Journal of Chemical and Pharmaceutical Research, 2015, 7(7):1184-1187



Research Article

ISSN : 0975-7384 CODEN(USA) : JCPRC5

Anti-inflammatory activity of acacia catechu bark extract-in vitro study

Ghayathri¹ and Lakshmi T.*²

¹Bachelor of Dental Surgery, Department of Pharmacology, Saveetha Dental College, Chennai ²Department of Pharmacology, Saveetha Dental College, Chennai

ABSTRACT

Rheumatoid arthritis (RA) is a chronic inflammatory disease characterized by joint swelling, joint tenderness, and destruction of synovial joints, leading to severe disability and premature mortality. Acacia catechu Willd. Commonly known as karungali in Tamil and Khadira in Sanskrit .Chemical constituents are well known for their potential health benefits and have been reported to possess valuable biological activities such as antibacterial and antifungal, antioxidant, antiurolithiatic, anticonvulsant and anxiolytic, and hepato protective properties. The aim of this article is to investigate the Anti inflammatory activity of Acacia catechu Bark ethanolic extract.

Keywords: Acacia catechu Bark, Anti-Inflammatory, Biological, Rheumatoid arthritis, inflammation.

INTRODUCTION

Inflammation is the body's immediate response to damage tissues and cells by pathogens, certain stimuli such as chemicals or even by physical injury.^{1,2} Acute inflammation is a short term response that usually results in healings. For example ; leukocytes infiltrate the damaged region , removing the stimulus and thus repairing the tissues.³ On the other hand, chronic inflammation is a prolonged, and unregulated response that involves acute inflammation , tissue destruction and also attempts at tissue repair. ⁴⁻⁶Such persistent inflammation is associated with many chronic human diseases , including allergy , rheumatoid arthritis , and also autoimmune diseases. In specific , rheumatoid arthritis is an autoimmune disease that can cause chronic inflammation of the joints and other areas of the body. Moreover, inflammatory mediators are released from tissues during injury or any inflammatory reactions.⁷ They are capable of producing edema when injected locally. Stimulation of smooth muscles contraction *in vitro* and even in the alternating blood pressure when injected systemically. Histamine, major basic protein, complement, and arachidonic acid metabolites are examples of inflammatory mediators. Furthermore, NSAID's are prescribed for anti inflammatory actions. Because all NSAIDs can produce side effects, patients, especially if they are elderly, the drug should be selected carefully for the corresponding treatment.

Acacia catechu belongs to the family *leguminosae*. The common names includes cutch, black catechu, black cutch, wattle bark, black cattle . ⁸⁻¹⁰*Acacia* bark is an herbal plant used mainly for digestive disorders and to treat diarrhea. it is a natural astringent rich in tannic acid .11- Recent studies reported that *Acacia catechu* bark extract may aid to block the bodies pain trigger mechanism. it possess significant pharmacological , nutritive value. Acacia bark is hardened woody with a rusty appearance, brown color it contain tannins and Gallic acid. It is also employed in tanning industry. ¹⁵⁻¹⁸*Acacia* bark is commonly used in maintaining dental and oral hygiene. The fresh twigs is used for the protection of gums and teeth. ¹⁹⁻²² It is also considered useful as an external application for mouth ulcers , it reduces gingival inflammation. Used in case of leprosy in rural areas.

Plant material

EXPERIMENTAL SECTION

Acacia catechu Bark is obtained as an gift sample from Green Chem Herbal Extracts & Formulations, Bangalore.

Chemicals

Diclofenac sodium is obtained from sigma Aldrich (USA), All the chemicals used were of analytical grade.

EVALUATION OF INVITRO ANTI-ARTHRITIC ACTIVITY

Inhibition of Protein Denaturation method²³,²⁴

Concentration of test substance: 1000 to 200µg/ml Standard: Diclofenac sodium Chemicals Required : Bovine serum albumin, 1N HCl, Phosphate buffer (pH 6.3)

Instrument : Incubator, Spectrophotometer - 660nm

The following 4 solutions is used

Test solution (0.5ml) consists of 0.45ml of bovine serum albumin (5% w/v aqueous solution) and 0.05ml of test solution.

1. **Test control** solution (0.5ml) consists of 0.45ml of bovine serum albumin (5% w/v aqueous solution) and 0.05ml of distilled water.

2. Product control (0.5ml) consists of 0.45ml of distilled water and 0.05ml of test solution

3. **Standard solution** (0.5ml) consists of 0.45ml of bovine serum albumin (5%w/v aqueous solution) and 0.05ml 0f Diclofenac sodium (200µg/ml).

