

**EVALUATION OF ANTHELMINTIC ACTIVITY OF INDIAN HERBS**

Kulkarni Chitrarekha Girish, Lohar Bhagwan Namdev, Jadhav Shital Tanaji*, Salunkhe Satyajeet Sunil

Government College of Pharmacy, Vidyanagar, Karad (India) 415124

***Corresponding author e-mail:** shitaljadhav67@yahoo.com

ABSTRACT

Helminthic infections are major cause of chronic ill health among tropical people. Traditional medicines contain several plant products for eliminating helminthic worms. However, standardization of individual ingredients is lacking in many cases. Present investigations are undertaken to ascertain anthelmintic activity of aqueous and ethanolic extracts of *Terminalia belerica* fruit, *Boerhavia diffusa* leaf and *Saxifraga granulata* root against *Pheretima posthuma* and *Eisenia foetida*. Albendazole 20mg/ml is used as a reference standard drug and normal saline as control. The result is expressed in terms of time for paralysis and time for death. The study indicated significant anthelmintic activity of aqueous as well as ethanolic extracts of *Terminalia belerica* and *Boerhavia diffusa*. Ethanolic extract of *Saxifraga granulata* at concentration of 50mg/ml showed comparable anthelmintic activity with reference standard against *Pheretima posthuma*.

KEYWORDS: Anthelmintic activity, *Pheretima posthuma*, *Eisenia foetida*, *Terminalia belerica*, *Boerhavia diffusa*.

INTRODUCTION

Helminthic infestations are very common in human beings, affecting a large proportion of population. In tropical developing countries they pose a big threat to community health by contributing to prevalence of anaemia, malnutrition, pneumonia, etc⁽¹⁾. Helminths also affect millions of livestock resulting in considerable economic losses in domestic and farm yard animals.

An ideal anthelmintic should have a broad spectrum activity & be free from toxicity to the host & should be cheap. Also, it should achieve the cure with a single therapeutic dose. Because of limited availability and affordability of modern medicines, financially weaker population depends greatly on traditional remedies. Multidrug resistance shown by most of the parasitic worms necessitates search of newer anthelmintic drugs. Since many effective drugs find their origin in traditionally exploited herbs, a systematic study of traditionally used herbs and exploration of their medicinal properties becomes inevitable.

Three Indian herbs are therefore studied with respect to their anthelmintic activity.

MATERIALS AND METHODS**Collection and identification of Plants**

Information regarding vernacular name, plant parts used, etc. was collected and authentic identification of plants was done with the help of different flora and monographs⁽²⁻¹⁶⁾.

Extraction Procedure

Plant parts (Refer table 1) were air dried and reduced to fine powder in a pulveriser. Aqueous and alcoholic extracts of respective plant parts were concentrated⁽¹⁷⁾.

Experimental animals

Pheretima posthuma (Indian Earthworm) were collected from moist soil, near Government College of Pharmacy, Vidyanagar Karad (India) and *Eisenia foetida* were collected from Vermiculture plant, Saidapur, Karad (India). The collected worms were washed with normal saline (0.9% w/v NaCl) to remove soil and dirt. The collected *Pheretima*

posthuma measured 14cm in length and 3mm in diameter whereas, *Eisenia foetida* were 17cm long and 6mm wide approximately. Identity of worms was confirmed by a registered veterinary practitioner at Kanfe Globuz pet polyclinic, Kolhapur (India).

Evaluation of anthelmintic activity

Adult *Eisenia foetida* & *Pheretima posthuma* were used to evaluate anthelmintic activity of various phytoextracts because of their anatomical and physiological resemblance with intestinal round worm parasites of human beings⁽¹⁸⁻²²⁾. Test samples of various extracts were prepared at the concentrations, 20mg/ml and 50mg/ml in double distilled water

Anthelmintic activity of extracts was assessed by comparing the paralysis time and death time required for worms in presence of various extracts and comparing it with 20mg/ml Albendazole as a reference standard and normal saline as control⁽²³⁻²⁵⁾. All experiments were done in triplicate. The time for paralysis was noted when no movement of any sort could be observed except when the worms were shaken vigorously. Time for death of worms were recorded after ascertaining that the worms neither moved when shaken vigorously nor when dipped in warm water (50°C)^(21,25-27).

