In vitro anthelmintic activity of Cassava (Manihot esculenta) extract on Trichostongyloid larvae

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Abstract

Anthelmintic resistance has become a serious world-wide problem and has affected ruminant productivity in many countries. Thus the search for alternative methods of worm control has been an important priority in many small ruminant industries. Manihot esculenta had been shown to reduce trichostrongyloid eggs in infected sheep fed with such fresh leaves for prolonged period of time. Thus, this study was carried out to evaluate the anthelmintic activity of Cassava (Manihot esculenta) leaves extract against larvae of Trichostrongyloid nematodes using larval paralysis assay. Phytochemical analysis was carried out to determine the presence of tannin, saponin, phenols and alkaloids. The results of this study showed that Manihot esculenta extract can be used to control the infective stage of trichostrongyloid parasites of small ruminants.

Keywords: trichostrongyloid, larvae, Manihot esculenta (Cassava), phytochemical analysis.

Introduction

In Malaysia, trichostrongyloid worms showed a resistance to wide range of anthelmintics in sheep and goats (Dorny et al., 1994 and 1995; Rahman, 1994; Pandey and Sivaraj, 1994; Chandrawathani et al., 1999; Sivaraj et al., 1994). In recent years a lot of research had focused on using of traditional medical herbs to control these worms and most of them showed significant effects on the egg-hatching and larval development of Haemonchus contortus and other trichostrongyloids (Ademola et al., 2004, Al-Shaibani et al., 2009, Marie-Magdeleine et al., 2010 ).

Manihot esculenta (Cassava) and locally known as Ubi kayu is a perennial shrub of the Euphorbiaceae and native to tropical and subtropical regions. It is used in human and animal food because it is rich in carbohydrates and it is considered as one of the main source of starch together with rice and corn (Grace, 1977). Manihot esculenta hyad been shown to reduce gastrointestinal nematodes eggs in infected sheep fed with fresh leaves for prolonged time (López, 2007, Sokerya, 2009). The present research will study the effect of cassava leaf extracts against the infective larvae of trichostrongylids worms in infected goats.

Materials and Methods

(i) Plant collection and extracts preparation

Leaves of M. esculenta were collected in Bayan Lepas, Pulau Pinang, northwest Peninsular Malaysia. Voucher specimens were identified by botanists and deposited at the herbarium of School of Biological Sciences, Universiti Sains Malaysia, Penang and given a reference number 11182. The leaves were washed and dried at room temperature and the milled to powder using an electrical blender and stored in dark tightly closed glass bottles. The crude leaf extract were prepared with four organic solvents: hexan, chloroform, ethyl acetate and methanol (80%) using Soxhlet's apparatus. Solvents were evaporated using a rotary evaporator and the residue extracts were dried at room temperature and then stored at 4ºC in small tightly closed universal glass bottles. In vitro assays were conducted at concentrations of 3.1, 6.2, 12.5 and 25 mg/ml with three replicates for each concentration after dissolving the dried extract in Tween 20 to improve the solubility of extract in water (Maciel et al., 2006).
(ii) Larvae collection and identification

Faecal samples were collected from the rectum of naturally infected goats in a private farm in Bayan Lepas, Penang, northwest Peninsular Malaysia. Larvae were cultured according to MAFF (1986), and 100 larvae were identified according to MAFF (1986). Five replicates were carried out and the percentage of each species calculated.

(iii) In vitro assay

In vitro assays were carried out using Larval Paralysis Assay (LPA) according to the recommendations of the World Association for Advancement of Veterinary Parasitology (WAAVP) (Taylor et al., 2002; and Coles et al., 2006). Briefly, 100µl containing approximately 100 larvae were added to 100µl of extract in 96-well plate and kept at room temperature for 24 h. The living (motile) and dead larvae were counted. Two controls were used; one positive (Tween 20) and one negative (ivermectin 0.01 mg/ml).

(iv) Phytochemical study

Phytochemical tests were carried out to detect the presence of alkaloids, flavonoids, steroids, phenols and tannins in the methanol extract of M. esculenta leaves (Jack & Okorosaye-Orubite 2008; Egwaikhide & Gimba, 2007).

Statistical Analysis

The lethal concentration (LC\textsubscript{50}) of methanol extract was calculated from the linear regression ($y = ax + b$). Data from larval paralysis assay were compared using ANOVA and Tukey’s test (p<0.05 using SPSS).

Results and Discussion

The examination of fecal samples revealed that Haemonchus was the predominant genus (72 ± 2.20%), followed by Oesophagostomum (20.2 ± 1.00%), Trichostrongylus (6.2 ± 0.86%) and Ostertagia (1.6 ± 0.68%).

