

Original Research Article

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**Anti Helminthis and Molluscicidal Activity of *Momordica dioica* against *Pheretima posthumous* and *Indoplanorbis exustus***

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The present study was intended to delineate the Anti Helminthis and Molluscicidal activity of *Momordica dioica* fruit extract. The crude extract was obtained by the employment of Soxhlet extraction method. The extract was used to screen the antagonistic activity *in vitro* by taking *Pheretima posthumous* and *Indoplanorbis exustus* as the study subjects. The results obtained proved the capability of extract to downregulate the biological activities of subjects. Mortality was induced by varying concentrations (100-1000 ppm/ml). Interestingly the antagonistic activity was in direct proportional with the extract administrated. The analysis of phyto-constituents revealed that the extract is composed of polyphenols, Saponins, Flavonoids etc. Therefore, it is assumed that the biological activities are complimented by the phyto-components. The results obtained from the current studies can be further exploited in the field nutraceuticals.

**Introduction**

Despite of several biological advantages of Helminthis and Molluscans it is also known for its diversified infection and vector abilities (Rabiu and Subhasish, 2011). Therefore, it is essential to develop efficient control agents against them. Trematodes, the causal agents of schistosomiasis and fascioliasis are important parasites of economic and public health implications in. Schistosomiasis affects over 240 million people worldwide, with up to 700 million individuals living at risk of infection (Kumarasingha *et al.*, 2016). The

disease caused up to 250,000 deaths per year in the last decade.

Similarly, Helminthis infection causes huge revenue in agro farming. There have been several synthetic chemical compounds and nanoparticles proven their anti Helminthis and Molluscicidal activities. However, it is also observed that these compounds are mostly toxic to the environment. In this juncture it is essential to develop suitable ethnobotanical agents against this threat (Rajeswari, 2014). *Momordica dioica* Roxb. is a perennial, dioecious climber included in *Cucurbitaceae*

family. *Momordica* genus contains about 80 species. According to the latest revision of Indian *Momordica*, there are six well identified species of which four are dioecious and two are monoecious (Bawara *et al.*, 2010). Although this genus is originated from Indo Malayan region, it is now found to grow in India, Bangladesh, Srilanka, Myanmar, China, Japan, South East Asia, Polynesia, Tropical Africa, and South America (Pusuloori *et al.*, 2017)

Its cultivation up to an altitude of 1500 meters in Assam and Garo hills of Meghalaya is reported. It is commonly known as spine gourd, teasel gourd or small bitter gourd worldwide whereas in Bangladesh it is known as kakrol and in India as kankro, kartoli, kantoli, kantola, kantroli, ban karola, or jangleekarela. Kakrol is about 5–7 meters in length, a popular summer vegetable of which its fruit, young twigs and leaves are used as vegetable (Ali *et al.*, 1991).

It is evident that the plant possess diuretic, laxative, hepatoprotective, antivenomous, antihypertensive, anti-inflammatory, antiasthmatic, antipyretic, antileprosy, antidiabetic, and antidepressant properties (Jeyadevi *et al.*, 2012). In this study I am delineating the Molluscicidal Activity and Anthelmintic activity of *Momordica dioica*.

## Materials and Methods

### Plant collection and extracts preparation

The fruit of *Momordica dioica* were collected from Botanical garden of St. Mary's College, Thrissur and the identity of plant was confirmed with the help of Dr. Sr. Meena K Cheruvathur, Assistant professor, Department of Botany, St. Mary's College, Thrissur, Kerala. The dried fruits were ground to a coarse powder and 50g of the powdered plant materials is subjected to extraction by using Soxhlet apparatus using 95% ethanol as the

solvent. The filtrate is collected after 72 hours. It is twice filtered using Whatman no.1 filter paper and extracts were concentrated by evaporation at room temperature. The dried residue is kept in refrigerator for further studies (Cheruvathur *et al.*, 2015).

### Collection of study subjects

*Pheretima posthumous* were collected from vermicomposting unit of St. Mary's College, Thrissur, Kerala and washed with normal saline to remove all fecal matter. Healthy adult *Indoplanorbis exustus* were collected from in and around Thrissur which was brought to Laboratory. The identity of collected organisms was confirmed by the help of Dr. Dalie Domnic, Head of the department, Department of Zoology, St. Mary's College, Thrissur, Kerala. Study subjects were acclimatized in aquaria for a minimum period of four days in holding tanks containing aerated, de-chlorinated tap water and washed sand (Williams *et al.*, 2014).