RESULTS

1. Study with respect to *Eisenia foetida*:

Aqueous and alcoholic extracts of *Terminalia belerica* took 35 and 30 minutes respectively, for *Eisenia foetida* death at the concentration of 20 mg/ml whereas 19 and 16 minutes were required at the concentration of 50 mg/ml (Refer figure 1-4). Comparable activity with *Terminalia belerica* was shown by *Boerhavia diffusa* at corresponding extract concentrations. It took longer period by *Saxifraga granulata* to paralyse and kill *Eisenia foetida* as compared to the reference standard Albendazole.

2. Study with respect to *Pheretima posthuma*

Aqueous and alcoholic extracts of tested plant parts against *Pheretima posthuma* indicated nearly equivalent to better activity in case of *Terminalia belerica* and *Boerhavia diffusa* (Refer figure 5-8). *Saxifraga granulata* alcoholic extract showed

equivalent death activity at 50mg/ml in comparison with reference standard.

DISCUSSION

Percent decrease study with respect to paralysis and death time of *Eisenia foetida* clearly suggested a dose dependent result under the set of conditions applied during evaluation. A sharp decrease over 66% was shown by ethanolic extract of *Terminalia belerica* at the concentrations of 50mg/ml as compared to 37.5% at 20 mg/ml. An equivalent decrease in death time was observed by equivalent concentration of aqueous extract of *Terminalia belerica*. A similar dose dependent activity was observed for *Boerhavia diffusa* extract. The best time dependent activity was shown by ethanolic extract of *Terminalia belerica* at 50mg/ml concentration (Refer table 2).

Paralysis and death time study with respect to *Pheretima posthuma* showed interesting features with *Boerhavia diffusa* ethanolic extract at 50mg/ml concentration showing the best killer (Refer table 3). Ethanolic extract of *Terminalia belerica* at equivalent concentration took up the second position. Among the extracts and concentrations of *Saxifraga granulata* only ethanolic extract at 50 mg/ml concentration had anthelmintic activity comparable with reference standard.

CONCLUSION

From the above results it is concluded that ethanolic extract of fruit of *Terminalia belerica* at 50mg/ml concentration has showed best anthelmintic activity among all studied plants against *Eisenia foetida*. Ethanolic extracts of plants are proved to be better paralyzers and death inducers as compared to aqueous extracts.

Ethanolic extract of *Boerhavia diffusa* proved to be the best death inducer for *Pheretima posthuma* at concentration of 50mg/ml.

The present study can be considered as step towards rationalisation of herbal drugs. *In vivo* studies need to be done as a next step towards the same.

ACKNOWLEDGEMENTS

Authors are thankful to all who helped us for this study directly or indirectly.

Table 1: Indian herbs and their parts under current study

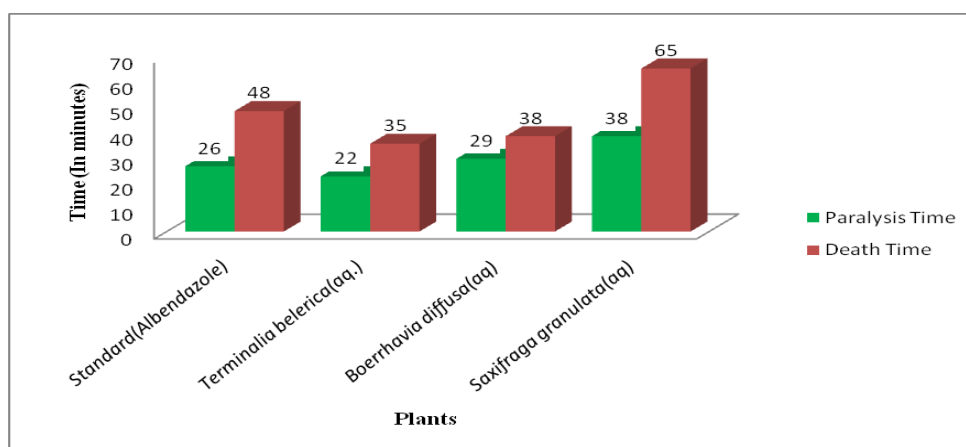
Sr. No.	Plant names	Vernacular name	Part used
1	<i>Terminalia belerica</i>	Baheda	Fruit
2	<i>Boerhavia diffusa</i>	Punarnava	Leaves
3	<i>Saxifraga granulata</i>	Pashanbheda	Roots