Table 1 shows the mean efficacy of M. esculenta leaf extracts on trichostrongyloid larvae using larval paralysis assay. Methanol extract had higher efficacy against the larvae than others (p<0.05); concentration of 25 mg/ml was more effective than other concentrations (p<0.05).

<table>
<thead>
<tr>
<th>Concentrations</th>
<th>M. esculenta leaves extracts</th>
<th>Hexan</th>
<th>Chloroform</th>
<th>Ethyl acetate</th>
<th>Methanol 80%</th>
</tr>
</thead>
<tbody>
<tr>
<td>25</td>
<td>10.33 ± 1.20</td>
<td>4.33 ± 2.33</td>
<td>5.00 ± 2.31</td>
<td>59.33 ± 8.96</td>
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<tr>
<td>12.5</td>
<td>2.33 ± 0.67</td>
<td>1.67 ± 0.88</td>
<td>1.33 ± 0.67</td>
<td>57.33 ± 6.89</td>
<td></td>
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<tr>
<td>6.2</td>
<td>0.67 ± 0.67</td>
<td>1.33 ± 0.33</td>
<td>0.33 ± 0.33</td>
<td>24.00 ± 3.60</td>
<td></td>
</tr>
<tr>
<td>3.1</td>
<td>0.33 ± 0.33</td>
<td>0.33 ± 0.33</td>
<td>0.00 ± 0.00</td>
<td>12.67 ± 3.18</td>
<td></td>
</tr>
<tr>
<td>1.6</td>
<td>0.00 ± 0.00</td>
<td>0.33 ± 0.33</td>
<td>0.33 ± 0.33</td>
<td>1.67 ± 0.33</td>
<td></td>
</tr>
<tr>
<td>Ivermectin 0.01</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>99.67 ± 0.33</td>
</tr>
<tr>
<td>Tween 20 (3%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.33 ± 0.40</td>
</tr>
</tbody>
</table>

The relationship between the concentration of methanol extract and the number of dead larvae was seen as linear despite a concentration of 25mg/ml. As result of this linear regression, it has been used to determine the equation that governs the relation. The equation is given as $y = 4.981x - 5.268$ with $R^2 = 0.993$. The effective concentration of methanol extract (LC\textsubscript{50}) that killed half of infective larvae was 11.1 mg/ml.
Phytochemical tests revealed the presence of alkaloids which were detected by producing of orange red precipitate after adding of Dragendorff’s reagent. The presence of tannins were detected by the dark green colour formed after adding of 2 drops of 5 % ferric acid solution whereas saponins were detected by using frothing test.

Results and Discussion

Cassava decreased in the number of nematodes eggs and Coccidia oocysts in the faeces (Tien Dung et al., 2005). However, there are no previous reports on the efficacy of the plant on larvae of trichostrongyliod nematodes in goats.

In the present research, 11mg/ml of methanol extract killed half of the larvae and that was less effective than the control because the control was a synthetic anthelmintic which included pure active substances, while the extract contained several compounds and the active compound may be present in small amounts.

Phytochemical tests revealed the presence of phenols, tannins, alkaloids, and saponins. These compounds revealed anthelmintic activity in some plants (Maciel et al., 2006, Costa et al., 2008, Olivirra et al., 2008). The activity of M. esculenta against trichostrongyliod larvae may differ from that of other plants like Melia azedarach (Maciel et al., 2006), Fumaria parviflora (Al-Shaibani et al., 2009), Azadirachta indica (Costa et al., 2008) and Tabernaemontana citrifolia (Marie-Magdeleine et al., 2010) and thus may differ in their respective active compounds.

Generally, the presence of condensed tannins in cassava extracts gives a good anthelmintic activity because it has direct and indirect effects against gastrointestinal nematodes (Athanasiadou et al., 2001, Iqbal et al., 2007). The mode of action of tannins as anthelmintic is attributed to their capacity to bind to some proteins of the metabolism or larva's organs and muscles causing a change in their functions and resulting in the paralysis or death (Martínez et al., 2010). The mechanism of saponin action can cause changes in the permeability of the cell membrane and pore formation, resulting in the disintegration of teguments of the parasites. Nevertheless, the mode of action of these organic substances still needs to be further investigated.

Conclusion

In this study, M. esculenta revealed anthelmintic activity due to the active substances in methanol leaf extract. However, further work should be carried out to isolate the active compounds responsible for anthelmintic activity and to understand their mode of action as well.

References


