

Scholars Research Library

J. Nat. Prod. Plant Resour., 2013, 3 (5):64-70 (http://scholarsresearchlibrary.com/archive.html)



Comparison of Anti-inflammatory activity of *Ampelocissus latifolia* (Roxb.) root extract: Oral administration Vs. Topical application

Jitendra G. Patel*¹, Natvarlal M. Patel², Amit A. Patel², Amish J. Patel² and Sohan Patel²

¹Hemchandracharya North Gujarat University, Patan ²Shri B M Shah College of Pharmaceutical Education &Research, Modasa, Gujarat

ABSTRACT

Ampelocissus latifolia (fam. Vitaceae) is climber with annual stems and fasciculated tuberous roots found in India. Traditionally people uses root in skin disease, fracture, as a tonic, for wound healing, diuretic, in eye disease, gonorrhoea, syphillis, menstrual troubles, rheumatic affection. A. latifolia contains flavonoids, saponins and reducing sugar. In present study freshly collected and dried root powder of Ampelocissus latifolia (Roxb.) was subjected to hydro alcoholic extraction. The extracts were tested for anti-inflammatory activity in carrageenan induced paw edema in rat by oral and topical application. The result found marked inhibition of inflammation in orally given and topically applied extracts. The results also suggest that hydro alcoholic extract administered orally is more effective than topically applied extract.

Key words: Ampelocissus latofolia, Roots, Anti-inflammatory, Topical, Hydro alcoholic.

INTRODUCTION

Ampelocissus latifolia(fam. Vitaceae), is climber with annual stems found mainly in sub-Himalaya tract from sutlej eastwards to Kumaon up to 4000 ft., Aasam, Konkan, W. ghats from bombay to Nilgiris and Anamallis Deccan, and throughout in india. Roots are arising adventitiously from root stock. Roots found fasciculated with many tuberous roots in cluster having irregular shape, 7 to 15 cm in length and 1.5 to 4 cm in diameter. Roots are tapering at both the end with bulging in middle. Roots are reddish brown in colour and having no any characteristics taste. Dried roots are having scaly skin. Transverse section of A. latifolia root found circular in outline and contains Epiblema, Cortex,Pericyclic fibres, xylem, phloem and 6 to 10 seriate medullary rays. Acicular and Spharophide calcium oxalate crystals and starch is also present. A. latifolia in ayurveda reported to be used as Kustha, Kamala, Sotha, and Vrana [1]. Traditionally plant is used for Wound healing [2]. The stem bark is used in stomach Pain [3]. Stem is used in bene fracture [4]. Root finds use in skin disease [3], [4], [5], [6]. Roots are used in fracture and as a tonic [2], [7]. Root is used in menstrual troubles [8]. Root is used for wound healing [9]. Root is used as diuretic and in eye disease [10]. Root is used in gonorrhoea, syphillis [11]. Root used in rheumatic affection [12].

MATERIALS AND METHODS

Plant material

Fresh & fully grown plants of Ampelocissus latifolia collected in month of August, 2011 from the near places of Modasa city, Sabarkantha, Gujarat, India. It was authenticated by Dr. H. B. Singh Scientist and Head of Raw Materials Herbarium & Museum Dept of National Institute of Science and Communication and Information Resources, New Delhi (NISCAIR). The herbarium of this plant is deposited (voucher specimen no. BMCPER/HNGU/11-12), in Dept. of Pharmacognosy, Shri B. M. Shah College of Pharmaceutical Education and Research, Modasa

Preparation of extract

Locally collected whole plants (3.0 kg) were shade-dried and its roots and aerial parts separatedfrom each other and then roots waspowdered. Powdered rootsextracted with Hydro alcohol (1:1). The extracts were dried under reduced pressure yielding reddishbrown solid mass. These hydro alcoholic extracts was dissolved in respective solvents and used for Anti-inflammatory activity.

Animals

Male Albino rats of either sex, weighing 180-250 g were used. Theywere housed in standard environmental conditions of temperature, humidity, and light and provided with standard rodent food and water *ad libitum*.

Anti-Inflammatory Activity [13], [14], [15].

