



EVALUATION OF IN VITRO ANTIUROLITHIATIC ACTIVITY OF *ALOE VERA*

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ABSTRACT

The present study was undertaken to evaluate the in vitro antiurolithiatic activity of the medicinal plant *Aloe vera*. Aqueous extract showed its maximum efficiencies in the dissolution of calcium oxalate crystals. Our results have clearly indicated that the Aqueous leaf extract of *Aloe vera* was quite promising for further studies in this regard. In this study Neeri was used as standard drug.

KEYWORDS: In vitro antiurolithiatic activity, Aqueous extract, urolithiasis, *Aloe vera*, Neeri.

1. INTRODUCTION

Plants provide food, raw materials for medicine and various other requirements for the very existence of life from the origin of human

Beings.^[1] Even the current conventional medicine is using a lot of plant derived chemicals as therapeutic agents. The overuse of synthetic drugs results in higher incidence of adverse drug reactions has motivated humans to return to nature for safe remedies. Herbs and herbal drugs have created interest among the people by its clinically proven effects.^[2] Therefore, there is a compelling need for detailed scientific validation of all traditional medicinal plant drugs to establish their efficacy and safety in light of modern science.

Kidney stone disease is a multi-factorial disorder resulting from the combined influence of epidemiological, biochemical and genetic risk factors.^[3] Urolithiasis is considered as the third most common affliction of the urinary tract. It refers to the solid non-metallic minerals in the urinary tract. It is a complex process that is a consequence of an imbalance between promoters and inhibitors in the kidney. The formation of kidney stones involves several

phytochemical events beginning with crystal nucleation, aggregation and end with retention within the urinary tract. Among the several types of kidney stones, the most common are calcium oxalate stones representing up to 80% of the analyzed stones.^[4] Calcium containing stones may be in the form of pure calcium oxalate(50%) or calcium phosphate(5%) and a mixture of both(45%) followed by magnesium phosphate(15-20%), uric 5 acid(10%) and cystine(1%).^[5]

It is estimated that at least 10% of the population in the industrialized part of the world is afflicted by urinary tract diseases and among these kidney stones are common with an annual incidence of 0.5 -1.9%. About 12% of the population of India is expected to have urinary stones and out of that 50% of cases encounter loss of one or both 2 kidneys with or without renal damage up to some extent.^[2]

Stone disease is 2-3 times more common in males, than in females. It 5 has a reoccurrence rate of 70-81% in males and 47-60% in females.^[5] In spite of substantial progress in pathophysiology and treatment of urolithiasis, there is no satisfactory drug being used in clinical therapy. Kidney dialysis, endoscopic stone removal and extra corporeal shock wave lithotripsy are prohibitively costly and reoccurrence is quite common with these procedures.^[1]

Data from in vitro and in vivo clinical trials revealed that phytotherapeutic agents could be useful as alternative therapy in the management of urolithiasis. Medicinal plants and their products are more useful, because they promote the repair mechanism in natural way^[1] Pharmacological and phytochemical prospecting of medicinal plants based on traditional knowledge can lead to the discovery of new drug and development of pharmacologically important products for human health care.^[6] Green medicines were safe and more dependable than the costly synthetic drugs, many of which have side effects.^[7]

Aloe vera has been widely grown as an ornamental plant. The species is popular with modern gardeners as a putatively medicinal plant and due to its interesting flowers, form, and succulence. This succulence enables the species to survive in areas of low natural rainfall, making it ideal for rockeries and other low-water use gardens.^[8] Aloe vera juice is used for consumption and relief of digestive issues such as heartburn and irritable bowel syndrome, although it bears significant potential to be toxic when taken orally.^[9]

2. MATERIALS AND METHODS

Plant Material

The leaves of *Aloe vera* were collected in the month of March 2018 from Sangareddy town, Sangareddy dist. of Telangana, India. The plant was authenticated by D.Venkateshwara Rao, Deputy Director, Telangana. Forest Academy, Dullapally, Hyderabad, Rangareddy District. The leaves were washed with tap water and dried under shade.

Preparation of Plant Extract

The leaves were shade dried and powdered. The crude plant extract was prepared by Soxhlet extraction method. 50g of powdered plant material was extracted with 500ml of water. The process of extraction was carried out up to 6 cycles, till the solvent in siphon tube of an extractor became colorless. The extracts were filtered, and evaporated to dryness using rotary evaporator. Further the dried extract was maintained in a refrigerator at 4°C for further antiurolithiatic activity.

Chemicals Used

Neeri, Sodium oxalate, Tris buffer, calcium chloride, Potassium permanganate (KMnO₄), Sulphuric acid (H₂SO₄).

