A preliminary investigation on the effects of polpala (*Aerva lanata*) on the structure and function of urinary tract of rats

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Abstract

Polpala (*Aerva lanata*) is a medicinal plant that is extensively used for the treatment of renal disease in Ayurvedic medicine. This preliminary investigation was carried out to evaluate the diuretic effect of the crude aqueous plant extract and high and low molecular mass fractions of this herb, using the hydrated rat assay technique. Possible toxicological effects on renal structure and function, mediated by long term administration of aqueous plant extract was also studied using two concentrations of both fresh and dried plant material, made according to Ayurvedic system of medicine in Sri Lanka. Urine output, osmolality and electrolyte excretion were measured to evaluate diuretic activity. Creatinine clearance and histopathological studies were done to evaluate toxicological effects on renal structure and function.

Crude aqueous plant extract showed a diuretic activity with marked increase in urine output, urine osmolality and urinary potassium excretion. Low molecular mass fraction of dried plant material showed a significant diuretic effect while the high molecular fraction did not have the same effect. Investigations also showed that long term administration of the aqueous extract of this plant had no toxicological effects on renal structure and function of rats.

Key words: polpala, *Aerva lanata*, diuretic activity, toxicological effects.

Introduction

Aerva lanata Linn. (Sinhala: polpala, polkudupala, pahanabaya, puswenna. Tamil: sirupulai, thengapookeerai) which belongs to the family Amaranthaceae is a medicinal plant which is used by Ayurvedic practitioners in Sri Lanka to treat various conditions such as lithiasis, headache and renal disease (1). This plant is used in other countries too as a diuretic and anthelmintic. It is also used in malarial fever, diabetes and gonorrhea (2, 3, 4, 5). The whole plant or parts of it is used as a diuretic, herbal drink, tea, porridge, extract or as a decoction with other herbs (5).

Diuretic activity of this herb had been subjected to research by several research groups in the recent past. Two had been carried on humans and one on rats (6, 7, 8). Experiments carried out by two research groups showed the diuretic effect of *A. lanata* when given to humans and rats. Also *A. lanata* is not prescribed by Ayurvedic physicians as a decoction or as a herbal drink for more than 7 days as there is a general belief that it can damage the renal structure. However, this is not scientifically proved

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and people use *A. lanata* as a herbal drink or porridge according to their own interest.

This preliminary investigation was therefore conducted to evaluate the diuretic activity of crude aqueous extracts of dried and fresh plants and high and low molecular mass fractions of the dried plant using the hydrated rat assay technique (9). Also investigations were conducted to study any possible toxicological effects on renal structure and function mediated by the long term administration of aqueous extract of *A. lanata*.

Material and Methods

Animals

Rats for the experiments were obtained from an inbred colony maintained at the animal house of the Faculty of Medicine, Colombo. They were Sprague-Dawley male rats, in the age of 3-4 months, weighing 200 ± 15 g. The animals were housed in cages under tropical laboratory conditions (room temperature $28 \pm 3^\circ\text{C}$) and provided a standard laboratory diet and water. Ethical review for the experiments was obtained from the Ethical Review Committee of the Faculty of Medicine, Colombo.

Plant material and sample preparation

Whole plants of *A. lanata* were collected from waste and cultivated grounds in the Gampaha District, Western Province, Sri Lanka. Botanical identity of the plant was determined using herbarium specimens available at the Department of Botany, University of Colombo, Sri Lanka.

