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Full Length Research Article

## Anti-Inflammatory and Anti-nociceptive Effects of Ethanolic Extract of *Setaria megaphylla* Leaves in Rodents

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### ABSTRACT

The ethanolic leaf extract of *Setaria megaphylla* (100-300 mg/kg) was investigated for pharmacological properties against egg white albumin - induced inflammation, Chemical as well as thermal- induced pain. The extract demonstrated a dose – dependent anti- inflammatory and antinociceptive activities. These activities were comparable to that of ASA (100mg/kg). The leaf extract possess anti inflammatory and analgesic properties, which can be exploited in health care

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**Keywords:** *Setaria megaphylla*, anti inflammatory, analgesic.

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## INTRODUCTION

*Setaria megaphylla* (steud) Dur schinz (family-Poaceae) also called broad leafed brittle grass is a very tall, robust, tufted, perennial grass used mainly as pasture. It occurs in tropical and subtropical areas of Africa, America and India where there is high rainfall (Van oudtshoorn, 1999, Lowe, 1989). The plant is used traditionally by the Ibibios in Akwa Ibom State of Nigeria, in the treatment of various ailments such as hemorrhoids, Urtheritis and diabetes. The plant has been reported to possess antiplasmodial activity in vitro (Clarkson *et al*, 2004) as well as antidiabetic activity (Okokon and Antia, 2005). The leaves have been reported by Okokon and Antia (2005) to contain flavonoids, terpenes, saponins, tannins, anthraquinones, cardiac glycosides, while alkaloids are reported to be absent. However, no anti-inflammatory and analgesic effect of the leaves has been reported earlier in the literature. The present study, therefore, was to establish if the leaves of *S. megaphyglia* have any analgesic/ant-inflammatory effect especially because of its ethnomedical uses in the treatment of inflammatory cases

## MATERIALS AND METHODS

### Plant materials

Fresh leaves of *Setaria megaphylla* were collected in November, 2004 at Anwa forest in Uruan, Akwa Ibom State, Nigeria . The plant was identified and authenticated by Dr. Margaret Bassey, a taxonomist in the Department of Botany, University of Uyo, Uyo, Nigeria. Hebarium specimen was deposited at Faculty of Pharmacy Hebarium with voucher no. FPHUU 221. The fresh leaves (2kg) of the plant were dried on laboratory table for 2 weeks and reduced to powder. The powder 100g was macerated in 95% ethanol (300ml) for 72 hours. The liquid filtrate obtained was concentrated in vacuo at 40°C. The yield was 2.08% w/w. The extract was stored in a refrigerator at 4°C until used for experiment reported in this study.

### Animals:

Albino swiss mice (21-28g) of either sex were obtained from the University of Uyo animal house.

They were maintained on standard animal pellets and water ad libitum. Permission and approval for animal studies were obtained from the College of Health Sciences Animal Ethics committee, University of Uyo.

### Anti-Inflammatory Test

The test was carried out using a phlogistic agent – induced mouse hind paw oedema as a model of acute inflammation (Winter *et al*, 1963) The phlogistic agent employed in this study was fresh egg-albumin (Akah and Nwambie, 1994). Adult Swiss mice of either sex (21 - 28g) were used after a 12h fast. Animals were deprived of water only during the experiment. Inflammation of the hind paw was induced by injection of 0.1ml of fresh egg white into the subplantar surface of the right hind paw of the mice. Paw diameters were measured immediately before the administration of the phlogistic agent and 3 hours thereafter. For routine drug testing, the increase in paw diameter 3 hours after administration of the phlogistic agent was adopted as the parameter for measuring inflammation (Winter *et al*, 1962). Thus (inflammation) was assessed as the difference between zero time paw diameter and that 3 hours after administration of phlogistic agent (Hess and Milonig, 1972). The extract (100,200 and 300 mg/kg) were administered i.p 1 hour before inducing inflammation. Control mice received equivalent amount of normal saline and the reference group administered Acetic salicylic acid (ASA) 100mg/kg. Average oedema ( $C_t - C_0$ ) and percent inhibition of oedema were calculated for each dose (Oriowo,1982; Akah and Njike, 1990).

### Acetic acid – induced writhing in mice

The analgesic activity of ethanolic leaf extract of *Setaria megaphylla* was measured against acetic acid induced writhmic movements in mice (Collier, 1968; Santos *et al*, 1994), consisting of the contraction of abdominal muscle together with the stretching of hind limbs. The extract at doses of 100, 200 and 300mg/kg and ASA 100mg/kg and normal saline 5ml/kg were administered intraperitoneally to the respective groups (n=5) of the 18hours fasted mice. Thirty minutes later, 0.5ml of 2% v/v acetic acid solution was given to each animal intraperitoneally.

The animals were then placed in separate plastic cages and closely observed at 10 minutes interval for 50 minutes. The number of writhes for each animal was counted. Percent inhibition of pain for each group was calculated by comparing the total writhetic number of writhes in the group over the 50 minutes period with the number of writhes in the control group over the same time period. Data were calculated according to the following formula.

$$\% \text{ Inhibition} = \frac{W_t - W_c}{W_c} \times 100$$

Where,

$W_t$  = Mean number of writhes for the test group

$W_c$  = Mean number of writhes for the control group

### Thermally – induced pain in mice

The effect of extract on hot plate – induced pain was investigated in adult mice. The hot plate test was used to measure response latencies according to the method of Vaz *et al.*, (1996, 1997). The animals were divided into 5 groups of 5 mice each. Group 1 mice served as the control and received only saline. Groups 2, 3 and 4 were pre-treated with 100, 200 and 300mg/kg *S. megaphylla* extract i.p respectively, 30min prior to the placement on the hot plate, while group 5 animals received 100mg/kg of ASA by i.p route. The hot plate was set at  $45 \pm 1^\circ\text{C}$  and animals were placed into a glass beaker of 50cm diameter on the heated surface and the time(s) between placement and shaking or licking of the paws or jumping was recorded as the index of response latency.

