The Antihelminthic Activities of Dita (A lstonia scholaris) Bark and Tamarind (Tamarindus indicus) Seeds: A Comparative Study

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Abstract

The study was done to compare the antihelminthic activities of dita (A lstonia scholaris) bark and tamarind (Tamarindus indicus) seeds using adult earthworms, Pheretima posthuma. It was found out that the dita bark extract showed more potent antihelminthic activity as compared to the tamarind seeds. Further, the antihelminthic activity of the two plant extracts were also compared to the antihelminthic activity of a known standard drug, mebendazole Results showed that the dita bark crude extract has stronger antihelminthic action against the worm than mebendazole.

Introduction

Plants provide the principal ingredients in medicines used in most medical traditions globally. According to the World Health Organization (WHO), 80% of people in the world use herbal medicine as their primary form of health care. In the past, our ancestors made new discoveries of the healing power of plants through trial and error. Nowadays, more scientific approaches are commonly applied to understanding the medicinal properties of plants and developing new products.

Modem pharmaceutical drugs have been discovered through research into the physiological effects of chemicals found in plants. Some of these drugs are still derived directly from plants while others are now synthesized. Chemicals in some plants are extracted and transformed, providing the building blocks of drugs.

Antihelminthic drugs are medicinal drugs used to rid humans or other host animals of infestations by parasitic worms such as tapeworms, roundworms, pinworms, trichinae, flukes, whipworms, schistosomes and filariae. These drugs are also known as vermifuges. They act by attacking the worms' neuromuscular or respiratory systems, interfering with their metabolism, or making them more susceptible to attack by the host's macrophages. Antihelminthics include mebendazole, pyrantel pamoate, diethylcarbamazine and niclosamide, among others. The drugs are generally more effective against gastrointestinal infestations than muscular infestations (Microsoft Encarta, 2007).

Alstonia scholaris, commonly known as dita plant, is found throughout tropical Eastern Asia and the Malayan Archipelago. It is a large tree, with smooth, entire, thick leaves disposed in whorls. The flowers resemble those of *Alstonia constricta*, but differ in having corolla tubes about three times as long as the calyx and shorter pubescent lobes. The pods are slender and over a foot long. The tree is a native of the East Indies and the Moluccas, the bark which is sold in commerce comes from the Philippines and neighboring islands and is the portion used in medicine.

Dita bark is ½ inch thick, and is found in the market in irregular sizes rom 1 to 2 inches wide and from 3-6 inches long. Externally, it is of a mottled pinkish or brownish and white color, rather smooth but marked by shallow fissures which are raised upon the edges and scarcely extend through the corky layer. The cork, a very thin layer, is brownish in color. Internally, the color of the bark is light, slightly striated with yellowish layers or grains. In texture, it is granular and brittle, resembling wild cherry bark from old trees. The taste is slightly bitter, free from astringency, not unpleasant and maybe compared to the after taste of wild cherry bark and in like manner, the bark is gritty between the teeth.

Tamarind (*Tamarindus indicus*) is a slow-growing, massive tree that can reach a height of 80 or even 100 feet (24-30 m), spread a crown of 40 ft (12 m) and develop a very large trunk of 25 \checkmark 7.5 m) in circumference.

It is a long-lived tree with high resistance to wind, dark-gray and rough bark and strong, supple branches that are gracefully drooping at the ends. The mass of bright-green, fine, feathery foliage is composed of pinnate leaves, each having IO to 20 pairs of oblong leaflets, which fold at night. The leaves are normally evergreen but may shed briefly in very dry areas during the hot season. Inconspicuous, inch-wide flowers, borne in small racemes, are 5-petalled (2 reduced to bristles), yellow with orange or red streaks. The flower buds are distinctly pink due to the outer color of the four sepals, which are shed when the flower opens. The fruits are curved and bulged pods, borne in great abundance along the new branches. The pods are cinnamon-brown or grayish-brown and tender-skinned with green, highly acid flesh and soft, whitish and under developed seeds. As they mature, the pods fill out and the juicy, aciduous pulp turns brown or redish brown. Then, the skin becomes a brittle, easily cracked shell and the pulp dehydrates naturally to a sticky paste enclosed by a few coarse strands of fiber. The seeds are hard, glossy-brown and each is enclosed in a parchment like membrane.