In this investigation, *A. lanata* was used in the form of fresh plant as well as in the dried form. Dried form was prepared by keeping

fresh plant (washed and cut) in an oven at 60°C until a constant weight was obtained. The plant extract was prepared according to the method used by ayurvedic practitioners by boiling the samples with water until the volume is reduced to half. Amounts of 50 g and 100 g of *A. lanata* used in the present study were based on amounts used in previous investigations by Udupihille and Jiffery (6). The extracts were therefore prepared by boiling either 50 g or 100 g of the plant in 400 ml of water until the volume was halved. Then the extracts were cooled and filtered through Whatman filter paper No. 1. Four solutions were prepared according to the above procedure. They were

Solution 1 – Extract of *A. lanata* using 50g of fresh plant in 200 ml (AL50FR)

Solution 2 – Extract of *A. lanata* using 100g of fresh plant in 200 ml (AL100FR)

Solution 3 – Extract of *A. lanata* using 50g of dried plant in 200 ml (AL50DR)

Solution 4 – Extract of *A. lanata* using 100g of dried plant in 200 ml (AL100DR)

Findings of the experiments with rats treated with *A. lanata* were compared with those from animals administered water, (negative control) and frusemide (used as a reference diuretic drug, positive control).

Separation of high and low molecular mass fractions

One liter of 100 g/200 ml dried plant extract was prepared according to the method mentioned above. Freeze-dried product of this extract was obtained by means of a freeze-dryer and yielded 30 g. Ten grams of the freeze-dried powder was reconstituted in 250 ml of distilled water and transferred into a previously prepared dialysis tube in 50 ml portions and dialysed against distilled

water (dialysis medium) contained in a flask, and continuously stirred using a magnetic stirrer. The dialysis medium was replaced several times with distilled water until it remained colorless. Pooled samples of the dialysis medium collected during the dialysis period were freeze-dried to obtain the low molecular dry powder (6.28 g). The high molecular mass fraction, which remained inside the tube, was also freeze-dried to obtain a dry powder of high molecular mass compounds (2.94 g). Two solutions were prepared after reconstituting high and low molecular dry powder fractions in water. These solutions contained dry powder equal to 100 g/200 ml extract.

Experiment 1 – Screening for diuretic activity of crude aqueous plant extract

The method used to study the diuretic activity of the extract was similar to the method used by Sripanidkuichi *et al* (10). Rats were fasted overnight with free access to drinking water. They were given 18 ml/kg body weight of water one hour prior to the following experiment. This dose was calculated on the basis of weight ratio between human and rats (11). Water or food was not allowed during the experimental period (8, 9).

Thirty six rats were randomly divided into six groups (n=6 for each group) and rats in the different groups were treated as follows:

Group 1 – Distilled water (control)

Group 2 – AL 50 FR

Group 3 – AL 100 FR

Group 4 – AL 50 DR

Group 5 – AL 100 DR

Group 6 – Frusemide (20 mg/kg body weight of rat)

All solutions were administered orally via a metal oro-gastric tube. All were administered doses of 18 ml/kg body weight of rat (11). Each animal was then placed in separate metabolic cages and urine was collected into tubes fitted to the funnel at the bottom of the cage. Urine output was determined hourly for a period of four hours. Urine osmolality was measured using a micro-osmometer. Urinary electrolytes (Na^+ and K^+) were determined by flame photometry.

Experiment 2 – Screening for diuretic activity of high and low molecular mass fractions

Forty rats were randomly divided into 4 groups. Rats in the 4 groups were treated as follows.

Group 1 – Distilled water

Group 2 – Low molecular mass fraction

Group 3 – High molecular mass fraction

Group 4 – Frusemide

These animals were subjected to the same experimental procedure as described under Experiment 1.

Experiment 3 – Screening for the effect of long term administration of *A. lanata* on renal structure and function of rats

Eighteen male rats were randomly assigned to six groups (sample size was limited to 3 rats in each group due to limited availability of resources). The animals in each group were fed with the different solutions as given in Experiment 1. All treatments were given doses of 18 ml/kg of rat, which was comparable to that given to humans on the basis of relative weights (11). All treatments were given once daily orally for 30 days by a metal oro-gastric tube (12).

Effect on renal function

On the 30th day of the experiment, test animals were kept separately in metabolic cages and urine was collected for 24 hours. On the 31st day before the dissection, blood was drawn from the tail vein. Blood was centrifuged and serum was separated. Serum and urine samples were used to determine the creatinine clearance of each rat. Creatinine levels in serum and urine were determined spectrophotometrically using a creatinine reagent kit (Cat. no. CR 510, RANDOX Laboratories Ltd. UK). Creatinine clearance was calculated to study the effect of long term administration of *A. lanata* on the renal function of rats.