Table 2: Anthelmintic activity of plant extracts against *Eisenia foetida*

Sr. No.	Plant names	Type of Extract	Concentration of extract (mg/ml)	Percentage decrease in Paralysis time	Percentage decrease in Death time
1	<i>Terminalia belerica</i>	Aqueous	20	15.38	27.08
			50	34.62	60.17
		Ethanollic	20	7.69	37.5
			50	46.15	66.67
2	<i>Boerhavia diffusa</i>	Aqueous	20	-11.54	20.83
			50	19.23	50
		Ethanollic	20	15.38	39.38
			50	42.30	58.33
3	<i>Saxifraga granulata</i>	Aqueous	20	-46.15	-35.41
			50	-26.93	-8.33
		Ethanollic	20	-50	-29.16
			50	-38.46	-4.17

Table 3: Anthelmintic activity of plants against *Pheretima posthuma*

Sr. No.	Plant names	Type of Extract	Concentration of extract (mg/ml)	Percentage decrease in Paralysis time	Percentage decrease in Death time
1	<i>Terminalia belerica</i>	Aqueous	20	-3.33	2.38
			50	23.33	16.67
		Ethanollic	20	10	7.14
			50	30	19.4
2	<i>Boerhavia diffusa</i>	Aqueous	20	-26.66	-7.14
			50	23.33	23.80
		Ethanollic	20	13.33	26.19
			50	46.67	40.48
3	<i>Saxifraga granulata</i>	Aqueous	20	-30	-30.95
			50	-6.67	-11.90
		Ethanollic	20	-16.67	-23.80
			50	3.33	2.38

**Figure 1: Anthelmintic activity of aqueous plant extracts at 20mg/ml against *Eisenia foetida***

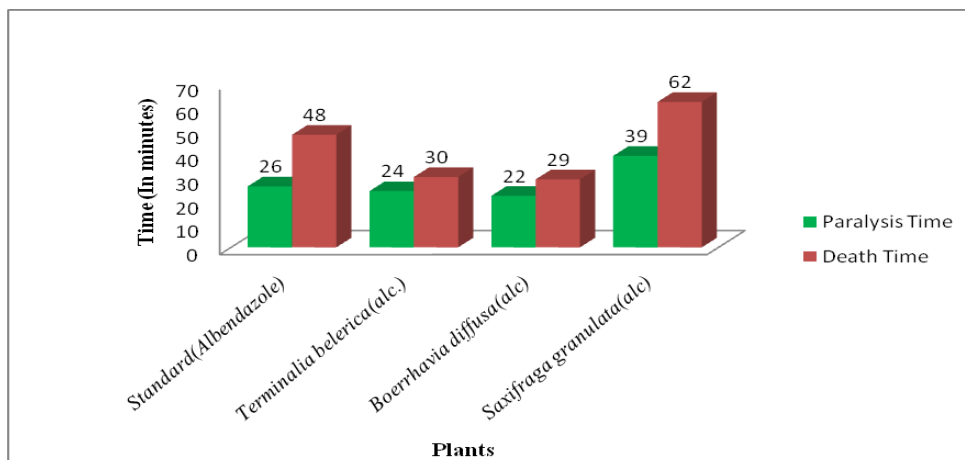


Figure 2: Anthelmintic activity of ethanolic plant extracts at 20mg/ml against *Eisenia foetida*