Investigation of In Vitro Antiurolithiatic Activity Test By Titrimetry

The experimental kidney stones of calcium oxalate (CaOx) were prepared in the laboratory by taking equimolar solution of calcium chloride dehydrate in distilled water and sodium oxalate in 10 ml of 2N H₂SO₄. Both were allowed to react in sufficient quantity of distilled water in a beaker, the resulting precipitate was calcium oxalate. The precipitate was freed from traces of sulphuric acid by ammonia solution, washed with distilled water and dried at 60°C. The dissolution percentage of calcium oxalate was evaluated by taking exactly 1 mg of calcium oxalate and 10 mg of the extract, packed it together in semi permeable membrane of egg as shown in the model designed given below. This was allowed to suspend in a conical flask containing 100 ml of 0.1M Tris buffer. First group served as blank containing only 1 mg of calcium oxalate. The second group served as positive control containing 1 mg of calcium oxalate and along with the 10mg standard drug, i.e. Neeri. The 3rd group along with 1 mg of calcium oxalate contain aqueous extract. The conical flasks of all groups were kept in an incubator preheated to 37°C for 2 h. Remove the contents of semi permeable membranes from each group into separate test tubes, add 2 ml of 1N sulphuric acid to each test tube and titrated with 0.9494 N KMnO₄ till a light pink colour end point obtained. The amount of remaining

undissolved calcium oxalate is subtracted from the total quantity used in the experiment in the beginning to know the total quantity of dissolved calcium oxalate by various solvent extracts.^[10]

3. RESULTS AND DISCUSSION

Drug therapy has developed in response to population health care^[11] needs. There are many crucial areas in medicine such as liver diseases, arthritis, old age related problems, certain viral infections and cancer where the conventional medicine is devoid of satisfactory treatment. These are among the promising areas of research and development of medicines from the vast highly potential plant resources. Plants are also attractive sources for the development of novel and very effective and safe therapeutic agents against kidney procumbens. Herbal medicines are also in great demand in the developed world for primary health care because of their efficacy, safety and lesser side effects.^[12] Unlike allopathic medicines which target is only one aspect of urolithiatic pathophysiology, most of plant based therapy have been shown to be effective at different stages of stone pathophysiology.^[13] About 80% of the world populations rely on the use of traditional medicine which is predominantly based on plant materials.^[14] Plant based drug discovery programmes continue to provide an important source of new drug leads.^[15] Lithiasis (stone formation) is an important cause for acute and chronic renal failure, includes both nephrolithiasis (stone formation in kidney) and urolithiasis (stone formation in ureter or bladder or both). Among the various kinds of stones identified, calcium stones occur mainly in Men, while phosphate stones formation is more in women.^[16]

This study evaluates the antiurolithiatic activity of Aqueous extract of *Aloe vera*. The highest percentage i.e. 98.1% of calcium oxalate {CaOx} dissolution was observed in Aqueous extract . Aqueous extract of *Aloe vera* was found to be more effective in dissolution of calcium oxalate than standard drug Neeri. This study has given primary evidence for *Aloe vera* as the plant which possess lithotriptic property. This in vitro study has given lead data and shown that Aqueous extract are quite promising for further studies in this regard.

Table 1: Shows % dissolution of calcium oxalate (CaOx) by *Aloe vera* leaves extracts.

% of dissolution of calcium oxalate		
S.No	GROUPS	<i>Aloe vera</i>
1.	Blank	0
2.	Positive Control	81
4.	Aqueous extract	98.1



Figure 1(a): Decalcification of egg shell in 10% Acetic acid overnight.



Figure 1(b): Decalcified Eggs.

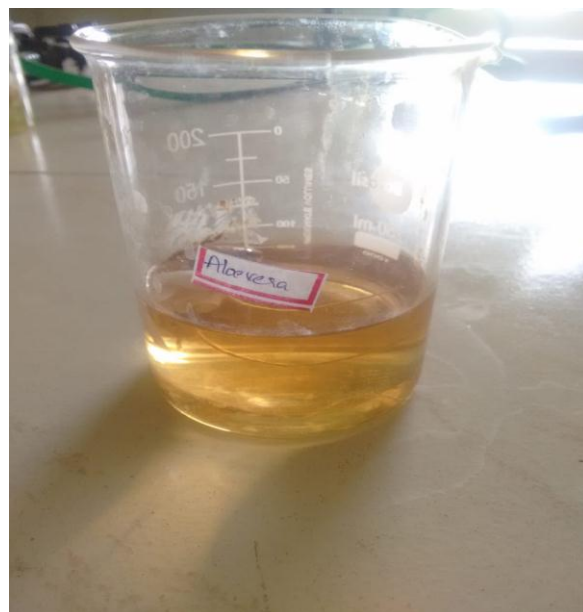


Figure 1(c): Egg membrane along with the contents suspended into the 0.1 M Tris buffer.

