



IN VIVO ANTI INFLAMMATORY AND ANTI ARTHRITIC ACTIVITY OF ETHANOLIC EXTRACT OF *ASPARAGUS RACEMOSUS* ROOTS

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ABSTRACT

The present study is aimed to appraise the anti-inflammatory and anti-arthritic activity of ethanolic extract of *Asparagus racemosus* roots belonging to family Liliaceae. Carrageenan is used to induce inflammation and Freund's Complete Adjuvant is used to induce arthritis. The result of this study revealed that *Asparagus racemosus* show potent effect on both the condition at a dose of 200mg/kg and 400mg/kg respectively.

Key Words: *Asparagus racemosus*, Anti-inflammatory, Anti-arthritis, Carrageenan.

INTRODUCTION

If infectious diseases are troubling the general population on one hand, the pace of modern life is adding to the woes of the middle class¹. Numerous physiological and biochemical processes in the human body may produce oxygen-centered free radicals and other reactive oxygen species as by products. Overproduction of such free radicals can cause oxidative damage to biomolecules (e.g. lipids, proteins, DNA), eventually leading to many chronic diseases, such as atherosclerosis, cancer, diabetes, aging, and other degenerative diseases in human².

When cells in the body are damaged by microbes, physical agents or chemical agents, the injury is in the form stress. The response to the stress of tissue damage is called as inflammation³. Inflammation is a complex process, which is frequently associated with pain and involves occurrences such as: the increase of vascular permeability, increase of protein denaturation and membrane alteration⁴. Inflammatory response is a series of well coordinated dynamic mechanism consisting of specific vascular, humoral and cellular events that is characterized by the movement of fluids, plasma and inflammatory leukocytes (neutrophils, eosinophils, basophils and macrophages) to the site of inflammation⁵. Lysosomal enzymes released during inflammation produce a variety of disorders which leads to the tissue injury by damaging the macromolecules and lipid peroxidation of membranes which are assumed to be responsible for certain pathological conditions as heart attacks, septic shocks and rheumatoid arthritis etc. The extra cellular activity of these enzymes is said to be related to acute or chronic inflammation⁶. Although rheumatism is one of the oldest known diseases of the mankind and affects a large percentage of population of the world⁷. The management of rheumatoid arthritis (RA) rests on several principles such as drug treatment, which comprises diseasemodifying anti rheumatic drugs (DMARDs) and also non-steroidal anti-inflammatory drugs and gluco corticoids (GCs), as well as nonpharmacological measures, such as physical, occupational and psychological therapeutic approaches, together may lead to therapeutic success⁸. Most of the anti-inflammatory drugs are steroidal or nonsteroidal anti-inflammatory drugs. Though they are very useful, they have a number of severe adverse effects such as gastrointestinal disturbances and body fat redistribution⁹. To

overcome these problems the preparations from plant origin become important in modern medicine and widely prescribed in traditional medicinal systems¹⁰. Numerous types of herbs have been well recognised and catalogued by botanist from the high ranges of the Himalayan tract up to the sea-shores of Kanyakumari. According to WHO nearly 80 % of the global population still rely upon the herbal drugs for their primary health care¹¹. The present study was focused to evaluate the anti-arthritis and anti-inflammatory potentials of ethanolic extract of *Asparagus racemosus* roots by widely accepted methods.

MATERIALS AND METHODS

Collection of plant: The part of the plant (roots) has been collected from local market of Lucknow (U.P) and identified by the University of Rajasthan, Jaipur. After procurement, the roots were dried in shade and ground mechanically into a coarse powder and kept into an air-tight container for use in the study.

Preparation of extract: The powdered plant material was successively extracted using ethanol in a soxhlet extraction apparatus. The extract was concentrated and traces of the solvent were completely removed under reduced pressure and stored in vacuum desiccators for further use. The extract was named as ethanol extract of *Asparagus racemosus* (AREE).

Chemicals: Carrageenan and Freund's Complete Adjuvant were purchased from UGO Basil, Sigma Aldrich (USA) and the entire reagents used in the study were analytical grade.

Selection of Animals: Wistar rats (150-200 g) of either sex, procured from Jaipur College of Pharmacy, Jaipur, were used. Animals were housed at $25 \pm 1^\circ\text{C}$ and at standard environmental conditions (12 h light and 12 h dark cycle) in the institutional animal house. The animals were fed with standard pellet rodent diet and water was provided *ad libitum*. All the experiments were carried out between 09.00 am to 5.30 pm and approved by Institutional Animal Ethical Committee of Jaipur College of Pharmacy, Jaipur.