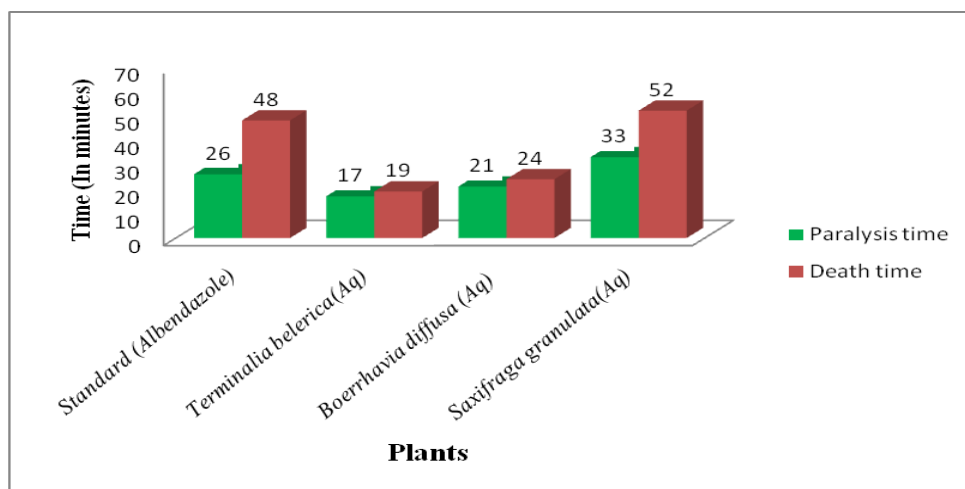


Figure 3: Anthelmintic activity of aqueous plant extracts at 50mg/ml against *Eisenia foetida*

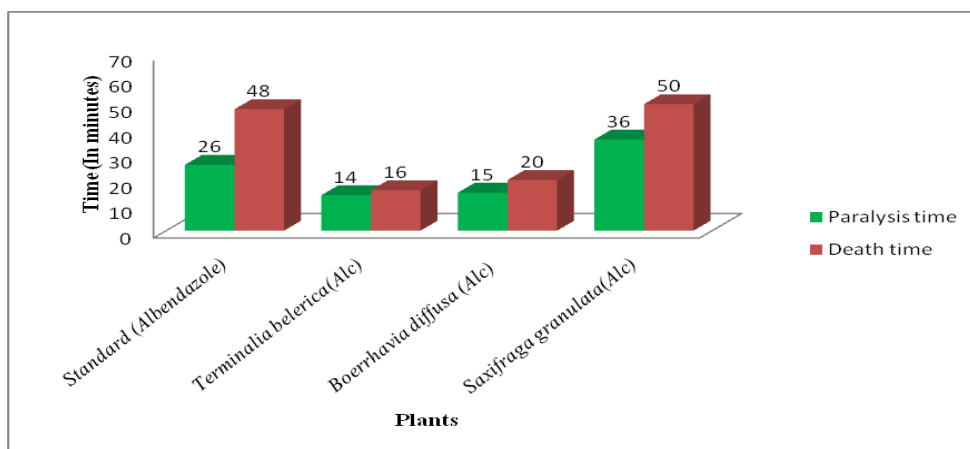


Figure 4: Anthelmintic activity of ethanolic plant extracts at 50mg/ml against *Eisenia foetida*

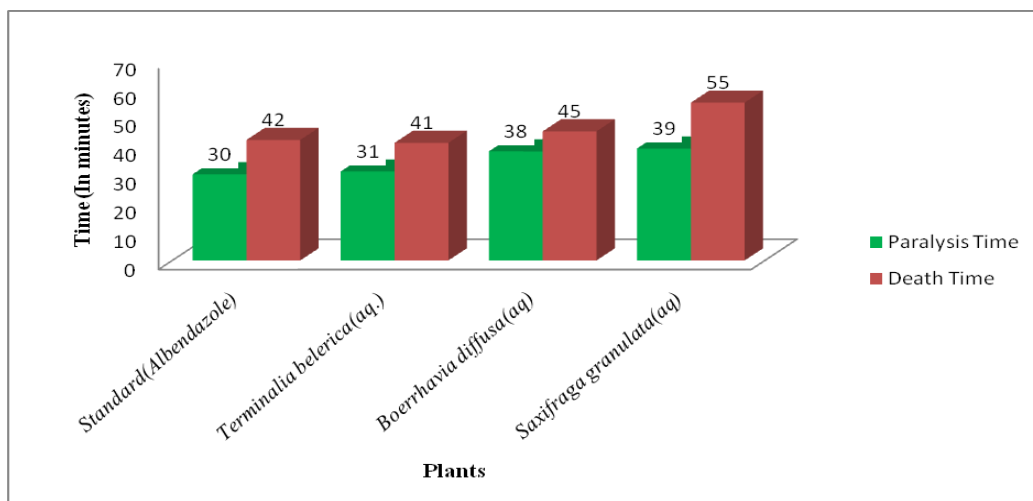


Figure 5: Anthelmintic activity of aqueous plant extracts at 20mg/ml against *Pheretima posthuma*