In native practice, the pulp is applied on inflammations, used as a gargle for sore throat and mixed with salt, as a liniment for rheumatism. It is administered to alleviate sunstroke, digitalis poisoning and alcoholic intoxication. The pulp is said to aid in the restoration of sensation in cases of paralysis. Tamarind leaves and flowers, dried or boiled, are used as poultices for swollen joints, sprains and boils. Lotions and extracts made from them are used in treating conjunctivitis, dysentery, jaundice, hemorrhoids and various other ailments, because of their antiseptics and vermifuge properties.

In the Philippines, folk people claim that there are some plants that help in the elimination of helminthes. After drinking a decoction of the bark of dita or the powdered seeds of tamarind seeds for three or more days, helminthes were found evacuated from the stools of children and adults.

These premises led the researchers to compare the anthelminthic properties of these two plants, dita (*Alstonia scholaris*) bark and tamarind (*Tamarindus indicus*) seeds.

Objectives of the Study

The study aimed to compare the antihelminthic activities of dita (Alstonia scholaris) bark and tamarind (Tamarindus indicus) seeds.

Specifically, it sought to:

- I. determine the antihelminthic activities of the two plant extracts in terms of paralysis time and death time
- 2. compare the two extracts to the antihelminthic drug, mebendazole.

Scope and Delimitation

This study was limited to the determination of the antihelminthic activities of dita bark and tamarind seeds in terms of the paralysis time and death time they had caused to the earthworms. Comparisons of the antihelminthic properties of these two plants were also determined against the standard drug, mebendazole.

Only the bark of the dita tree and the seeds of the tamarind were used as the extracts under investigation.

Ninety percent (90%) ethanol was used as the solvent in the extraction process.

Adult earthworms (*Pheretimaposthuma*) were the helminthes used in the study.

The experiment was conducted at the Natural Products Chemistry and Biotechnology Laboratory, Science Complex I, University of Northern Philippines, Vigan City from June 8-15, 2009.

Review of Related Literature

Helminth infections are among the most common infections in man, affecting a large proportion of the world's population. In developing countries, they pose a large threat to public health and contribute to the prevalence of malnutrition, anemia, eosinophilia and pneumonia. Although the majority of infections due to the worms are generally limited to tropical regions, they can occur to travelers who have visited those areas, and some of them can be developed in template climates (Bundy, 1994).

Parasitic diseases causing severe morbidity include lymphatic filariasis (a cause of elephantiasis), onchocerciasis (river blindness) and schistosomiasis. These infections can affect most populations in endemic areas with major economic and social consequences.

The limited availability and affordability of pharmaceutical medicines mean that the world's population depends to a great extent on traditional medical remedies, and some 20,000 species of higher plants are used medicinally throughout the world. Many well-known drugs listed in the modem pharmacopoeia have their origins in nature, including for example, quinine from the bark of Cinchona tree for the treatment of malaria, which has been followed by the subsequent development of the synthetic derivatives chloroquine, amodiaquine, primaquine and mefloquine. More recently, the wider recognition of the antimalarial activity of artemisinin from the herb *A temisia annua* has led current research to focus on the development of a large nwnber of synthetic and semisynthetic compounds, which are more active than artemisinin (Tagboto and Townson, 2001).

In the study of the antiparasitic activity of *Melia azedarach*, a native tree growing in Argentina, Szewczuk,et.al. found that the extracts obtained from the drupes of M. azedarach **are** active against both the tapeworm *Taenia solium* and the earthworms tested. The drupe extracts are comparatively more active than piperazine phosphate against *Taenia solium*, The antiparasitic activity against this tapeworm was better than that of piperazine phosphate (80 minutes at 0.1% and 56 minutes at 0.2%) at drupe extract concentrations of 0.1 % and 0.2% (mean death values of 52 and 32 minutes, repectively). The findings support the use *of Melia azedarach* drupes as antiparasitic in the traditional medicine. (Szewczuk et. al, 2003)

A similar study on the antihelmintic activity of a polyherbal preparation containing the herbs *Thespesia populnea* (bark), *Terminalia alata* (bark), *Clematis triloba* (roots) and *Ceratophyllum demersum* (leaves) was conducted using adult earthworm Pheretirna posthuma as test organisms. The aqueous and ethanolic extract of the crude drug of different concentration were tested which involved detennination of paralysis time and time to kill the worms. Piperazine citrate was used as standard and it was found that the ethanolic extract of the polyherbal preparation formula is better than the aqueous extract of the polyherbal preparation formula.