The anti-inflammatory activity of Hydro alcoholic extract of A. latifolia root wasevaluated by the carrageenan-induced rat hind paw edemamethod. The experimental protocol (No: IAEOBMCPER/11/2011-12) wasdesigned and approval of Institutional AnimalEthics Committee (IAEC) (Reg. No. /date: 194/CPCSEA/1st June 2001) wasobtained. Healthy male albino rats weighing between 180-250 g were obtained from the diseasefree animal house of ZRC, Ahmedabad. Theanimals were housed in institutional animal houseunder standard conditions with free access to foodand water. Anti-inflammatory activity of Hydro alcoholic extract of A. latifolia root was compared with Standard Indomethacin (20mg/kg) for extract given orally. Hydro alcoholic extract of A. latifolia root applied topically was compared with the marketed gel of diclofenac (Diagesic gel).

Twenty eight albino rats were divided into sevengroups of four animals each as follows:

Group 1 (Negative Control): Water

Group 2 (Positive Control group): 0.1ml of 1 % Carageenan

Group 3 (Standard (Oral)): Indomethacin (20 mg/kg)

Group 4 (Test 1(Oral)): Hydro alcoholic extract of A. latifolia root (500mg/kg).

Group 5 (Test 2(Oral)): Hydro alcoholic extract of A. latifolia root (750mg/kg).

Group 6 (Standard (Topical)): 1.16 % Diclofenac Gel (0.2 gm).

Group 7 (Test (Topical)): Paste of hydro alcoholic extract of A. latifolia root (0.2 gm)

Afterone hour of the above respective (oral or topical) administration, carrageenan (1%, 0.1ml) was injected subcutaneously in the sub plantar tissue of the right hind paw ofeach rat. The inflammation was measured using plethysmometer immediately afterinjection of carrageenan and then 1,2,3 and 4h. Theaverage foot swelling in drug treated animal as well asstandard was compared with that of control and the percent inhibition (anti-inflammatory activity) of edema was determined using the formula.

Percentage inhibition = [(C-T)/C] * 100

Where C = control paw edema, T = test paw edema.

Statistical analysis

All the results are expressed as mean \pm standard error of mean. The data was analyzed statistically using ANOVA followed by Dunnett's Multiple Comparison Test.

RESULTS AND DISCUSSION

• The anti-inflammatory activity of hydro alcoholic extract of *Ampelocissus latifolia* revealed significant inhibition of inflammation as compared to control group in orally administered and topically applied extract.

- Result focus that in oral administration inhibition is increases with increase in dose.
- Initially oral administration is giving faster effect than topical application because of slow permeability of topical applied extract. But after 3 hour orally administered and topically applied groups are showing significant inhibition of inflammation.(**Tab 1 and Fig 1 to 6**)

Table 1: Anti-inflammatory activity of Ampelocissus latifolia (Roxb.) Root (Oral Administration Vs. Topical Application)

	Group A	Group B	Group C	Group D	Group E	Group F	Group G
	Control (-ve)	Control (+ve)	Standard (Oral) Indomethacin 20 mg/kg	Test 1 (Oral) 500 mg/kg	Test 2 (Oral) 750 mg/kg	Std. (Topical) Diclofenac gel	Test (Topical)
	Mean	Mean	Mean	Mean	Mean	Mean	Mean
Paw Volume	\pm SEM	\pm SEM	\pm SEM	\pm SEM	\pm SEM	\pm SEM	\pm SEM
	ml	ml	ml	ml	ml	ml	ml
Time (Hour)							
0	1.625	1.625	1.625	1.650	1.650	1.575	1.600
	± 0.02500	± 0.02500	±0.04787	±0.02887	±0.02887	± 0.04787	± 0.04082
1	1.625	2.45	2.025	2.15	2.100	2.025	2.325
	± 0.02500	± 0.08660	±0.04787**	±0.09574	±0.09129*	±0.07500**	±0.08539
2	1.625	3.35	2.075	2.65	2.4	2.100	2.9
	± 0.02500	±0.1041	±0.06292**	±0.1555**	±0.1080**	$\pm 0.04082^{**}$	$\pm 0.1080^*$
3	1.625	4.525	2.150	2.925	2.525	2.175	3.325
	± 0.02500	±0.2016	±0.06455**	±0.04787**	±0.1652**	±0.04787**	$\pm 0.1250^{**}$
4	1.625	3.175	1.925	2.325	2.225	1.975	2.45
	± 0.02500	±0.1181	±0.06292**	±0.09574**	±0.1493**	±0.04787**	$\pm 0.06455^{**}$