Effect on renal structure

On the 31st day of the experiment animals were anaesthetized by ether inhalation. Then animals were dissected through a mid ventral laparotomy and their kidneys, bladder and ureters were harvested and fixed separately in Duboseq Brazil Solution (DBS). These specimens were processed, embedded in wax and 5 μ m sections were cut. These thin sections of harvested organs were stained with haematoxylin and eosin. The stained sections were examined light microscopically for histological changes.

Results

Diuretic activity of crude plant extract

The hourly mean total urine output, urine osmolality and electrolyte excretion during the 4 hours of experimental time period are summarized in Tables 1 and 2. It is evident that there is an increase in the urine output in the groups of rats fed with aqueous plant extracts compared to those in the negative control group (water). When compared with the negative controls fed only water, in groups of rats fed with 100 g/200 ml of dried and fresh plant extracts, a significant

increase in the mean total urine output was evident after the 3rd hour. In the frusemide treated rats, a significantly increased urine output was evident at the end of the 2nd and 3rd hours.

Animals treated with plant extracts and frusemide showed higher urinary osmolality compared to the negative control group during each hour. In the 1st hour and 3rd hour, rats fed with 100 g/200 ml of both fresh and dried plant extracts showed a significant increase in urine osmolality. The highest urine osmolality was observed in the 4th hour in all groups except the rats in the group fed with AL50DR. However, it was not statistically significant.

Na⁺ excretion was significantly high only in animals treated with frusemide. In the 4th hour AL50FR and AL100FR showed slight increase in Na⁺ excretion but that was not significant. There was a significant increase in K⁺ excretion compared to the negative control during 1st hour following different treatments. Also there was a significant increase in excretion of K⁺ during the 3rd hour in rats who were given AL100FR, AL50DR and AL100DR solutions.

Diuretic activity of high and low molecular mass fractions

The hourly mean total urine output, urine osmolality and electrolyte excretion during the 4 hours of experimental time period are summarized in Tables 3 and 4. Rats fed with low molecular mass fraction had significant increase in urine output after 2nd, 3rd and 4th hours, whereas rats fed with high molecular mass fraction had a significant increase in urine output only after 2nd and 3rd hours compared to the rats in the negative control group. Urine osmolality was also significantly high in the group of rats fed with low molecular mass fraction. Significantly increased electrolyte excretion was also observed in the group of rats fed with the low molecular mass fraction.

Effect of long term administration of *A. lanata* on renal structure and function

Mean creatinine clearance and the light microscopic appearance of the specimens following long term administration of different treatments are shown in Table 5. Following long term administration of the different treatments there was no significant

difference in serum creatinine clearance, compared to values in the negative control group.

When histological sections of the specimens (kidneys, ureters and bladder) of rats treated with *A. lanata* extracts were compared with the specimens of the control groups, there was no remarkable difference.

Table 1. Mean total urine output (UOP) after each hour and urine osmolality (Uos) in rats after feeding of crude aqueous plant extracts of *A. lanata*