### Screening of anthelmintic activity

Test samples of plant extract was prepared at the concentrations 100 - 1000 ppm/ml in 10 ml of distilled water and five worms of approximately equal size (same type), about 0.32 g weight were placed in each petridish containing 10 ml of above test solution of extracts.

All the test solutions were prepared freshly before starting the experiments. Observations were made for the time taken for paralysis was noted when no movement of any sort could be observed except when the worms were shaken vigorously. Time for death of treating worms were recorded after ascertaining that worms neither moved when shaken vigorously nor when dipped in warm water. Standard drug Albendazole is taken as control for comparison (Murugamani *et al.*, 2012)

### **Screening of Molluscicidal activity**

Evaluation of molluscicidal activity of *Momordica dioica* against snails was investigated as recommended by the World Health Organization. The solution was prepared by adding varying concentration (100-1000 ppm/ml) of plant extract in distilled water. The snails were exposed in groups of five for 24 to 48 hours to 10 ml of each concentration of ethanolic fruit extract of *Momordica dioica*. Control organisms were similarly immersed in dechlorinated water. Snail mortality was identified by the contraction of the body within the shell; no response to a needle probe was taken as evidence of death (Omobhude *et al.*, 2017).

### **Phytochemical analysis of plant extract**

Phytochemical screening was carried out to assess the qualitative chemical composition of crude ethanolic extracts of *Momordica dioica*. Standard screening tests using conventional protocol, procedure, and reagents were conducted using standard procedures to identify the constituents as described previously (Talukdar and Hossain, 2014).

### **Results and Discussion**

#### **Effect of plant extract in causing death and paralysis in *Pheretima posthumous***

Ethanolic fruit extract of *Momordica dioica* possess dose dependent anthelmintic. At higher concentrations, it produces paralytic effect much earlier and the time to death was shorten. At higher doses, the plant extract induces 100% mortality (Figure 1). It was observed that time required for inducing paralysis and causing death is in directly proportional that with the concentration of extract administrated. Time taken for inducing paralysis was only 20 minutes when the concentration was 1000 ppm/ml whereas the

time of survival was prolonged to 90 minutes in lower concentration of antagonist.

A similar kind of activity was reported in aqueous leaf extract of *Annona muricata* L. (Annonaceae) against *Haemonchus contortus* from sheep (Ferreira *et al.*, 2013). Similarly plant extracts from Brazilian savanna also possessed Anthelmintic activity (Oliveira *et al.*, 2017). Generally, Albendazole eliminate worm chiefly by inducing flaccid paralysis interestingly plant extract also showed same clinical symptoms therefore we assume that the mechanism of plant extract is similar to that of Albendazole (Haque *et al.*, 1993).

#### **Effect of plant extract in causing death in *Indoplanorbis exustus***

The toxic effect of active plant extracts was evident in treated snails with ethanolic fruit extract of *Momordica dioica*. There was a visible swelling in the cephalopodal mass. High doses of active plant extract caused the cephalopodal mass of each snail to become severely swollen, turgid and failing to mechanical stimulus with blunt needle. Mucous secretion was observed over most of the foot. Hundred percentage mortality is observed in all treated concentrations (Figure 2). Similarly *Solanum* Species showed Molluscicidal activity against *Biomphalaria alexandrina* (El-Sherbini *et al.*, 2009) as well as (Abdel-Haleem, 2013) these observations suggests that the phytoconstituents of plants constitute its activity and it is not depending upon its trait.

