaldi, the fruits are named Gunguleiz. Baobab leaves, bark, pulp and seeds are used as food and for multiple medicinal purposes in many parts of Africa (Diop et al., 2005).

The baobab fruit pulp can be considered a highly valuable source containing levels of vitamin C ranging from 2.8-3 g/kg and this was six times more than the contents of an orange which was considered the best source of ascorbic acid. Higher levels of pro vitamin A were detected in the young leaves, especially when they are used as dried material (Sidibé, et al., 1996; El-Kamali, et al., 1999).

Diarrhoea is one of the main water-borne diseases, endemic in many regions of the world and considered to be the major health threats to the world populations, both in tropical and subtropical poor countries (Damiki and Siva, 2011).

People customarily using the plants or plant-derived preparations consider them to be efficacious against diarrhoeal disorders without any scientific basis to explain the action of such plants. WHO has encouraged studies for treatment and prevention of diarrhoeal diseases using traditional medicinal practices (Atta and Mounair, 2004)

Diarrhoea is a common gastrointestinal disorder characterized by increased frequency of bowel, wet stool and abdominal pains (Ezekwesili et al., 2004; Farthings, 2002). Diarrhoea resulting from an imbalance between the secretory and absorptive forces in the intestine and it is an important symptom and complications of many disease of great public health importance. (Tijani et al., 2009) On the other hand, the WHO has contributed a Diarrhoeal Disease Control Programme including prevention, approach evaluations of health

education and studies on traditional medicine uses for the management of the disease suggesting that the herbal treatment is still important (Synder and Merson, 1982).

The present study was aimed to verify the traditional claim that, *Adansonia digitata* possess antidiarrhoeal activity.

Materials and Methods

Plant Materials

The fruit pulp of *A. digitata* was purchased from a local Market in Khartoum, Sudan. The fruit pulps were made into powder, extracted at the Medicinal and Aromatic Plants Institute (MAPRI), the extract weight was 26.18 gm and its yield was 13.9%.

Phytochemical Screening

The freshly prepared extract was subjected to a standard phytochemical screening test for various constituents (Trease and Evans, 1993). The extract was screened for the presence of flavonoids, alkaloids, saponins, tannins, coumarin, triterpenes and glycosides.

Animals

White Albino rats weighing 100-120 g were obtained from the Medicinal and Aromatic Plant, Research Institute, National Centre for Research, Khartoum, Sudan, where they were maintained in a room under standard environmental condition, controlled temperature ($22\pm 25^{\circ}\text{C}$), and relative humidity (60%) and maintained on standard animal pellets and free access to water. Food was withheld for 24 hrs prior to each experiment.

Effect of A. Digitata on Castor Oil – Induced Diarrhoea in Rats

The method described by (Sunil et al., 2001), was followed. Twenty four male and female rats were distributed and divided randomly to 4 groups each group containing

6 rats. Group 1 (control) received 1ml normal saline/rat orally. Group 2 and 3 were treated orally with ethanolic extract of *A. digitata* at doses 250 and 500 mg/kg body weight/rat, respectively. Group 4 received Loperamide at 3mg/kg body weight /rat, it was used as standard antidiarrhoeal drug. After sixty minutes all groups were received castor oil 1ml/rat. Each rat was then housed separately in a cage over clean filter paper and diarrhoea episodes were observed for a period of 24 hours. First defecation time and frequency of defecation were also recorded. Total weight of the feces and inhibition percentage was recorded 4 and 6 hours after the last treatment.

Effect of A. Digitata on Castor Oil – Induced Fluid Accumulation and Electrolyte Secretion

Following the method of (Dicarlo et al., 1994), rats were divided into 5 groups of 6 rats each. Groups 1 and 2 received normal saline, 1 ml/rat, groups 3 and 4 received the ethanolic extract of *A. digitata* at 250 and 500 mg/kg respectively and group 5 received Loperamide at 3mg/kg. One hour after the last treatment castor oil was administered to all groups except group 1, at 1 ml/rat. Two hours later all rats were sacrificed and the small intestines were removed from pylorus to the cecum after ligating the ends. Intestinal contents was collected into graduated cylinder and the volume was measured and reduction percentage was calculated.

Blood was collected from ocular plexus of rats immediately before sacrificed and serum was separated for detection of electrolyte using flame photometer.

Statistical Analysis

All the results were expressed as mean \pm S E. One way analysis of variance (ANOVA) was used for the statistical analysis of data. Duncan's multiple range tests was used for determining the

significance. A probability value of $p < 0.05$ was considered as significant (Snedecor and Cochran, 1989).