Treatment	No. of animals		UOP (ml/rat) and Uos (mOsm/kg)			
			1 h	2h	3h	4h
Water	6 ^a	UOP	1.72 ± 0.25	2.45 ± 0.27	2.45 ± 0.27	2.84 ± 0.26
	6 ^b	Uos	70.2 ± 5.49	172 ± 22.7	272.3 ± 22.5	465 ± 13
AL50FR	6 ^a	UOP	1.85 ± 0.43	2.74 ± 0.20	3.37 ± 0.2	0.03 ± 0.41
	6 ^b	Uos	113.2 ± 13.3*	234 ± 8.84	435 ± 12	760 ± 8.0
AL100FR	6 ^a	UOP	2.12 ± 0.37	2.52 ± 0.31	3.83 ± 0.44*	3.83 ± 0.44
	6 ^b	Uos	143.3 ± 12.3*	397 ± 13	625 ± 70.6*	1096
AL50DR	6 ^b	Uos	2.20 ± 0.49	2.98 ± 0.31	3.37 ± 0.19	3.37 ± 0.18
	6 ^b	Uos	82.6 ± 1.69	310.3 ± 95	478 ± 32.2*	♦
AL100DR	6 ^b	Uos	2.02 ± 0.46	3.28 ± 0.78	3.95 ± 0.87*	4.90 ± 0.56
	6 ^b	Uos	176.2 ± 22.0*	371.3 ± 23.1	852.2 ± 169*	853.3 ± 15.4
Frusemide	6 ^a	UOP	2.48 ± 0.24	4.90 ± 0.41*	6.03 ± 0.66*	6.76 ± 0.86
	6 ^b	Uos	161.7 ± 29.1*	259.2 ± 21.1*	344.4 ± 12	532 ± 62.4

1. * Significantly different from the negative control (water) - $p < 0.05$, Mann-Whitney U test.

2. AL50FR – 50 g/200 ml of fresh *A. lanata* crude extract

AL100FR – 100 g/200ml of fresh *A. lanata* crude extract

AL50DR – 50 g/200 ml of dried *A. lanata* crude extract

AL100DR – 100 g/200 ml of dried *A. lanata* crude extract

3. ♦ Rats of this group did not pass urine at all.

4. ^a The urine output of some rats was negligible at times and some did not pass urine at all in which case the urine output of such rats was considered as zero.

5. ^b Initially number of rats in each group was six ($n = 6$). The number of samples considered in the analysis of urine osmolality due to the reason mentioned above was

Water – 1h ($n = 6$), 2h ($n = 4$), 3h ($n = 4$), 4h ($n = 2$)

AL50FR – 1h ($n = 5$), 2h ($n = 4$), 3h ($n = 2$), 4h ($n = 2$)

AL100FR – 1h ($n = 4$), 2h ($n = 2$), 3h ($n = 3$), 4h ($n = 1$)

AL50DR – 1h ($n = 5$), 2h ($n = 4$), 3h ($n = 4$)

AL100DR – 1h ($n = 5$), 2h ($n = 3$), 3h ($n = 4$), 4h ($n = 3$)

Frusemide – 1h ($n = 6$), 2h ($n = 6$), 3h ($n = 5$), 4h ($n = 4$)

Table 2. Urinary electrolyte excretion (Na⁺ and K⁺) in rats after feeding of crude aqueous plant extracts of *A. lanata*

Treatment	No. of animals	Na ⁺ and K ⁺ (mmol/l)				
			1 h	2h	3h	4h
Water	6 ^a	Na ⁺	6.3 ± 2.6	13 ± 8.4	23.7 ± 12.7	3.7 ± 3.8
		K ⁺	7.82 ± 1.0	26.2 ± 2.7	34 ± 2.8	55.8 ± 2.2
AL50FR	6 ^a	Na ⁺	10.6 ± 2.2	16.8 ± 5.9	38.5 ± 0.5	69.5 ± 5.5
		K ⁺	16.2 ± 2.8*	45.3 ± 1.4	82 ± 2.0	139.5 ± 19
AL100FR	6 ^a	Na ⁺	14 ± 3.7	19 ± 4	25 ± 4.9	67
		K ⁺	25.6 ± 5.6*	27.1 ± 15.4	119.8 ± 13*	194
AL50DR	6 ^a	Na ⁺	3.6 ± 0.50	6 ± 1.5	16 ± 2.4	♦
		K ⁺	23.3 ± 0.95*	122.8 ± 27	111.3 ± 0.3*	♦
AL100DR	6 ^a	Na ⁺	7.8 ± 3.0	6.3 ± 3.4	16.8 ± 8	14.7 ± 5.4
		K ⁺	47.2 ± 9.2*	131.3 ± 12.9	192.3 ± 1.7*	173 ± 21
Frusemide	6 ^a	Na ⁺	40.5 ± 13*	67.7 ± 9.6*	111.7 ± 12*	03 ± 8.5*
		K ⁺	17 ± 0.9*	33.8 ± 4.3	39.9 ± 4.9	81.9 ± 29

1. * Significantly different from the negative control (water) – p<0.05, Mann-Whitney U test.