Pharmacological Study

Carrageenan induced inflammation: Carrageenan-induced rat paw oedema has been used for assessment of the anti-inflammatory activity of many plant extracts¹². The animals were divided into 5 groups of 6 animals each (one normal, one control, one standard and two test groups). Edema in the rats were induced by injection of 0.05 ml of carrageenan (prepared as 1% w/v suspension in saline) locally injected into sub plantar region of the left hind paw of rats.

Treatment Groups

Normal: 1% aqueous solution of Tween80, p.o.

Control: Carrageenan + 2% Tween80 (10ml/kg).

Standard: Carrageenan + Diclofenac sodium (4mg/kg)

Test group (Low dose): Carrageenan + AREE (200mg/kg)

Test group (High dose): Carrageenan + AREE (400mg/kg)

The extracts were administered orally into the rats 1 hour prior to carrageenan injection. Diclofenac sodium (4 mg/kg) was given to standard group. The volume of the paw was measured 1 h before the injection and at 1, 2, 3, 4, 5, 6 and 24 hrs after the injection of carrageenan. Edema was expressed as the increment in paw thickness due to carrageenan administration¹³. The percentage inhibition of paw volume in extract treated groups was compared with control.

Reduction in the paw volume were compared with the vehicle treated controlled animals with that of the test groups and the anti-inflammatory activity were carried out on the basis of the percentage (%) inhibition of edema. The percent inhibition of edema was calculated by using the formula¹⁴.

$$\% \text{ inhibition of edema} = (V_c - V_t / V_c) \times 100$$

Where V_t = Paw volume in test group animals,

V_c = Paw volume in control group.

Freund's Complete Adjuvant induced Arthritis

Arthritis was induced by a single intra-dermal injection (0.1 ml) of Complete Freund's adjuvant (CFA) containing 1.0 mg dry heat-killed *Mycobacterium tuberculosis* per milliliter sterile paraffin oil into a foot pad of the left hind paw of rats¹⁵. The animals were divided into 5 groups of 6 animals

each (one normal, one control, one standard and two test groups).

Treatment groups

Normal: 1% aqueous solution of Tween80

Control: FCA + 2% Tween80 (10ml/kg).

Standard: FCA + Dexamethasone (5mg/kg)

Test group (Low dose): FCA + AREE (200mg/kg)

Test group (High dose): FCA + AREE (400mg/kg)

The dosing of all the groups was started from day 12th once daily orally. Various parameters i.e. body weight, joint diameter, paw volume, arthritic score, motor incoordination, analgesic have been evaluated on day 0th, 4th, 7th, 10th, 12th, 14th, 17th, 19th, 21th, and day 28th. On last day (day 28th), blood was withdrawn by retro-orbital puncture for assessment of hematological parameters i.e. WBC, RBC, Hb, ESR.

Behavioral assessment

Paw Volume

Paw volumes of both hind limbs were recorded on the day of FCA injection, and again measured on day 0th till day 28th using mercury plethysmometer¹⁸. The change in paw volume was measured as the difference between the final and initial paw volumes¹⁶.

Joint Diameter

Paw thickness was measured by compressing the joint by rotating the screw of micrometer screw gauge till the pain elicited as indicated by squeaking or leg withdrawal. The distance moved by the screw gauge was recorded¹⁷.

Arthritic Score

The arthritic severity in each paw was graded from 0 to 4:

0= paws with no swelling and focal redness.

1= paws with swelling of finger joints.

2= paws with mild swelling of ankle or wrist joints.

3= paws with severe inflammation of the entire paw.

4= paws with deformity or ankylosis.

Each paw was graded and the four scores were totalled so that the possible maximum score per rat was 16¹⁸.

Table.1: Effect of AREE on paw edema

S. No.	Time (Hour)	Control	Normal	Standard	AREE (200mg/kg)	AREE (400mg/kg)
1.	1	0.2±0.00	0.00 ± 0.000	0.200 ± 0.000	0.2±0.00	0.2±0.00
2.	2	0.4±0.00	0.00 ± 0.000	0.333 ± 0.067	0.33±0.06	0.2±0.00
3.	3	0.53±0.06	0.00 ± 0.000	0.333 ± 0.067	0.4±0.11	0.4±0.00
4.	4	0.6±0.00	0.00 ± 0.000	0.267± 0.067**	0.33±0.06*	0.33±0.06**
5.	5	0.53±0.06	0.00 ± 0.000	0.067±0.067***	0.26±0.06*	0.26±0.06**
6.	6	0.6±0.11	0.00 ± 0.000	0.000 ± 0.000***	0.2±0.00***	0.06±0.06***
7.	24	0.6±0.11	0.00 ± 0.000	0.000 ± 0.000***	0.2±0.00***	0.00±0.00***

Values were expressed Mean ± SEM. *P < 0.05; **P < 0.01 and ***P < 0.001 significant as compared to control group animals.