### Statistical analysis

Data are expressed as mean  $\pm$  SEM for n numbers of experiment. Statistical comparisons and significance levels were analyzed with student's t – test. A 'p' value less than 0.05 was considered as significant

## RESULTS

### Fresh Egg Induced Inflammation in Mice

The extract showed significant ( $P < 0.05$ ) anti-inflammatory activity against acute inflammation. (Table 1). It suppressed in a dose related manner the increase in the mice paw edema caused by egg albumin. The inhibition by the extract was maximal after 3 hours of administration of phlogistic

agent. The effect which was significant when compared to control was comparable to that of the standard drug, ASA.

### Acetic Acid – Induced Writhing In Mice

The extract (100 – 300mg/kg) dose – dependently reduced acetic acid induced abdominal contractions and stretching of hind limbs. The reduction was significant ( $P < 0.05$ ) (Table 2) when compared to control. The analgesic effect was comparable to that of ASA.

**Table 1:**

Effect of *Setaria megaphylla* on fresh egg albumin induced inflammation in rats.

Treatment	Dose mg/kg	Paw Diameter cm	Inhibition %
Control		$0.69 \pm 0.03$	-
<i>S. megaphylla</i> extract	100	$0.29 \pm 0.02^*$	57.97
	200	$0.28 \pm 0.03^*$	59.42
	300	$0.27 \pm 0.03^*$	60.86
ASA	100	$0.26 \pm 0.01^*$	62.31

Results are expressed as mean  $\pm$  SEM (n=5) \* $P < 0.05$  significantly different from control.

**Table 2 .**

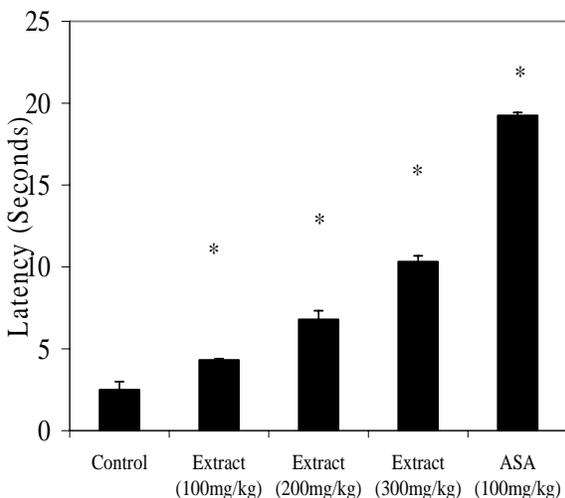
Analgesic activity of ethanolic leaf extract of *Setaria megaphylla* on acetic acid induced writhing in mice.

Treatment	Dose mg/kg	No. of Writing	Percent activity against acetic acid induced pain
Control		237.3 $\pm 18.50$	
<i>S. megaphylla</i> extract	100	$161.7 \pm 5.06^*$	31.85
	200	$119.4 \pm 3.18^*$	49.96
	300	$40.5 \pm 7.21^*$	82.93
AS A	100	$53.0 \pm 2.80^*$	77.66

Results are expressed as mean  $\pm$  S.D (n=5) \* $P < 0.05$  significantly different from control

### Thermally- Induced Pain In Mice

Administration of *S. megaphylla* extract (100 – 300mg/kg i.p) elicited a dose – dependent increase in the latency response in the hot plate test. These increases in latency responses (analgesic effect) were statistically significant ( $P < 0.05$ ) (Fig. 1), when compared to control.



**Fig. 1.**

Effect of *S. megaphylla* on thermally induced pain in rats.  
\* $P < 0.05$  significantly different from control

### DISCUSSION

The ethanolic leaf extract of *S. megaphylla* significantly reduced edema of the mouse hind paw induced by fresh egg albumins. This dose - dependent action; was comparable to that of an acetylsalicylic acid, a cyclo-oxygenase inhibitor (Singh *et al*, 1996). Flavonoids which are some of the constituents of the extract have anti-inflammatory property (Trease and Evans, 1989; Parmer and Gosh, 1978), Edema is attributed to the release of histamine, 5-HT, Kinins and prostaglandins (Vane and Booting, 1987, Larsen and Henson, 1983) and the anti-inflammatory action of this extract may be due to the inhibition of the release of the above mentioned autocooids.

The leaf extract was also found to possess significant ( $P < 0.05$ ) dose and time - dependent analgesic activity against chemical and thermal –

induced pains. Acetic acid causes inflammatory pain by inducing capillary permeability (Amico-Roxas *et al*, 1984) while hot plate-induced pain indicates narcotic involvement (Turner, 1965; Besra *et al*, 1966). The ability of the extract to show significant effect in these two types of pain induction suggest that its analgesic effect may in part be related to its anti-inflammatory and narcotic properties.

Therefore, the result obtained in this study shows that *S. megaphylla* possess anti-inflammatory and analgesic properties which are probably mediated via inhibition of various autocooids formation and release. Further studies are needed to elucidate the exact mechanism by which *S. megaphylla* inhibits inflammation and pains

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