2. ♦ Rats of this group did not pass urine at all.

3. ^a Initially number of rats in each group was six (n= 6). The urine output of some rats was negligible at times and some did not pass urine at all in which case the urine output of such rats was considered as zero. These samples were not considered in the analysis.

Table 3. Mean total urine output (UOP) after each hour and urine osmolality (Uos) in rats after feeding of low and high molecular mass fractions of *A. lanata*

Treatment	No. of animals	UOP (ml/rat) and Uos (mOsm/kg)				
			1 h	2h	3h	4h
Water	10 ^a	UOP	1.67 ± 0.3	2.44 ± 0.20	2.91 ± 0.20	3.30 ± 0.19
	10 ^b	Uos	92 ± 10.7	223.2 ± 30.9	351.2 ± 113	421 ± 44.6
Low mol. mass fraction	10 ^a	UOP	2.46 ± 0.58	4.14 ± 0.38*	4.78 ± 0.34*	5.38 ± 0.35*
	10 ^b	Uos	209.3 ± 26*	447.9 ± 53*	749 ± 47	899.8 ± 41.1*
High mol. mass fraction	10 ^a	UOP	2.25 ± 0.32	3.38 ± 0.26*	4.05 ± 0.34*	4.19 ± 0.38
	10 ^b	Uos	106 ± 6.3	188.6 ± 22.6	497.5 ± 106	631.3 ± 40.4
Frusemide	10 ^a	UOP	1.73 ± 0.34	3.94 ± 0.27*	5.05 ± 0.48*	6.04 ± 0.38*
	10 ^b	Uos	141.2 ± 4.03*	238.4 ± 16	352.8 ± 25	58.5 ± 172

1. * Significantly different from the negative control (water) – p<0.05, Mann-Whitney U test.

2. ^a The urine output of some rats was negligible at times and some rats sometimes did not pass urine at all in which case the urine output of such rats was considered as zero.

3. ^b Initially number of rats in each group was ten (n= 10). The number of samples considered in the analysis of urine osmolality due to the reason mentioned above was

Water – 1h (n= 8), 2h (n= 6), 3h (n= 4), 4h (n= 3)

Low molecular mass fraction – 1h (n= 8), 2h (n= 7), 3h (n= 2), 4h (n= 5)

High molecular mass fraction – 1h (n= 9), 2h (n= 7), 3h (n= 6), 4h (n= 2)

Frusemide – 1h (n= 8), 2h (n= 8), 3h (n= 6), 4h (n= 6)

Table 4. Urinary electrolyte excretion (Na⁺ and K⁺) in rats after feeding of low and high molecular mass fractions of *A. lanata*

Treatment	No. of animals		Na ⁺ and K ⁺ (mmol/l)			
			1 h	2h	3h	4h
Water	10 ^a	Na ⁺	3.6 ± 0.98	5.5 ± 0.96	27 ± 9.3	28 ± 7.7
		K ⁺	9.02 ± 1.0	26.2 ± 5.8	37.6 ± 3.0	6.2 ± 22.5
Low mol. mass fraction	10 ^a	Na ⁺	37.3 ± 3.7*	52.3 ± 10.3*	44.5 ± 4.5	43.2 ± 6.6
		K ⁺	42.6 ± 2.58*	112.9 ± 17.9*	147.5 ± 2.5*	238.5 ± 19.7*
High mol. mass fraction	10 ^a	Na ⁺	7.44 ± 1.6	18.9 ± 5.24	40.4 ± 9.5	64 ± 24
		K ⁺	8.62 ± 0.95	19.8 ± 3.4	62.3 ± 9.2	66.7 ± 1.4
Frusemide	10 ^a	Na ⁺	31.8 ± 10.5*	60.3 ± 10*	116.3 ± 9.6*	113.8 ± 6.5*
		K ⁺	14.7 ± 1.4*	24.4 ± 4.1	40.3 ± 3.5	90.9 ± 19.2

- * Significantly different from the negative control (water) - p<0.05, Mann-Whitney U test.
- ^a Initially number of rats in each group was ten (n= 10). The urine output of some rats was negligible at times and some did not pass urine at all in which case the urine output of such rats was considered as zero. These samples were not considered in the analysis.