Table.2: Effect of AREE on average body weight

S.No.	Days	Control	Normal	Standard	AREE (200 mg/kg)	AREE (400 mg/kg)
1.	12	94.16± 3.27	101.66 ± 13.017	161.66± 6.00	75.00± 0.000	83.33± 8.333
2.	14	88.33± 2.10	108.33 ± 8.333	168.33± 4.41***	90.00± 5.773	86.66± 9.279
3.	17	77.50± 2.14	108.33 ± 8.333	171.66± 1.66***	106.66± 6.666*	98.33± 13.017
4.	19	76.66± 2.47	108.33 ± 8.333	176.66± 1.66***	116.66± 3.333**	103.33± 14.813*
5.	21	72.50± 1.11	108.33 ± 8.333	178.33± 1.66***	130.00± 5.000***	128.33± 11.666***
6.	28	72.50± 1.11	116.66 ± 8.333	188.33± 3.33***	143.33± 6.666***	133.33± 8.333***

Values were expressed Mean ± SEM. *P < 0.05; **P < 0.01 and ***P < 0.001 significant as compared to control group animals.

Table.3 Effect of AREE on average arthritic score

S.No.	Days	Control	Normal	Standard	AREE (200 mg/kg)	AREE (400 mg/kg)
1.	12	2.00± 0.365	0.00± 0.00	2.00± 0.000	2.33± 0.333	2.66± 0.333
2.	14	1.83± 0.166	0.00± 0.00	1.66± 0.333	2.00± 0.000	2.33± 0.333
3.	17	1.83± 0.166	0.00± 0.00	0.00± 0.000***	1.33± 0.333	1.00± 0.333*
4.	19	2.16± 0.307	0.00± 0.00	0.00± 0.000***	1.33± 0.333	1.00± 0.333*
5.	21	2.33± 0.210	0.00± 0.00	0.00± 0.000***	1.00± 0.000**	0.33± 0.333***
6.	28	1.83± 0.166	0.00± 0.00	0.00± 0.000***	0.66± 0.333***	0.33± 0.333***

Values were expressed Mean ± SEM. *P < 0.05; **P < 0.01 and ***P < 0.001 significant as compared to control group animals.

Table.4 Effect of AREE on average paw volume

S.No.	Days	Control	Normal	Standard	AREE (200 mg/kg)	AREE (400 mg/kg)
1.	12	0.46± 0.04	0.00± 0.00	0.40± 0.115	0.533± 0.066	0.667± 0.067
2.	14	0.50± 0.0	0.00± 0.00	0.33± 0.066	0.400± 0.115	0.46± 0.066
3.	17	0.50± 0.04	0.00± 0.00	0.20± 0.000**	0.333± 0.066	0.46± 0.066
4.	19	0.50± 0.04	0.00± 0.00	1.33± 0.066***	0.266± 0.066*	0.266± 0.066*
5.	21	0.46± 0.04	0.00± 0.00	0.06± 0.066***	0.133± 0.066**	0.067± 0.067***
6.	28	0.46± 0.06	0.00± 0.00	0.06± 0.066***	0.133± 0.066**	0.067± 0.067***

Values were expressed Mean ± SEM. *P < 0.05; **P < 0.01 and ***P < 0.001 significant as compared to control group animals.