Statistical analysis by **ANOVA** and **Dunnet's Multiple comparison Test**. Results are expressed as mean \pm standard error of mean, n = 4 in each group. *Less significant difference compared to control group at p < 0.05. ** Significant difference compared to control group at p<0.01

Figure 1

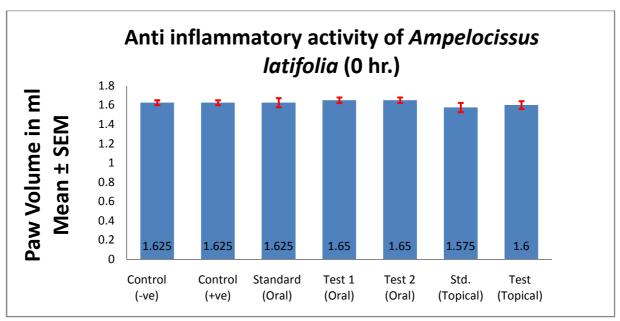


Figure: 2

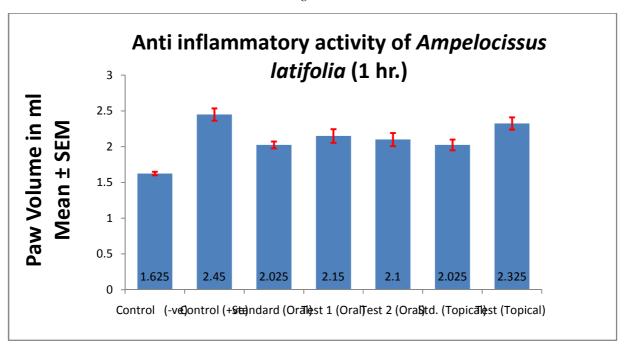


Figure: 3

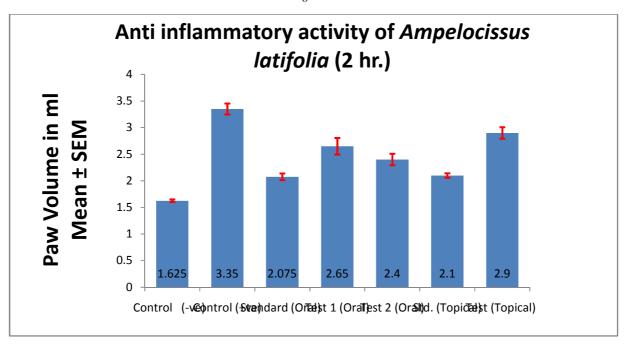


Figure: 4

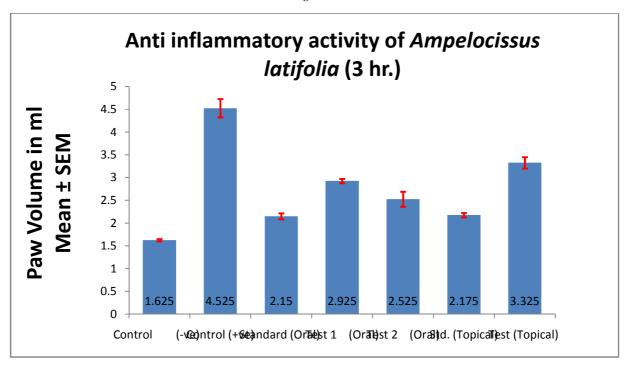
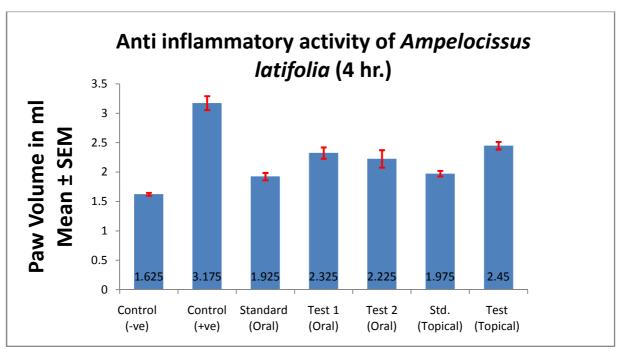
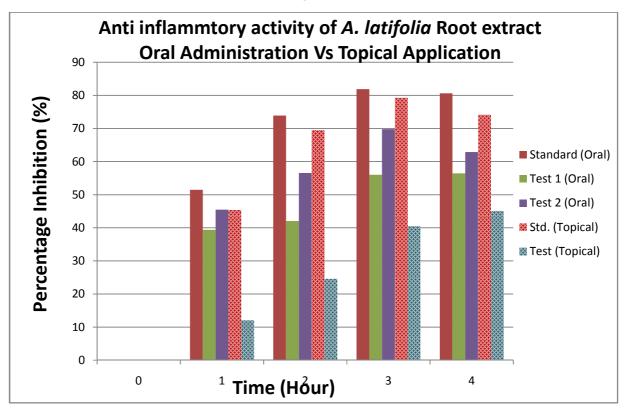


