Anthelmintic activity of *Cassia occidentalis* L. methanolic leaf extract on *Ascaridia galli* and *Heterakis gallinarum* and its acute toxicity

John N. Kateregga*, Maria Nabayunga, Patrick Vudriko, James G. Ndukui

INTRODUCTION

Poultry is a major source of high quality protein for the fast growing human population and provides income to resource-poor small-scale rural farmers; especially women.¹ Rural poultry require low levels of inputs and contribute significantly to food security and poverty alleviation. Chicken, which make up around 80% of poultry stocks in many developing countries of Africa and Asia, are also highly valued in the religious and socio-cultural lives of local communities and provide employment to many rural folk.²

Although global poultry meat production has increased in the last 10 years from 20% to 30%, infections such as helminthosis remain a major constraint to rural poultry production.³,⁴ Free ranging poultry production systems which predominate in rural communities expose birds to...
heavy worm burdens. The most frequently diagnosed nematodes of poultry are Acaridia galli and Heterakis gallinarum, with prevalences of 51.6% and 31% in chicken and turkeys respectively.5 Heterakis gallinarum is estimated to cause production losses in a range 10% to 20% due to impaired feed conversion, reduced egg production and increased mortality.6 This nematode also plays a role in histomoniasis (black head disease) epidemiology in turkeys where it causes severe disease and high mortality, sometimes up to 100% of a flock.7,8 Acaridia galli causes ascaridiosis with reduced feed conversion and mortality in birds.

The use of modern anthelmintics is not common among rural poultry farmers in developing countries because of high costs of drugs, their low availability and the high rates of illiteracy of the farmers. According to Uganda Bureau of Statistics (UBOS), 60.7% of rural people live below the poverty line and cannot afford conventional drugs.9 Furthermore, there is limited availability of drugs in rural areas despite the helminth burden prevalent in these areas due to the free range production system practiced there. In addition, drug residue problems and the development of resistance by target parasites have made alternative remedies imperative.10,11 Cassia occidentalis has for long been used by rural communities in Uganda against both human and domestic animal helminth infestations and, in the Luganda language spoken in central Uganda, it is known as “Omutanjoka” (‘worm killer’). Other studies indicate that the plant also possesses antibacterial, antiparasitic and antifungal effects.12 Exploitation of medicine plants provides alternative disease control options that are culturally acceptable; ecologically and environmentally sound.13 This study sought to evaluate the anthelmintic properties of Cassia occidentalis leaves as well as its acute toxicity in mice.

METHODS

Plant materials

Fresh leaves from healthy Cassia occidentalis plants free from plant pests were collected in Uganda from Nakabago village (0°24’ North 32°42’ East, VR 67, 34) and dried in the oven at 50°C for two days to a constant weight. A mature branch with flowers and fruits was picked from a plant and dried between plant presses under direct sunlight. The plant was identified, authenticated and a voucher specimen Maria Nabayunga No. 1 was deposited at the Botany Herbarium of Makerere University.

Preparation of the extract

The cold maceration method was used for extraction. Dried leaves were thoroughly crushed using motor and pestle and 350g of the powder was soaked in two liters of 70% methanol in a brown bottle at 25°C for three days and kept in a wooden cabinet. The mixture was shaken periodically and then filtered using cotton wool. The filtrate was poured in labeled kidney dishes of known weight and dried off in the oven at 50°C for two days, leaving a dark green residue which was weighed and placed in a glass bottle and kept at 4°C for further use. A stock solution of the extract was prepared immediately with distilled water before the start of the anthelmintic or acute toxicity evaluation experiments.

Worm collection

Acaridia galli worms were collected from the small intestines of freshly killed indigenous chicken from a chicken slaughter house in Bwaise market, Kampala, while Heterakis gallinarum worms were collected from the caeca of freshly slaughtered turkeys. Intestines and caeca were then dissected and worms isolated while placing them into a vacuum flask containing Goodwin’s physiological solution before being transported to the lab. They were then placed in two separate glass beakers containing Goodwin’s physiological solution and kep at 37°C immediately.