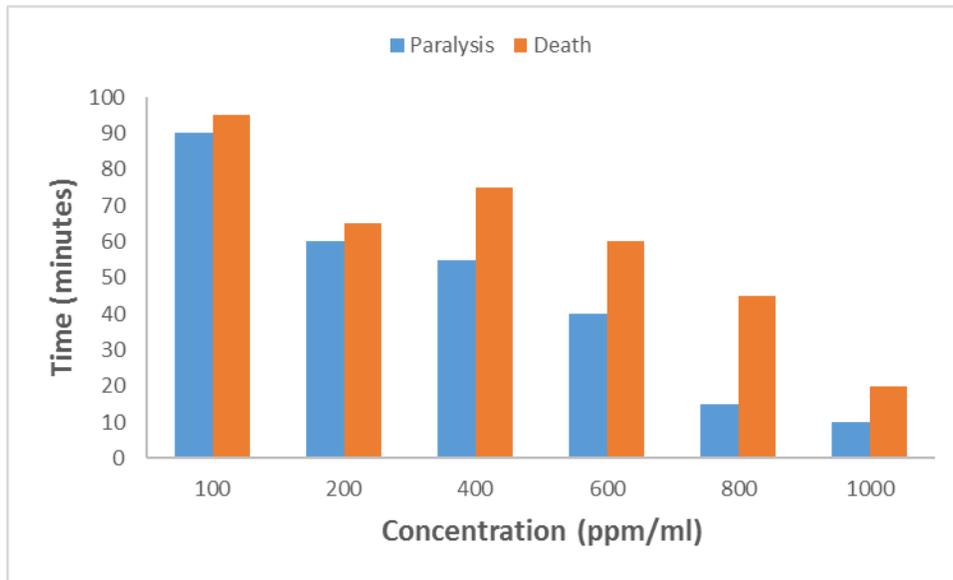
### **Phytochemical analysis of plant extract**

Phytochemical screening of ethanolic fruit extract of *Momordica dioica* showed the presence of alkaloids, coumarins, terpenoids, cardioglycosides, flavonoids, saponins, quinine and phenols as major chemical constituents.

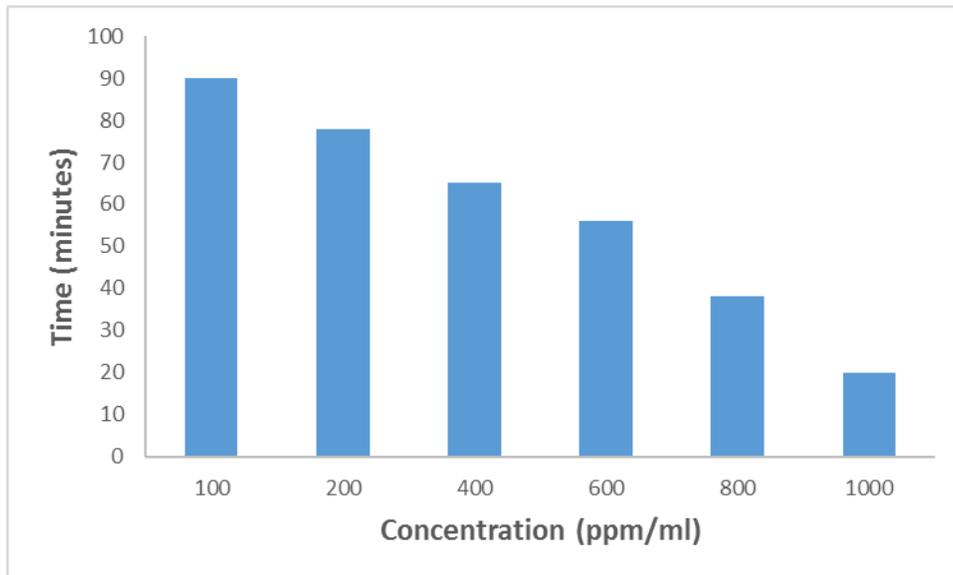
Among these, poly phenols, flavonoids are said to have anthelmintic activity (Nayak *et al.*, 2012). Therefore, it is assumed that the activity is contributed by the presence of the polyphenols and flavonoids. Similarly

saponins show Molluscicidal activity and the presence of saponins might have contributed the activity in the current study (Lemmich *et al.*, 1995).

**Fig.1** Anthelmintic activity of extract in varying concentration



**Fig.2** Molluscicidal activity of extract in varying concentration



From the results, it is evident that ethanolic fruit extract of *Momordica dioica* possess both anthelmintic and Molluscicidal properties in a dose dependent way. *M. dioica* extract kills the worm even in low concentrations within short duration of time compared to standard drug Albendazole. It justifies the claims of its potential anthelmintic and Molluscicidal activity. It can be also exploited as a neutrasuitical agent in our body as the frequent inclusion of this vegetable in our food, eliminate intestinal parasitic worms and purifies our blood by free radical scavenging activities.

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