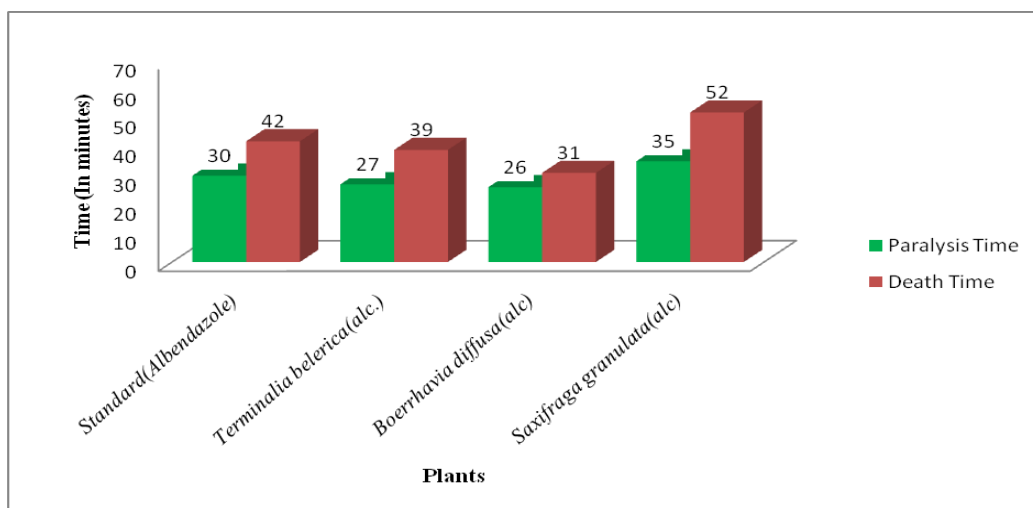


Figure 6: Anthelmintic activity of ethanolic plant extracts at 20mg/ml against *Pheretima posthuma*

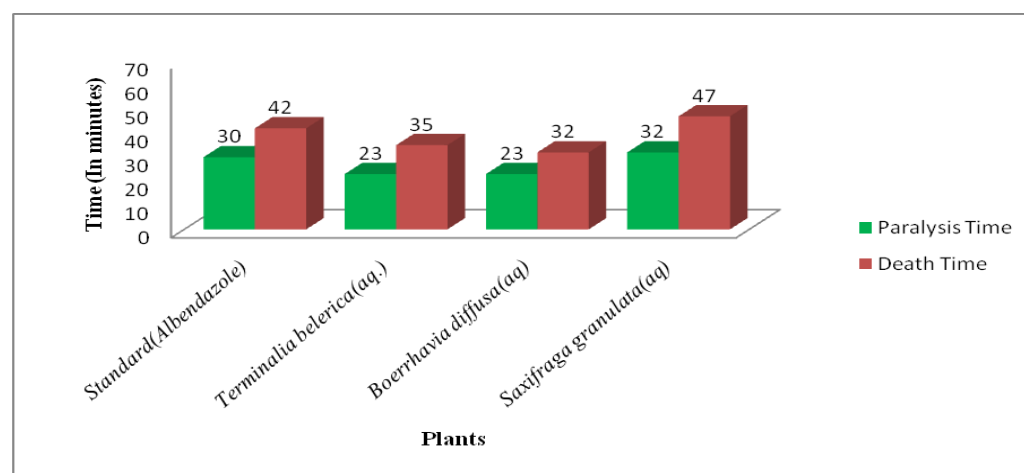


Figure 7: Anthelmintic activity of aqueous plant extracts at 50mg/ml against *Pheretima posthuma*

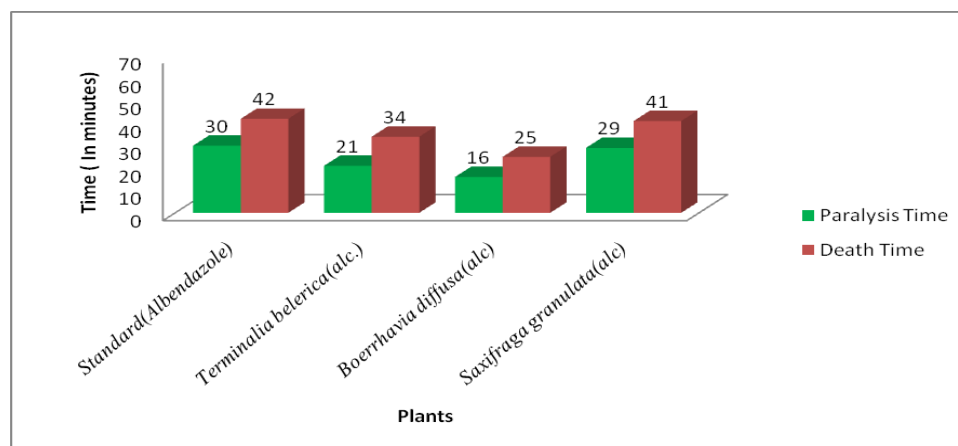


Figure 8: Anthelmintic activity of ethanolic plant extracts at 50mg/ml against *Pheretima posthuma*