Evaluation of anthelmintic activity

The worms were placed in 250ml conical flasks containing 100ml of Goodwin’s solution at 37°C with each flask having 6 worms. For each worm species, 5 extract concentrations of 8, 12, 16, 20 and 24 mg/ml were used in duplicates. The negative control was Goodwin’s solution with no extract added. Piperazine citrate (20mg/ml) was used as positive control for Acaridia galli while ivermectin (0.5mg/ml) was used for Heterakis gallinarum. The extract concentrations were prepared from the stock solution using the formula C1V1 = C2V2 where C1 was the extract concentration; C2 was the stock concentration; V1 was the volume of Goodwin’s solution in each flask and V2 was the volume of the stock solution to be added to each flask.

The experimental setups for the two worm species were carried out separately in a water bath at 39°C for A. galli and at 41°C for H. gallinarum. The worms were then observed for mortality at time intervals of 12, 24, 36 and 48 hours and were recorded as dead when they did not express sinusoidal movement upon exposure of their extreme ends to a water temperature of 50°C. The number of dead and live worms for each flask were recorded and mean mortality ± SEM was calculated from the duplicates. Percent mortality for each concentration was determined and mean half-maximal effective concentration (EC50) obtained from a plot of percent mortality versus log concentration using GraphPad Prism. ANOVA was used for statistical comparison of mortality for the different extract concentrations. The differences were considered significant at p<0.05 or p<0.01.

Evaluation of the acute toxicity of the extract

Swiss mice aged 6-7 weeks old and weighing 20-29g were used for the acute toxicity study. They were fed on mice feed and water was provided ad libitum. Before the
experiment they were allowed to acclimatize for two weeks. The staircase (up and down) method was used to determine the different doses to be used for the different mice groups during the acute toxicity study. Two mice were randomly selected and orally administered with a limit dose of 5000mg/kg of the extract using an intragastric tube and observed for mortality after 24 hours. Basing on the results of the preliminary investigation, doses of 5,000, 10,000, 15,000, 20,000 and 25,000mg/Kg were used in the actual study.

Thirty (30) mice were placed in 5 groups of six mice each i.e. three females and three males with separate cages for each sex. The animals were fasted for 3 hours before the experiment and were weighed before dosing. They were then marked with unique marks on the tails using a water proof marker for proper identification and follow up during the experiment. Group I was used as a negative control and no extract was administered to this group. Group II to IV were orally administered with the extract using an intragastric tube at doses of 5,000, 10,000, 15,000, 20,000 and 25,000mg/Kg respectively. The volume of stock solution for oral dosing was calculated using the formula

\[ V = \frac{D \times W}{C} \]

where \( V \) (ml) is volume required from the stock solution; \( D \) (mg/kg) is the dose; \( W \) (kg) is mouse body weight and \( C \) (mg/ml) is stock solution concentration. The number of dead animals per group after 24 hours was recorded. Determination of the LD50 was done by plotting mortality probits against log dose. Ethical clearance was obtained from the Research and Ethics Committee of the College of Veterinary Medicine, Makerere University.

RESULTS

The ascaricidal activity of the \( C. \) occidentalis methanolic extract increased with incubation time (Figure 1). After 12 hours, only the positive control showed mortality. The 8, 12, 16, 20, and 24mg/ml concentrations showed mortality after 24 hours. The negative control did not show any mortality up to 48 hours. At 48 hours the 24mg/ml concentration showed a significantly higher mortality (p<0.01) than the positive control (piperazine citrate) when used at the recommended concentration of 2mg/ml (Table 1). No mortality was observed for the negative control.

The anti-\( Heterakis gallinarum \) activity of the extract increased with duration of incubation time (Figure 2). Only the 16 and 24mg/ml extract concentrations and positive control showed mortality after 12 hours. The 24mg/ml showed higher mortality than the positive control (Table 2) but this was not statistically significant (p>0.05). No mortality was observed for the negative control.