Carrageenan induced Inflammation

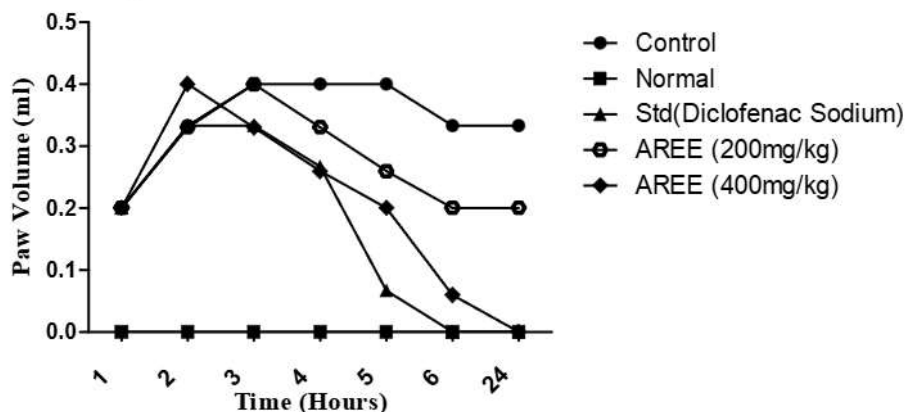


Figure 1: Effect of AREE on Carrageenan induced inflammation

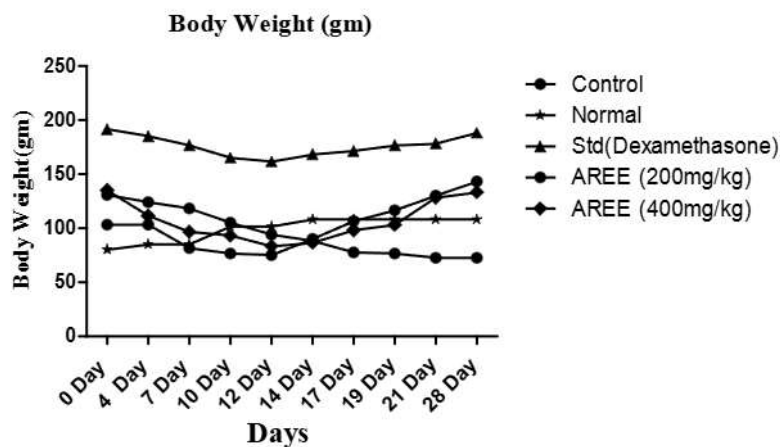


Figure 2: Effect of AREE on body weight

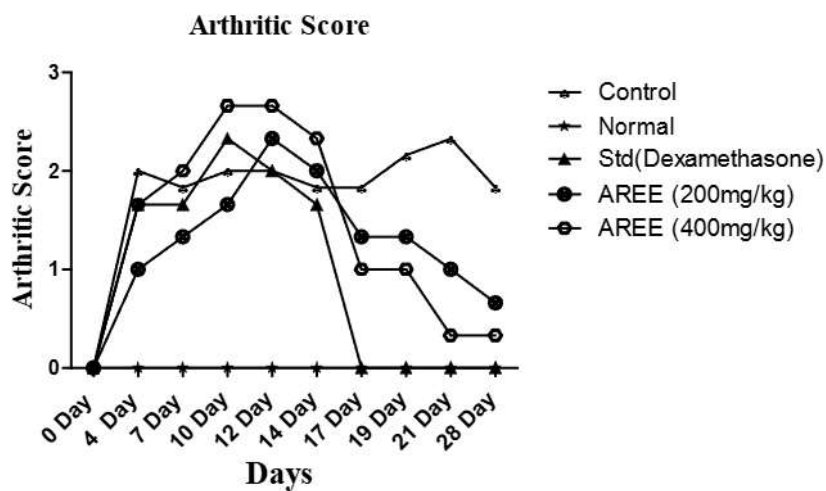


Figure 3: Effect of AREE on arthritic score

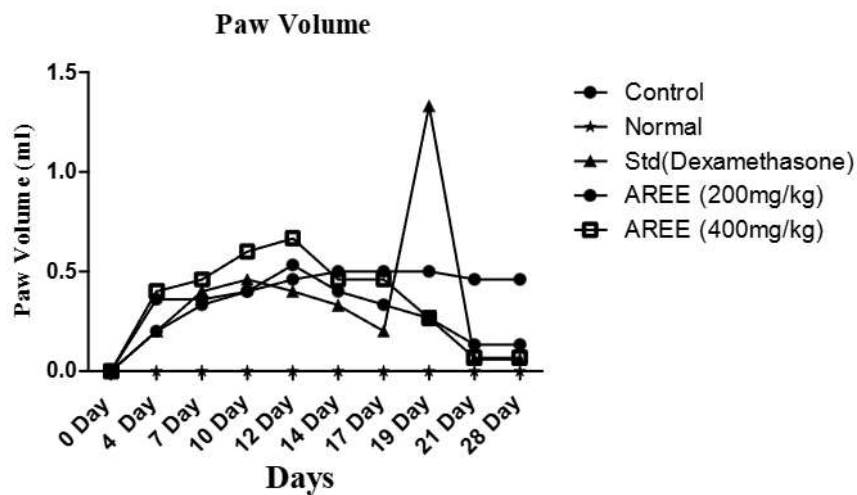


Figure 4: Effect of AREE on paw volume

RESULTS

Carrageenan induced inflammation

Anti-inflammatory effect of AREE of roots was evaluated after subplantar injection of carrageenan in animals. The standard Diclofenac (4 mg/kg) showed significant and dose-dependent decrease ($P < 0.01$ and $P < 0.001$) in paw edema on 4th, 5th, 6th and 24th hours as compared to control group animals. Whereas treatment with AREE (400 mg/kg) showed significantly ($P < 0.01$ and $P < 0.001$ respectively) decrease in paw edema as compared to control group animals on 4th, 5th, 6th and 24th hours. AREE (200 mg/kg) showed significant decrease in paw edema as compared to control group animals ($P < 0.05$, $P < 0.01$ and $P < 0.001$) on 4th, 5th, 6th and 24th hours. (Table.1, Figure 1)

FCA induced arthritis

Behavioral assessment

Effect of AREE on body weight

Standard group animals showed significant increased in body weight ($P < 0.05$, $P < 0.01$ and $P < 0.001$) from 14th day to 28th day as compared to the control group animals. Treatment with AREE (400 mg/kg) showed significant increase in body weight ($P < 0.05$, and $P < 0.001$) as compared to control group animals from 19th day to 28th day. Treatment with AREE (200 mg/kg) showed significant increase in body weight ($P < 0.05$, $P < 0.01$ and $P < 0.01$) as compared to control group animals from 17th day to 28th day. (Table.2, Figure 2)

Effect of AREE on arthritic score

All the groups of animals administered with FCA started showing signs of clinical inflammation in one or more hind paws, which was a biphasic response. The arthritic score was significantly increased from day 7th to 12th in control group animals which remained significantly increased till the end of the study i.e. up to 28th day. Animals treated with Standard drug showed significant and dose dependant decreased in arthritic score ($P < 0.001$) from day 17th onward till the end of the study as compared to control group animals. Treatment with AREE (400 mg/kg) showed significant decreased in arthritic score ($P < 0.05$, $P < 0.001$) as compare to control group animals from 17th day to 28th day. Treatment with AREE (200 mg/kg) showed significant decreased in arthritic score ($P < 0.01$ and $P < 0.001$) as compare to control group animals from 21th day to 28th day. (Table. 3, Figure 3)