Morever, the study reveals that the ethanolic extract of *Thespesia popu Zea* (bark), *Terminalia alata* (bark), *Clematis triloba* (roots) and *Ceratophyllum demersum* (leaves) showed marked and potent antihelminthic activity (although all these plants alone exhibit antihelminthic activity but when combined will give m ore potent activity) than the aqueous axtract of *Thespesia populnea* (bark), *Terminalia alata* (bark), *Clematis triloba* (roots) and *Ceratophyllum demersum* (leaves) as compared to standard drug piperazine citrate (Dwivedi, et.al., 2009).

Most antihelminthic drugs are only active against specific parasites, some are also toxic. Before treatment, the the parasites must therefore be identified using tests that look for parasites, eggs or larvae in feces, urine, blood, sputum, or tissues. Thus, niclosamide is used against tapeworms, but will not be effective for the treatment of pinwonn or roundworm infestations, because it acts by inhibiting ATP production in tapeworm cells. Thiabendazole (Mintezole) is the drug usually prescribed for treatment of threadworms but a similar drug, Mebendazole (Ver \neg ox) works better on whipwonn by disrupting the microtubules of this worm. Praziquantel is another drug that acts by altering the membrane permeability of the worms.

Common side effects of antihelminthic drugs include dizziness, drowsiness, headache, sweating, dryness of the mouth and eyes, and ringing in the ears. Side effects usually wear off as the body adjusts to the drug and do not usually require medical treatment. Thiabendazole may cause the urine to have an unusual odor that can last for **a** day after the last dose. Other side effects of antihelminthic drugs, such as loss of appetite, diarrhea, nausea, vomiting or abdominal cramps are less common. If they occur, they are usually mild and do not require medical attention (Gale Encyclopedia of Medicine, 2008.)

In the study of Supnet (2007), a phytochemical analysis on the dita bark and leaves were conducted. Results of the phytochemical analysis indicate that the leaf of dita contains alkaloids, sterols, flavonoids, glycosides, saponins and resins. The bark extract, on the other hand, are positive for alkaloids, flavonoids and resins.

Conceptual Framework

The researchers were guided by the paradigm presented below:

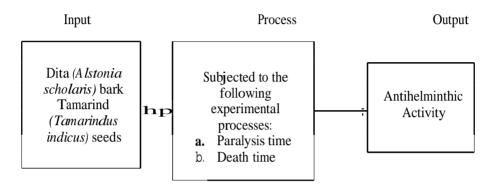


Figure 1. The Research Paradigm

The paradigm shows that dita (*Alstonia scholaris*) bark and tamarind (*Tamarindus indicus*) seeds were subjected to the two experimental processes to determine their paralysis time and death time as a measure of their antihelminthic activities.

Operational Definition of Terms

The following tenns were defined operationally for a clearer understanding of the study:

Antihelminthic activity. This refers to the property of the extract to immobilize and cause death to the living worms. In this study antihelminthic activity were measured in terms of paralysis time and death time.

Paralysis time. It refers to the time elapsed starting from the application of the different treatments to the time the earthworm loses muscle tonicity and no appreciable movement was observed.

Death time. It refers to the time elapsed starting from the application of the treatment to the time when complete loss of movement was observed on the earthwonn even after immersion in hot water was done.

Methodology

Collection of Plant Materials

The Dita (*Alstonia scholaris*) barks were collected around the vicinity of the University of Northern Philippines, Tamag, Vigan City while the tamarind (*Tamanidus indicus*) seeds were collected from the fruits eaten by residents of Vigan, llocos Sur.