The \( C. \) occidentalis methanolic extract showed a concentration-dependent relationship on both \( A. \) galli and \( H. \) gallinarum nematodes.

Table 1: Ascaridia galli mortality at 48h for the different treatments.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Mean mortality ± SEM</th>
<th>% mortality</th>
</tr>
</thead>
<tbody>
<tr>
<td>Extract 8 mg/ml</td>
<td>2.00± 0.91</td>
<td>33.3</td>
</tr>
<tr>
<td>Extract 12 mg/ml</td>
<td>2.63±1.14</td>
<td>50.0</td>
</tr>
<tr>
<td>Extract 16 mg/ml</td>
<td>2.75±1.00</td>
<td>66.7</td>
</tr>
<tr>
<td>Extract 20 mg/ml</td>
<td>3.00±1.34</td>
<td>66.7</td>
</tr>
<tr>
<td>Extract 24 mg/ml</td>
<td>3.50±1.24</td>
<td>83.3</td>
</tr>
<tr>
<td>Piperazine 2mg/ml</td>
<td>2.25±0.52</td>
<td>33.3</td>
</tr>
<tr>
<td>Goodwin’s solution</td>
<td>0.0±0.0</td>
<td>0.0</td>
</tr>
</tbody>
</table>

n=6, *p<0.05, **p<0.01(comparison of concentrations with positive control)

Figure 1: Effect of \( Cassia occidentalis \) methanolic leaf extract on \( Ascaridia galli \).

The worms were exposed to various concentrations of the \( C. \) occidentalis methanolic extract for a period of 48 hours. The extract showed a concentration and time-dependent wormicidal activity.

Figure 2: Effect of \( Cassia occidentalis \) on \( Heterakis gallinarum \).

The worms were exposed to various concentrations of the test extract. The extract showed a concentration and time-dependent wormicidal activity.

However, the mortality was higher for \( A. \) galli than for \( H. \) gallinarum especially at the higher doses (Figure 3). The
R (regression co-efficient) for *A. galli* was 0.8891 while that for *H. gallinarum* was 0.7814 indicating a stronger relationship between dose and response for *A. galli*. The EC\textsubscript{50} for the *A. galli* was determined to be 11.78mg/ml while that for *H. gallinarum* was 17.75mg/ml.

Table 2: *Heterakis gallinarum* mortality at 48h for the different treatments.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Mean mortality ± SEM</th>
<th>% mortality</th>
</tr>
</thead>
<tbody>
<tr>
<td>Extract 8 mg/ml</td>
<td>2.13±1.11</td>
<td>33.3</td>
</tr>
<tr>
<td>Extract 12 mg/ml</td>
<td>2.25±1.05</td>
<td>33.3</td>
</tr>
<tr>
<td>Extract 16 mg/ml</td>
<td>2.50±1.06</td>
<td>50.0</td>
</tr>
<tr>
<td>Extract 20 mg/ml</td>
<td>2.63±1.197</td>
<td>50.0</td>
</tr>
<tr>
<td>Extract 24 mg/ml</td>
<td>3.75±1.20</td>
<td>66.7</td>
</tr>
<tr>
<td>Ivermectin 0.5mg/ml (Positive control)</td>
<td>2.13±0.75</td>
<td>33.3</td>
</tr>
<tr>
<td>Goodwin’s solution (Negative control)</td>
<td>0.0±0.0</td>
<td>0.0</td>
</tr>
</tbody>
</table>

n=6, p>0.05 (comparison of concentrations and positive control)

Figure 3: Concentration-response relationship for *A. galli* and *H. gallinarum*.

The EC\textsubscript{50} of the extract for *A. galli* and *H. gallinarum* is the anti-log of the value on the x axis corresponding to 50% mortality.