REFERENCES

- Mali RG, Mehta AA. Nat Prod Rediance, 2008; 7(5): 466-475.
- Hooker JD. The flora of British India. L. Reeve and Co. Ltd. Kent. Vols 1-7; 1872-1897.
- Sharma BD, Balakrishnan NP and Sanjappa M. Flora of India. Vol II., BSI Calcutta ,Deep printers new Delhi : 1993
- Sharma BD and Sanjappa M, Flora of India. Vol III.,BSI Calcutta. Deep Printers New Delhi: 1993.
- Nadkarni AK. Indian Materia Medica. 3rd ed., Bombay; Popular Prakashan : 1954.
- International Library Association. Medicinal plants sourcebook, India. International Library Association, Switzerland:1996.
- Wilson DE and Reeder DM. Manual species of the World. 3rd ed., Johns Hopkin University press: 2005.
- Bhattacharjee SK and De LC. Medicinal Herbs and Flowers. Jaipur (India) ;Avishkar publishers: 2005.
- Small Indian civet- *Viverricula indica* http://www.vietlinh.vn/langviet/toilamnd/cogni/khac/cyhuong_muskcat.html. October 20, 2008.
- Ziziphus xylopyrus*. [http:// zipcodezoo.com/Plants/Z/Ziziphus_xylopyrus/](http://zipcodezoo.com/Plants/Z/Ziziphus_xylopyrus/). October 20, 2008.
- Nadkarni AK. Indian Materia Medica. 3rd ed., Bombay; Popular Prakashan: 1954.
- Indrajao- Flowers of India. [http:// www. Flowers of india.net/catalog/slides/Indrajao.html](http://www.Flowersofindia.net/catalog/slides/Indrajao.html). October 20, 2008.
- Wilson DE and Reeder DM, Mammal species of the world- A Taxonomic and Geographic Reference. 3rd ed., Johns Hopkins University Press: 2005.
- Blepharis-Wikispecies.<http://species.wikimedia.org/wiki/Blepharis>. October 20, 2008.
- Whitaker R. Common Indian Snakes- A Field Guide.1987.
- Medicinal Plants of Tamil Nadu. [http:// envis.frlht.org.in/checklist/TN.pdf](http://envis.frlht.org.in/checklist/TN.pdf). October 20, 2008.
- Mehta JH, Kulkarni CG, Jadhav ST, Deshpande AM, Bhise SB. Int J Res in Pharmacy & Chem,2013; 3(4):743-747.
- Kaur S, Kumar B, Puri S, Tiwari P, Divakar K. Int J Drug Dev & Res ,2010;2(4) : 758-763.
- Ahmed R, Sahu RK, Samele KK, Roy A, Dwivedi J. Appl Sci Res.,2010; 2(1):398-400.
- Khadse CD, Kakde RB. Int J Res Pharm Sci., 2010; 1(3): 267-269.
- Sangeetha J, Soundarya K, Santosh K, Sindhura C. Res J Pharm Biol & Chem Sci.,2010; 1(3) : 715-718.
- Patel HR, Fursule RA. Int J Phytomed & related Industries., 2009, 1(2):113-115.
- Ghule SC, Chaudhari SR, Chavan MJ. Int J Pharmacy & Pharm Sci., 2011; 3(1): 143-144.
- Aswar M, Aswar U, Watkar B, Vyas M, Wagh A, Gujar KN. Int J Green Pharm., 2008;2(3):170-172.
- Patil SH, Deshmukh PV, Sreenivas SA, Sangameshwar K, Vijapur LS. Int J Green Pharm,2013; 34-36.
- Sundeeep Kumar HK, Bose A, Raut A, Sahu SK, Raju MB. J Basic Clin Pharmacy.,2010; 1(2):103-105.
- Kosalge SB, Fursule RA. Asian J Pharm Clin Res., 2009; 2(2):69-71.