Preparation of the Extracts

The 200 grams of pulverized powder of tamarind (*Tamarindus indicus*) seeds were placed in a 500 ml Erlenmeyer flask and enough 90% ethanol was added to submerged the sample. Similarly, the bark of dita tree were chopped into small pieces, weighed (200 grams), were placed in another Erlenmeyer flask and enough ethanol was also added to submerged the bark. The Erlenmeyer flasks were stoppered and were allowed to stand for 48 hours. The ethanolic extracts of dita bark and tamarind seeds were then filtered using a Buchner funnel and concentrated using evaporation process until 20 ml of extract was obtained.

Experimental model

Adult earthworms *Pheretima posthuma* were collected (due to its anatomical and physiological resemblance with the intestinal roundworm parasites of human being) from moist soil obtained from the University of Northern Philippines grounds.

Experimental Procedures

These antihelminthic procedures were adapted from a similar study by Victor D. Szewczuk (2003) entitled "Antiparasitic Activity of Melia azedarach Growing in Argentina".

The antihelmiothic activities were done in four groups. The first group made use of the crude extracts of dita bark and tamarind seeds. The second group made use of the 0.5 % dilution of the dita bark and tamarind seeds extracts. The third group used the 0.75 % of the dita bark extract and tamarind seed extract. The fourth group made used of the 1% dilution of the dita bark and tamarind seed extracts.

In each of the groups, 27 equal sized earthworms (IO cm) were used in the experimental investigation. Triplicates were done for each treatment. Three earthworms were placed in each Petri dish

Mebendazole (Antiox) was taken as the standard drug and served as the positive control.

The dita bark extracts and tamarind extracts were diluted using distilled waler to obtain 0.5, 0.75 and 1.0% concentration. Two ml of each of these dilutions were used in the antihelminthic investigation

Normal saline solution was used as the negative control.

Observations were made for the time taken to paralyze the population of worms and time for the death of individual worms. Paralysis was said to occur when the worms do not revive even in nonnal saline solution. The death of the worms were ascertained by transferring the worms into a beaker containing hot water at 50 C which stimulated and induced movements if the worm was alive.

Procedures were repeated using the different concentrations of the dita bark and tamarind seed extract

Statistical Treatment of Data

Average mean of the time taken to paralyze the worms and time of death were taken and compared.

Results and Discussions

The results of the antihelminthic activity of the two extracts are presented in the following tables.

Table 1. Antihelminthic Activity of the crude extracts of Dita Bark and Tamarind Seeds

Treatments	Paralysis Time	Mean Average	Death Time	Mean _Average
Saline Solution				
Dita Bark Crude	5 mins	4.67mins	5 mins	5.33 mins
Extract	4 mins		6mins	
	5 m i s		5 mins	
Tamarind Seeds	12 mins	10.67 mins	14mins	15 mins
Crude Extract	10 mins		16 mins	
	10 mins		15 mins	
Mebendazole	9 mins	9 mins	10 mins	9.67 mins
	8 mins		10 mins	
	10 mins		9 mins	

Results on Table 1 shows that the crude dita bark extract gave the fastest time (x= 4.67 mins) to paralyze the wonns as compared to the crude tamarind seed extracts (x= 9 mins). When compared to the standard drug, crude dita bark extract showed a stronger activity against the earthwonn than mebendazole (x=9 mins). The crude tamarind seed extract (x = 10.67 mins) had less activity to paralyze the worms as compared to the standard drug, mebendazole.

With regards to the death time of the worms, crude dita bark extract caused a faster death to the worms with a mean time of 5.33 minutes as compared to the crude tamarind extract which had a mean time of IS minutes. When compared to the standard drug, the crude dita bark extract caused the death of the worns in 5.33 minutes as against mebendazole which is 9.67 minutes. The crude tamarind seed extracts showed the longest time to inflict death to the worns (15 mins).