**DISCUSSION**

Rural poultry farmers in Uganda use both modern anthelmintics and medicinal plants to control *Ascaridia galli* and *Heterakis gallinarum* infestations. Piperazine, the standard drug for *A. galli* acts by increasing chloride ion conductance leading to hyperpolarisation of the worm muscles leading to paralysis and death of the worm.\(^{17}\) Ivermectin, the drug mainly used for *H. gallinarum* binds selectively to glutamate-gated chloride ion channels in worm nerve and muscle cells causing hyperpolarization of the cells and paralysis and death of the worms. It is believed to act as an agonist at the receptor of the inhibitory neurotransmitter GABA, disrupting GABA-mediated central nervous system neurosynaptic transmission.\(^{18}\)

This study demonstrated that *C. occidentalis* extracts have incubation time and concentration-dependent anthelmintic activity against both worms. The dose-dependent relationship could be explained by exposure of the worms to higher quantities of the ascaricidal components of the plant extract leading to higher mortalities. The increase in activity with time may be due to a higher number of receptor sites being occupied by the active ingredients of the extract with increase in incubation time.\(^{29}\) An *in vitro* study with the ethanolic leaf extract of *C. occidentalis* show ascaricidal activity against *A. suum* another worm of the Ascarididae family.\(^{30}\)

The nematocidal activity of the plant could be attributed to phytochemicals in the leaves. The main phytochemicals in *C. occidentalis* leaf are carbohydrates, saponins, flavonoids, terpenes, sterols, alkaloids, anthraquinones and tannins.\(^{21}\) The methanolic leaf extract of this plant has been shown to have high quantities of tannins, saponins and alkaloids;\(^{22}\) phytochemicals with demonstrated anthelmintic properties.\(^{23}\) Tannins are water soluble polyphenolic compounds which interfere with energy generation in helminths by uncoupling oxidative phosphorylation just like synthetic phenolic anthelmintics like niclosamide, oxyclozamide, nitroxynil and bithionol and could have been majorly responsible for the activity observed in this study.\(^{25}\) They also bind free proteins in the gastrointestinal tract of the host or glycoprotein on the cuticle of the parasite disrupting the physiological functions such as motility, feed absorption and reproduction.\(^{25,26}\) They also cause death by interfering with the morphology and disrupt the proteolytic activity of microbes.\(^{27}\)

Alkaloids could have contributed to the paralysis and consequent death of the worms. The nematocidal activity of alkaloids has been demonstrated in two rat nematodes; *Strongyloides ratti* and *Strongyloides venezuelensis*.\(^{28}\) Alkaloid salts are competitive antagonists at muscarinic cholinergergic receptor preventing the binding of acetylcholine and their activity is similar to that of some anthelmintics such as pyrantel and morantel.\(^{29}\) Saponins are known to cause feed refusal and starvation of the parasites leading to their death from lack of energy. Such metabolites may work singly or in combination to impair worm motility (paralysis) and or cause death of the helminths. The synergistic interactions of the phytochemicals have been shown to be more effective than activity of individual constituents.\(^{30}\) Plant metabolites’ action may be additive, synergic, or antagonistic as they act at single or multiple target sites of the worms.\(^{31}\)

The acute toxicity study showed no mortality up to the extract dose level of 25,000mg/kg body weight. This indicates a high safety level since; according to OECD acute toxicity guidelines 423, doses higher than 5,000 mg/Kg are assumed to be experimentally safe.\(^{32}\) This study has demonstrated the anthelmintic activity of *C. occidentalis* and this could explain its use in traditional medicine as a remedy against helminth infections over the generations in many parts of Uganda. There is however
need to purify the crude extract further in order to identify the active chemicals responsible for the anthelmintic activity.

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**Conflict of interest:** None declared

**Ethical approval:** Ethical clearance was obtained from the Research and Ethics Committee of the College of Veterinary Medicine, Makerere University

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**REFERENCES**