Results

Phytochemical Screening

The percentage yield of the ethanol extract of *A. digitata* was 13.901% the results of the phytochemical screening revealed the presence of flavonoids, saponin, alkaloid, triterpenes, tannins, coumarin and glycosides (Table 1).

Effect of A. Digitata on Castor Oil – Induced Diarrhoea in Rats at 4 Hrs

The effect of the fruit pulp extract of *A. digitata* on weight of feces, frequency, accumulation, and inhibition rate at 4 hrs, are presented on table (2).

The ethanolic extract of *A. digitata* administered at the dose of 250, and 500 mg/kg showed 30% and 68% reduction in total weight of the feces, respectively. The reduction in diarrheal episodes was also significant and maximum effect is observed at the dose of 500 mg/kg.

Effect of A. digitata on Castor Oil – Induced Diarrhoea in Rats at 6 Hrs

Total weight of the feces, frequency of diarrhea and inhibition percentage was shown in table 3. After 6 hrs of the last treatment *A. digitata* ethanolic extract produced 60% and 78% inhibition of diarrhea at the dose rate of 250 and 500 mg/kg respectively where as the standard drug Loperamide showed significant reduction in diarrheal weight and recorded 84%. Similarly there was dose dependant decrease in frequency of diarrhoea.

Effect of A. Digitata on Castor Oil – Induced Fluid Accumulation and Electrolyte Secretion

Table 4 and 5 summarized the effect of *A. digitata* on castor oil –induced fluid accumulation and electrolyte secretion. *A.*

ANTIDIARRHOEAL ACTIVITY OF ETHANOLIC EXTRACT OF ...

digitata ethanolic extract at 250 and 500n mg/kg, was found to demonstrate a significant reduction (21% and 24% respectively) in intestinal fluid accumulation due to castor oil, when compared to control group 1, at the same time it produced significant reduction in weight of intestinal

contents. Treatment of rats with castor oil reduced the serum concentration of Na⁺ and K⁺, significantly when compared to the control while the concentration of these electrolytes increased significantly in groups treated with. *A. digitata* and Loperamide.

Table 1: Phytochemical analysis of ethanolic extract of *Adansonia digitata*.

Reaction	Phytochemical constituents	Results
Dragendorff test	Alkaloids	+
Acetic anhydride, Chloroform and sulphuric acid	Sterols	-
Acetic anhydride, Chloroform and sulphuric acid	Triterpens	+
Aluminum chloride and Potassium hydroxide	Flavonoids	+++
Frothing test	Saponins	+++
Potassium hydroxide	Cumarins	+
Gelatin and Ferric chloride	Tannins	+
Sodium picrate	Glycoside	+

+++ High concentration; ++ Moderate concentration; + Trace; - Negative.

Table 2: Effect of *A. digitata* on castor oil induced diarrhoea in rats after 4hrs.

Groups	Total weight of feces (g)	Frequency of diarrhoea	%inhibition of diarrhoea
G1 (castor oil 1ml/rat)	0.58±0.048 ^a	1.83±0.30 ^a	0
G2(<i>A. digitata</i> 250mg/kg+1ml castor oil/rat)	0.42±0.04 ^b	1.16±0.17 ^b	30%
G3(<i>A. digitata</i> 500mg/kg+1ml castor oil/rat)	0.12±0.08 ^c	0.33±0.21 ^c	68%
G4(3mg/kg lopermide+1ml castor oil/rat)	0.04±0.04 ^c	0.16±0.16 ^c	78%

Table 3: Effect of *A. digitata* on castor oil induced diarrhoea in rats after 6hrs.

Groups	Total weight of feces (g)	Frequency of diarrhoea	%inhibition of diarrhoea
G1(castor oil 1ml/rat)	0.41±0.026 ^a	1.66±0.14 ^a	0
G2(<i>A. digitata</i> 250mg/kg+1ml castor oil/rat)	0.30±0.067 ^{ab}	1.16±0.30 ^{ab}	60%
G3(<i>A. digitata</i> 500mg/kg+1ml castor oil/rat)	0.10±0.056 ^b	0.50±0.22 ^c	78%
G4(3mg/kg lopermide+1ml castor oil/rat)	0.055±0.037 ^b	0.33±0.21 ^c	84%

Table 4: Effect of *A. digitata* on castor oil induced fluid accumulation in rats.