All of the above solutions were adjusted to pH 6.3 using a small amount of 1N Hcl. The samples were incubated at 37°C for 20minutes and heated at 57°C for 3 minutes. After cooling, add 2.5ml of phosphate buffer to the above solutions. The absorbance of the solutions was measured using UV-Visible spectrophotometer at 416nm. The percentage inhibition of protein denaturation was calculated using the formula. The percentage inhibition of Protein denaturation will be calculated as follows.

OD of test solution – OD of product control Percent Inhibition = 100 – ------ X 100 OD of test control

The control represents 100% protein denaturation. The result is compared with Diclofenac sodium treated sample.

RESULTS AND DISCUSSION

Acacia catechu Bark exhibits significant anti inflammatory activity. The ethanolic solvent form of Acacia Bark shows an inhibitory activity when tested at 200-1000 μ g/ml by inhibiting denaturation of protein and its effect was compared with standard drug diclofenac sodium. The results are depicted in table 1 and represented in Fig 1. Auto antigen production in rheumatoid arthritis is due to denaturation of protein. From the results of the present study it can be stated that ethanolic bark extract of Acacia catechu is capable of controlling the production of auto antigen and inhibiting the protein denaturation in rheumatoid arthritis.



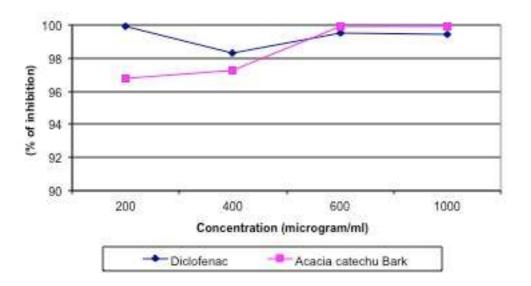


Table 1: invitro Anti- Inflammatory activity of Acacia catechu Bark Extract by inhibition of protein denaturation method

S.NO	Concentration µg/ml	Diclofenac % of Inhibition	Acacia catechu Bark extract % of Inhibition
1	200	99.88	99.92
2	400	98.27	99.91
3	800	99.52	99.88
4	1000	99.45	99.96

CONCLUSION

The *invitro* study by the inhibition of protein denaturation method emphasizes the anti-inflammatory/ anti arthritic efficacy of herbal extract similar to that of standard diclofenac sodium. The anti arthritic activity is due to the presence of flavonoids, phenols, polyphenols, and steroids. Further studies are mandatory, to identify the active constituent(s), that is responsible for the anti inflammatory efficacy.

Acknowledgements

The authors wish to thank Green Chem Herbal Extracts & Formulations, Bangalore for providing the extract as a gift sample for the study and the management, saveetha dental college and hospital for providing facility and kind support.

REFERENCES

1.Cruvinel Wde M, Mesquita D Jr, Araujo JA, Catelan TT, de Souza AW, da Silva NP, Andrade LE. Immune system – part I. Fundamentals of innate immunity with emphasis on molecular and cellular mechanisms of inflammatory response. Rev Bras Reumatol. 2010; 50(4): 434-61.

2.Acfarlane GJ, Paudyal P, Doherty M, Ernst E, Lewith G, MacPherson H. A systemic review of evidence for the effectiveness of practitioner – based complementary and alternative therapies in the management of rheumatic diseases : rheumatoid arthritis. Rheumatology (oxford). 2012; 51(9) : 1707-13.

3.McInnes IB, Schett G. The pathogenesis of rheumatoid arthritis. N Engl J Med. 2011; 365(23): 2205-19. 4.Tang F, Chen F, Ling X, Huang Y, Zheng X, Tang Q, Tan X, . Inhibitory effect of methyleugenol on IgE – mediated allergic inflammation in RBL-2H3 cells. Mediators inflamm. 2015; 463(30) : 530-17.

5.Chen Y, Paavola J, stegajev V, Stark H, Chazot PL, Wen JG, Konttinen YT. Activation of histamine H3 receptor decreased cytoplasmic Ca(2+) imaging during electrical stimulation in the skeletal myotubes. Eur J Pharmacol. 2015; 5(754): 173-8.

6.Preisner A , Albrecht S , Chui QL , Hucke S , Ghelman J , Hartmann C , Taketo MM , Antel J , Klotz L , Kuhlmann T. Non- steroidal anti inflammatory drug indomethcin enhances endogenous remylination. Acta Neuropathol. 2015; 20(4): 178-23.