4. CONCLUSION

In vitro urolithiasis has been performed on the selected plant *Aloe vera* by using the standard drug, Neeri. The work was performed by using in vitro antiurolithiatic model for calculating percentage dissolution of kidney stone. Ethanolic leaf extracts of *Aloe vera* shows highest dissolution than standard drug Neeri. This study has given primary evidence for *Aloe vera* as the plant which possess antiurolithiatic property.

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REFERENCES

1. Sumayyasikandari and Prathima Mathad. In vitro antiurolithiatic activity of *Butea monosperma* Lam. and *Nigella sativa* Linn. seeds. *Ukaaz-Annals of Phytomedicine*, 2015; 4(1): 105-107.
2. Sanjay kumar Gupta, Madhavsinghbaghel, Chaturbhuj Bhuyan, B.Ravi Shankar, Ashok. BK, Panchakshari D Patil. Evaluation of anti-urolithiatic activity of Pashanabhedadi Ghrita against experimentally induced renal calculi in rats. *AYU (An international Quarterly journal of Research in Ayurveda)*, 2012; 33(5): 429-434.
3. Atul Makasana, Vishavas Ranpariya, Dishant Desai, Jaymin Mendpara, Vivek Parekh. Evaluation for the anti-urolithiatic activity of *Launaeaprocumbens* against ethylene glycol-induced renal calculi in rats. *Elsevier-Toxicology Reports*, 2014; 1: 46-52.
4. Jagannath. N, Somashekara S. Chikkannasetty, Govindadas D, Devasankaraiah G. Study of anti urolithiatic activity of *Asparagus racemosus* on albino rats. *Indian Journal of Pharmacology*, 2012; 44(5): 576-579.
5. Radha singanallur Ramu, Ravi Doraiswamy and Hiran Mai Yadav. Antiurolithiatic activity of Aqueous bark extract of *Crateva Magna* Lour. (DC). *International Journal of Research in Ayurveda and Pharmacy*, 2017; 8: 271-278.
6. A.Subramoniam. Present scenario, challenges and future perspectives in plant based medicinedevelopment. *Ukaaz-Annals of phytomedicine*, 2014; 3(1): 31-36.
7. Subramoianm. A. *Phytomedicines for healthcare*. *Ukaaz-Annals of Phytomedicine*; 3(1): 1-3.
8. Yates A. *Yates Garden Guide*. Harper Collins Australia, 2002.

9. Cosmetic Ingredient Review Expert Panel. "Final report on the safety assessment of Aloe Andongensis Extract, Aloe Andongensis Leaf Juice, Aloe Arborescens Leaf Extract, Aloe Arborescens Leaf Juice, Aloe Arborescens Leaf Protoplasts, Aloe Barbadensis Flower Extract, Aloe Barbadensis Leaf, Aloe Barbadensis Leaf Extract, Aloe Barbadensis Leaf Juice, Aloe Barbadensis Leaf Polysaccharides, Aloe Barbadensis Leaf Water, Aloe Ferox Leaf Extract, Aloe Ferox Leaf Juice, and Aloe Ferox Leaf Juice Extract". *Int J Toxicol.*, 2007; 26(2): 1-50. doi:10.1080/10915810701351186. PMID 17613130.
10. Unnate Atodriya; Roshni Baard; Siddi Upadhyya and Umesh Upadhyay. Antiuro lithiatic activity of Dolichos biflorus seeds. *Journal of Pharmacognosy and Phytochemistry*, 2013; 2(2): 209-213.
11. Michael Dickson and Jean Paul Gagnon. Key factors in the rising cost of new drug discovery and development. *Nature Reviews Drug Discovery*, 2004; 3: 417-429.
12. Kamboj V.P. Herbal medicine. *Current Science Association (JSTOR)*, 2000; 78(1): 35-39.
13. Archana R. Dhole, Vikas R. Dhole, Chandrakant S. Magdum, Shreenivas Mohite. Herbal Therapy for Urolithiasis: A Brief Review. *Research Journal of Pharmacology and Pharmacodynamics*, 2013; 5(1): 6-11.
14. Subramoniam. A, P. Pushpangadan. Development of phytomedicines for liver disease. *Indian Journal of Pharmacology*, 1999; 31(3): 166-175.
15. Sanjay M, Jachak and Arvind saklani. Challenges and opportunities in drug discovery from plants. *Current Science Association (JSTOR)*, 2007; 92(9): 1251-1257.
16. Ramachandran. S, Vijayakumar. T.M, Saisandeep. V, Ramsai. K and Dhanaraju. M.D. Antilithiatic Activity of Poly Herbal Extracts on Ethylene glycol-Induced Lithiasis in Rats. *European Journal of Biological Sciences*, 2011; 3(2): 36-39.