Groups	Weight of intestinal contents (g)	Volume of intestinal contents (ml)	%reduction on Wt of intestinal contents
G1(control)	1.51±0.14 ^{ab}	1.10±0.14 ^b	100%
G2(castor oil 1ml/rat)	1.83±0.16 ^a	1.42±0.18 ^a	0
G3(<i>A. digitata</i> 250mg/kg+1ml castor oil/rat)	1.43±0.17 ^{ab}	1.10±0.19 ^{ab}	21%
G4(<i>A. digitata</i> 500mg/kg+1ml castor oil/rat)	1.38±0.14 ^{ab}	0.92±0.10 ^b	24%
G5(3mg/kg lopermide+1ml castor oil/rat)	1.25±0.16 ^b	0.94±0.14 ^b	31%

Table 5: Concentration of serum electrolytes in rats treated with *A. digitata* on castor oil induced diarrhea.

Groups	Na ⁺ (mmol L ⁻¹)	K ⁺ (mmol L ⁻¹)
G1(control)	143.16±1.22 ^a	5.90±0.20 ^a
G2(castor oil 1ml/rat)	111.91±1.7 ^{4b}	2.70±0.28 ^b
G3(<i>A. digitata</i> 250mg/kg+1ml castor oil/rat)	122.75±2.34 ^c	3.90±0.23 ^b
G4(<i>A. digitata</i> 500mg/kg+1ml castor oil/rat)	133.66±2.19 ^{ab}	4.34±0.61 ^c
G5(3mg/kg lopermide+1ml castor oil/rat)	140.00±1.11 ^a	5.79±0.49 ^a

Means in the same column with the same letter are not significantly different (p>0.05).

Discussion

The preliminary phytochemical screening of the extracts of *A. digitata* revealed the presence of flavonoids, triterpenoids, tannins, alkaloids and glycosides. The anti diarrhoeal activity of flavonoids has been ascribed to their ability to inhibit intestinal motility and hydro-electrolytic secretion, which are known to be altered in diarrhoea (Carlo et al., 1993; Rao et al., 1997). *In vitro* and *in vivo* experiments have shown that flavonoids are able to inhibit the intestinal secretory response, induced by PGE₂ (Sanchez et al., 1997). There are reports that flavonoids also modify mucosal permeability and inhibit intestinal peristalsis (Ghazouli and Holzer, 2004), hence helpful in controlling diarrhoea. The presence of tannins (phenolic glycosides) may also contribute to the anti diarrhoeal activity of ethanolic extracts,

since tannins may precipitate the proteins of enterocytes, reduce peristaltic movement and intestinal secretions (Okudo et al., 1989) The ethanolic extract of *A. digitata* administered at the dose of 250 mg and 500 mg/kg showed 30% and 68% inhibition of defecation, respectively, after four hours. The maximum significant (p < 0.05) effect is observed at the dose of 500 mg/kg comparable to control group (castor oil group), and at 6 hours time, the maximum dose of extract (500 mg/kg) showed almost similar antidiarrhoeal activity as that of loperamide (3mg/kg) 78% and 84% respectively. Castor oil induced diarrhoea by increasing the volume of intestinal content by prevention of reabsorption of water. The release of ricinolic acid stimulate release of prostaglandins, which results in stimulation of secretion (Pierce et al., 1971).

Wickens and Lowe (2008), stated that, two main factors attributed to the antidiarrhoeic action of baobab are thought to include the astringent action of the tannins causing an inhibition of osmotic secretions in addition to the anti-inflammatory action of the baobab mucilage on the intestinal mucous membrane. Similarly Gruenwald and Galizia (2005), reported that, the presence of tannins, mucilage, cellulose and citric acid present in the baobab may also have a role to play in the effects of baobab fruit pulp against diarrhoea.

Treatment of rats with castor oil significantly decrease the serum concentration of Na⁺ and K⁺ compared to the control where as treatment of rats with the ethanolic extract of *A. digitata* significantly increase the concentration of Na⁺ and K⁺ in the serum by increasing reabsorption of electrolytes. (Rouf et al., 2003), reported that, castor oil causes motility and secretory diarrhoea and the mechanism involved has been associated with dual effects on gastrointestinal motility as well as on water and electrolyte transport that decreasing Na⁺ and K⁺ absorption across the intestinal mucosa.

The results of this investigation revealed that *A. digitata* contains pharmacologically active substances with antidiarrhoeal properties. These attributes may provide the rationale for the use of *A. digitata* in diarrhoea management by traditional healers.