Treatments	Paralysis Time	Mean Average	Death Time	Mean Average
Saline Solution				
1 % Dita Bark	6 mins	6.33 mins	7mins	8 minus
Extracts' "	6 mins		9mins	
	7mins		8 mins	
1% Tamarind	16 mins	17 mios	20 mins	20.67 mins
Seeds Extract	20mins		22 mins	
	15 mins		20 mins	
Mebendazole	9 mins	9mins	10mins	9.67 mins
	8 mins		10 mins	
	10 mins •		9mins	

Table 2. Antihelminthic Act	ity of 1% Dita Bark and Tamarind Seed Ertract	S

Using I% dilution of the two extracts, it is gleaned on the table that the dita bark extract cause paralysis to the wonns at 6.33 mins as compared to the tamarind seed extract giving a 17 minutes paralysis time. Compared to mebendazole, dita bark extract still showed stronger activity to paralyze the worms at 6. 33 minutes as compared to mebendazole with a mean paralysis time of 9. The tamarind seed extract gave the longest paralysis time of 17 minutes.

The table also reveals that the 1% dita extract still cause mortality to the worms the fastest (x = 8 mins) as compared to the standard drug, mebendazole (x=9.67 mins) and tamarind seed extract (x=20.67 mins).

Treatments	Paralysis Time	Mean Average	Death Time	Mean Average
Saline Solution				
0.75 % Dita	9 mins	9.66 mins	12 mins	13 mins
Bark Extract	10 mins		13 mins	
	10 mins		13 mins	
0.75%	25 mins	29 ruins	42 mins	39 mins
Tamarind seeds	30 mins		35 mins	
extract	32 mins		40 mins	
Mebendazole	9mins	9 mins	10 mins	9.67 mins
	8 mins		IO mins,	
	10 mins		9 mins	

	Table 3.Antihelminthic Activi	tv of	0.75%	Dita Ba	rk and	Tamarind S	Seed Extracts
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Table 3 shows the results of the paralysis time the two extracts diluted to 0.75%. Dita bark extract gave a paralysis time of 9.66 minutes as compared to the 29 minutes paralysis time of the tamarind seed extracts and the 9 mins paralysis time shown by the standard drug, mebenclazole.

Point seventy-five percent (0.75%) dita bark extract gave a faster death time (x = 13 mins) as compared to 0.75% tamarind seed extract (x=39 mins).

Table 4. Antihelminthic Activit	y of 0.5%	Dita Bark and	Tamarind Seed extracts
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Treatment	Paralysis Time	Mean Average	Death Time	Mean Average
Saline Solution				
0.5% Dita bark	30 mins	32.33 mins	43 mins	41.33 mins
extract	32 mins		41 mins	
_	35 mins		40 mins	
0.5 % Tamarind	70 mins	74.33 mins	92 mins	90.33 mins
seeds extract	75 mins	_	89 mins	
	78 mins	-	90 mins	
Mebendazole	9 mins	9 mins	10 mins	9.67 mins
	8 mins		10 mins	
	10 mins		9 mins	

When the two extracts were diluted to 0.5%, results show that the extract of dita bark will paralyze the worms the faster with an average paralysis time of 32.33 minutes as compared to the tamarind seed extract (\checkmark 74.33 mins).

Similarly, the dita bark extract also gave a shorter death time to the wonns (x =41.33 mins) as compared to the death time showed by the tamarind seed extract (x = 90.33 mins.)

It can be inferred that in all the four groups of antihelminthic activity, dita bark extracts gave the most marked and potent antihelminthic activity in *terms* of paralysis time and death time as compared to the tamarind seed extract The tamarind seed extract only showed weak antihelminthic activities as evidenced by the longer duration time it gave to cause paralysis and death to the worms.

This implies that dita bark can be a good source of an antihelminthic drug comparable to the antihelminthic activity showed by the standard drug, mebendazole.

Conclusion

The dita bark contains substances that might prove potent as antihelminthic drugs **as** evidenced by the results obtained above.

Using the crude extracts, dita bark gave faster paralysis and death time as compared to the tamarind seed extract and the standard drug mebendazole.

In the 1%, 0.75% and 0.50% dilutions, dita extract again showed faster paralysis and death time compared to the same dilutions of tamarind seed extracts.

Recommendations

I. The antihelminthic activity of the two plant extracts should also be tested against parasitic organisms like pinworms, whipworms, tapeworms, amoebas, and protozoa.

2. Other plants should also be examined for their antihelminthic properties.

3. The antihelminthic substance of dita bark and tamarind seeds should be isolated and quantified.

4. Antihelminthic activity should also be tested to the different parts of the tamarind plant like its bark and its roots.

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