Effect of AREE on paw volume

Standard group animals showed significant decrease in paw volume ($P < 0.01$ and $P < 0.001$) 17th day to 28th day as compared to the control group animals. Treatment with AREE (400 mg/kg) showed significant decreased in paw volume ($P < 0.05$, and $P < 0.001$) as compare to control group animals from 19th day to 28th day. Treatment with AREE (200 mg/kg) showed significant decreased in paw volume ($P < 0.01$, $P < 0.001$) as compare to control group animals from 19th day to 28th day. (Table. 4, Figure 4)

DISCUSSION

Carrageenan induced inflammation:

The primary phase of edema has been attributed to the release of histamine and serotonin; the edema maintaining during the plateau phase, attribute to kinin like substances and the secondary accelerating phase of swelling is attributed to the release of prostaglandin¹⁹. Mediators like leukotriene, prostaglandins, PAF and cytokines are reported to be

responsible for the immediate hypersensitivity reaction, but it was observed that enhanced vascular permeability and leukocyte infiltration at the sites of allergen challenge²⁰.

FCA induced arthritis

The development of adjuvant-induced arthritis in the rat can be divided into three phases, just like human rheumatoid arthritis, starting with the induction phase without evidence of synovitis, followed by early synovitis, and finally late synovitis with progressive joint destruction²¹. Inhibition of COX-2 activity also modulated local and systemic cytokine production in arthritic rats. The development of arthritis was associated with increased levels of TNF- α and IL-6 mRNAs in affected paws and systemic IL-6 production. Both cytokines have been shown to be produced spontaneously by rheumatoid arthritis synovial cells²².

CONCLUSION

Treatment with ethanolic extract of *Asparagus racemosus* (200 and 400 mg/kg, p.o.) showed maximum reduction in paw volume as compared to vehicle treated animals in carrageenan induced rat paw edema. A significant increase in body weight, reduction in paw volume of both hind legs and reduction in total arthritic score were observed in FCA induced arthritis in rats. All these results thus envisage that the drug provide pharmacological rationale for the traditional use of the drug against inflammatory disorders such as rheumatoid arthritis.

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