References

Atta AH, Mounier SM (2004). Evaluation of some medicinal plant extracts for antidiarrhoeal activity. *Phytother. Res.*, 19: 481-485.
 Awouters F, Niemegeers CJE, Lenaerts FM and Janseen PAJ (1978). Delay of castor oil diarrhoea in wistar rats; a new way to evaluate

inhibitors of prostaglandin biosynthesis. *J. Pharm. Pharmacol.*, 30: 41-45.
 Laloo D, Hemalatha S (2011). Ethnomedicinal plants used for diarrhea by tribals of Meghalaya, Northeast India *Phcog. Rev.*, 5: 147-154.
 Carlo D, Autore G, Izzo G, Maibline AAP, Mascolo N, Viola P, Diurno MV, Capasso F (1993). Inhibition of intestinal motility and secretion by flavonoids in mice and rats: structure activity relationships. *J. Pharm. Pharmacol.*, 45: 1054-1059.
 Diop AG, Sakho M, Dornier M, Cissé M, Reynes M (2005). Le baobab African (*Adansonia digitata* L.): principes caractéristiques et utilisations. *Fruits.*, 61: 55-69.
 El-Kamali HH, El-Khalifa KF (1999). Folk medicinal plants of riverside forests of the Southern Blue Nile district, Sudan. *Fitoterapia.*, 70: 493-497.
 Ezekwesili CN, Obiara KA, Ugwu OP (2004). Evaluation of anti-diarrhoeal properties of crude aqueous extract of *Ocimum gratissimum* L. (Labiatae) in rats. *Biochem.*, 16: 122-131.
 Farthings MJG (2002). Novel targets for the control of secretory diarrhea. *Gut* 50: 15-18.
 Kamel G, Peter H (2004). Inhibition of Guinea Pig intestinal peristalsis by the flavonoids Quercetin, Naringenin, Apigenin and Genistein. *Int. J. Exp. Clin. Pharmacol.*, 70: 5-14.
 Gruenwald J, Galizia M (2005). *Adansonia digitata* Market Brief in the European Union for selected natural ingredients derived from native species. *The United Nations Conference on Trade and Development (UNCTAD)*. p. 35.
 Okudo T, Yoshoda T, Hatano T (1989). New methods of analyzing tannins. *J. Nat. Prod.*, 52: 1-31.
 Pierce NF, Carpenter CCJ, Elliot HZ, Greenough WB (1971). Effect of prostaglandins, theophylline and cholera exotoxin upon transmucosal water and electrolyte movement in canine jejunum. *Gastroenterol.*, 60: 22-32.
 Rao VSN, Santos FA, Sobreika TT, Souza MF, Melo LL, Silveira ER (1997). Investigations on the gastroprotective and antidiarrhoeal properties of ternatin, a tetramethoxy flavone from *Egletes viscosa*. *Planta Med.*, 63: 146-149.
 Rouf AS, Islam MS, Rahman MT (2003). Evaluation of antidiarrhoeal activity of *Rumex maritimus* roots. *J. Ethnopharmacol.*, 84: 307-310.
 Medina SDF, Galvez J, Gonzalez M, Zarzuelo A, Barrett KE (1997). Effects of quercetin on epithelial chloride secretion. *Life Sci.*, 61: 2049-2055.
 Sidibé M, Scheuring JF, Tembely D, Sidibé MM, Hofman P, Frigg M (1996). Baobab –

- Homegrown vitamin C for Africa. *Agroforestry Today*, 8: 13-15.
- Snedecor GW, Cochran WG (1989). *Statistical Methods*, 8th ed., Ames, Iowa State University Press, Iowa, USA.
- Sunil B, Bedi K, Singla A, Johri R (2001). Antidiarrhoeal activity of piperine in mice. *Planta Med.*, 67: 284-287.
- Snyder JD, Merson MH (1982). The magnitude of the global problem of acute diarrhoeal disease: a review of active surveillance data. *B. World Health Organ.*, 60: 605-613.
- Tijani AY, Okhale SE, Salawu TA, Onigbanjo HO, Obianodo LA, Akingbasote J, Aalawu OA, Okogun SJI, Kunle FO, Emeje M (2009). Antidiarrhoeal and antibacterial properties of crude aqueous stem bark extract and fractions of *Parkia biglobosa* (Jacq.) R. Br. Ex G. Don. *Afr. J. Pharm. Pharmacol.*, 3(7): 347-353.
- Trease GE, Evans MC (1983). *Text book of Pharmacognosy*, 12th ed. Bailliere, Tindail, London, pp. 343-383.
- Wickens GE, Lowe P (2008). *The Baobabs: Pachycauls of Africa, Madagascar and Australia*, Berlin, Germany, Springer.