Figure: 5







CONCLUSION

Hydro alcoholic extract of Ampelocissus latifolia (Roxb.) root extract was found effective in inhibition of inflammation. Furthermore this study was carried out for comparison of oral and topical application of hydro alcoholic extract. Result suggests that oral administration is faster effective than topical administration. But if any topical formulation of extract is prepared than it may be more convenient and safe than oral administration with same effectiveness.

REFERENCES

- [1] Kirtikar KR, Basu BD, Indian Medicinal Plant, International book Distributor, 1998, 1, 606.
- [2] Sikarwar RLS and Kaushik JP, Int. J. Pharmacog, 1993, 31, 283-287.
- [3] Joshi MC, Patel MB and Mehta PJ, Bull Med Ethnobot Res, 1980, 1, 8-24.
- [4] Singh KK, Kalakoti BS and Prakash A, J Bombay Nat Hist Soc, 1994, 91, 386-390.
- [5] Shah GL and Gopal GV, Acta Bot Indica, 1986, 14, 48-53.
- [6] Saxena HO, Bull Bot Surv India, 1986, 28, 149-156.
- [7] Jain SK and Tarafder CR, Econ Bot, 1970, 24, 241-278.
- [8] Shah GL, Menon AR and Gopal GV, J. Econ Tax Bot, 1981, 2, 173-182.
- [9] Chandra K, Pandey BN, Sinha GN and Pandey P, Bull Med Ethnobot Res, 1989, 10, 124-161.
- [10] Badhe PD and Pandey VK, Bull Med Ethnobot Res, 1990, 11, 1-39.
- [11] Sharma PC, Bull Med Ethnobot Res, 1988, 9, 89-85.
- [12] Jain SP. and Puri HS, *India. J Ethnopharmacol*, **1984**, 12, 213-222.
- [13] http://www.biomedcentral.com/1472-6882/12/5. Muhammadet al. BMC Complementary and Alternative Medicine 2012.
- [14] Vogel Gerhard H, et al. Drug discovery and Evalution Pharmacological Assay. Springer-Verlag Berlin Heidelberg New York. 2nd edition, p.no:759-761.
- [15] Goyal C, Ahuja M and Sharma Sk, Acta Poloniae Pharmaceutica Drug Research, 2011, 68, 147-150.

- [16] Rang H.P., Dale M.M. and Ritter, "Local hormones, inflammation and allergy", Chapter 12 In; Pharmacology, New York: Churchill Livingstone, 4^{th} Edition, **1999**, 199-225.
- [17] LJ Robert: JD Morrow; JG Hardman; LE Limbird; AG Gilman. Goodman and Gilman's The Pharmacological Basis of Therapeutics, McCraw Hill, New York, 200; pp 687
- [18]OECD Guidelines for Testing of Chemicals AcuteOral toxicity- Fixed Doseprocedure, Guideline no.420, 17thDecember**2001.**
- [19] M.N. Gosh; Fundamentals of experimental pharmacology, Hilton and company, Kolkata, 2005, 190-197.
- [20] ML Ferrandiz; MJ Alcaraz. Agents Actions, 1991, 32, 283.