7. Turkiewicz AM, Moreland LW. Rheumatoid arthritis. In: Bartlett SJ, Bingham CO, Maricic MM, Iversen MD, Ruffing V, eds. *Clinical Care in the Rheumatic Diseases*. 3rd ed. Atlanta, Ga: Association of Rheumatology Health Professionals; 2006.

8. Singh KN, Lal B, Note on traditional uses of Khair (*Acacia catechu Willd.*) by inhabitants of shivalik range of western Himalaya, Ethnobotanical Leaflets, 2006; 10;109-112.

 9. Naik GH, Priyadarsini KI, Satav JG, Banavalikar MM, Sohoni DP, Biyani MK, Mohan H, Comparative antioxidant activity of individual herbal components used in Ayurvedic medicine, 2003 ;63(1) ;97-104.
10. Anonymous, Indian Herbal Pharmacopoeia, Revised new edition 2002, Indian Drug Manufacturer's Association, Mumbai, 2002; 11(1): 13-16.

11.Lakshmi T., Anitha R., Geetha RV *Acacia catechu willd* -A gift from ayurveda to mankind – A Review. T. Ph. Res, 2011; 5(2); 273-93.

12.Lakshmi T, Aravindkumar S, Preliminary phytochemical analysis & Invitro Antibacterial activity of Acacia catechu willd Bark against Streptococcus mitis, Streptococcus sanguis & Lactobacillus acidophilus *International Journal of Phytomedicine* 2011; 3;579-84.

13. Evans WC. Trease and Evans' Pharmacognosy 14 Edn. W.B. Saunders Company, London; 1996.

14.Kalita, Tapan, B.K., Pal, T.K., Kalita, R.Estimation of total flavonoids content (TFC) and antioxidant activities of methanolic whole plant extract of Biophytum sensitivum Linn. J. Drug Delivery Ther. 2013;3 ;33–37.

15.Wan, C., Yu, Y., Zhou, S., Liu, W., Tian, S., Cao, S. Antioxidant activity and free radical-scavenging capacity of Gynura divaricata leaf extracts at different temperatures. Pharmacogn. Mag. 2011;7; 40–45.

16.Janakiraman N, Sahaya Sathish S, Johnson M. UV-VIS and FTIR Spectroscopic Studies on Peristrophe bicalyculata (Retz.) Nees. Asian Journal of Pharmaceutical and Clinical Research 2011; 4(4): 125-129.

17.Miller GL.Use of dinitro salicylic acid reagent for determination of reducing sugar. Anal. Chem 1959; 31 ;426-428.

18. Anis, Z., Sulaiman, O., Hashim, R., Mehdi, S.H., Ghalib, R.M. Radical scavenging activity: total phenol content and antifungal activity of Cinnamomum iners wood. Iranica J. Energy Environ. 2012; 3: 74–78

19.Deepa, S., Ramesh, K.P., Swarna, V.K., Rao, J.R., Chandrasekaran, B., 2013. Antioxidant and cytotoxic effects of methanolic extract of Salicornia brachiata L. in HepG2 cells. Int. J. Res. Pharm. Sci. 2013; 4; 512–517.

20.Kalita, Tapan, B.K., Pal, T.K., Kalita, R. Estimation of total flavonoids content (TFC) and antioxidant activities of methanolic whole plant extract of Biophytum sensitivum Linn. J. Drug Delivery Ther. 2013; 3, 33–37.

21. Mohammed Rahmatullah, Maraz Hossain, and Fatema Islam Antihyperglycemic and Antinociceptive Activity Evaluation of 'Khoyer' Prepared from Boiling the Wood of *Acacia Catechu* in WaterAfr J Tradit Complement Altern Med. 2013; 10(4): 1–5.

22. Singh KN, Mittal RK, Barthwal KC. Hypoglycemic activity of *Acacia catechu, Acacia suma*, and *Albizzia odoratissima* seed diets in normal albino rats. Indian Journal of Medical Research. 1976; 64:754–757.

23. Mizushima Y and Kbayashi m. interaction of anti-inflammatory drugs with serum proteins, especially with some biologically active proteins. J. Pharma Pharmacol, 20, 1968, 169-173.

24.Sadique J, Al Rqobahs WA, Bughaith, EI Gindi AR. The bioactivity of certain medicinal plants on the stabilization of RBC membrane system. Fitoterapia, 60, 1989, 525-532.