Table 5. Mean creatinine clearance and effects on renal structure of rats following long term administration of crude aqueous plant extracts of *A. lanata*

Solution	Creatinine Clearance (ml/min)	Effects on renal structure (Light microscopic appearance)
Water	0.425 ± 0.096	Kidneys, bladder and ureters were normal in appearance
AL50FR	0.363 ± 0.111	Kidneys, bladder and ureters were normal in appearance
AL100FR	0.412 ± 0.017	Bladder and ureters did not show any abnormality. Kidneys showed focally congested glomeruli
AL50DR ^a	0.414 ± 0.191	Bladder and ureters did not show any abnormality. Kidneys showed focally congested glomeruli
AL100DR	0.140 ± 0.042	Bladder and ureters did not show any abnormality. Kidneys showed focally congested glomeruli
Frusemide	0.424 ± 0.070	Kidneys, bladder and ureters were normal in appearance

- AL50FR – 50 g/200 ml of fresh *A. lanata* crude extract
AL100FR – 100 g/200ml of fresh *A. lanata* crude extract
AL50DR – 50 g/200 ml of dried *A. lanata* crude extract
AL100DR – 100 g/200 ml of dried *A. lanata* crude extract
- ^a One animal died on the 16th day of the experiment.

Discussion

Results of this study show that crude aqueous plant extract of *A. lanata* has a diuretic effect on rats when given orally. Our results are comparable with the findings of Prasad *et al.*, (8). Also our results show that the treatment with crude aqueous plant extract has a rapid onset of action and this effect remains for about three hours.

When the diuretic effect was studied further, using high and low molecular mass fractions of *A. lanata*, it shows that low molecular mass fraction is more effective as a diuretic substance compared to the high molecular mass fraction. This difference may be due to the presence of more ingredients responsible for the diuretic effect of *A. lanata* in the low molecular mass fraction.

The precise mechanism of the diuretic effect of the plant extract is not clear. However, the diuretic effect is not similar to the effect produced by frusemide. It may be similar to the effect produced by thiazides as the plant extract too has more effect on K^+ excretion than Na^+ excretion.

Further, the results of this preliminary investigation indicate that administration of *A. lanata* for one month has no toxic effect in the urinary tract of rats.

Conclusion

This study demonstrates that *A. lanata* plant extract has a diuretic effect. When low and high molecular mass fractions were used, it is evident that low molecular mass fraction has more diuretic activity compared to the high molecular mass fraction. Also administration of *A. lanata* for a period of one month has no significant toxic effects in the urinary tract of rats.

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Key words: *Aerva lanata*, diuretic, urinary tract, rats.

Introduction

Aerva lanata (Asteraceae) is a common plant in the tropics and subtropics. It is known for its diuretic and laxative properties. The plant is used in traditional medicine for the treatment of urinary tract infections and other ailments. The present study aims to evaluate the diuretic effects of *Aerva lanata* on the structure and function of the urinary tract of rats.

In Sri Lanka, *Aerva lanata* is commonly found in the wet zone and is used in traditional medicine for the treatment of urinary tract infections and other ailments. The plant is known for its diuretic and laxative properties. The present study aims to evaluate the diuretic effects of *Aerva lanata* on the structure and function of the urinary tract of rats. The study was conducted in the wet zone of Sri Lanka, where the plant is commonly found. The results of the study will be used to develop a standardized protocol for the use of *Aerva lanata* in the treatment of urinary tract infections and other ailments.