

NOVEL APPLICATION OF *TRICHILIA HEUDELOTII* PLANCH: EFFECTIVENESS OF DIFFERENT POLARITY OF ORGANIC SOLVENTS OF LEAF AND STEM BARK EXTRACTS ON THE CONTROL OF COWPEA BEETLE

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ABSTRACT. *Callosobruchus maculatus* (cowpea beetle) is a field-to-store pest causing over 90% losses of cowpea. The search for new plant-derived crop protectant that can be explored as alternative to synthetic pesticides is urgently needed. Thus, the phytochemical screening, as well as the bioactivity of different polarity of organic solvents of leaf and stem bark extracts of *Trichilia heudelotii* (Meliaceae), was investigated against the field-to-store insect pest *Callosobruchus maculatus* in laboratory bioassay. The non-polar (hexane) and polar (ethanol) extracts [0.0 (control), 0.5, 1.0 and 1.5 ml] of the leaf and stem bark were added to 100 g of cowpea. The following parameters of the cowpea weevil life cycle were analysed at the

various concentrations: adult mortality, oviposition rate, number of larvae and pupae, and emergence of first filial progeny. The phytochemical screening revealed the presence of flavonoids, terpenoids, alkaloids, saponins in both the non-polar and polar extracts. Furthermore, the various treatments used were significantly ($p < 0.05$) effective in controlling *C. maculatus* from 1 and also 30 days after treatment (DAT) for both the non-polar and polar extracts when compared to the control. However, the most effective treatment was observed with the polar (EtOH) stem bark extract at 1.5 ml, which caused more mortalities and fewer emergence of the insect. Our findings suggest that the non-polar and polar leaf and stem bark of *T. heudelotii*

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NOVEL APPLICATION OF TRICHILIA HEUDELOTII ON THE CONTROL OF COWPEA BEETLE

extracts could serve as a sustainable and potential alternative to synthetic chemicals in pest control.

Keywords: *Callosobruchus maculatus*; *Trichilia heudelotii* extracts; bio-pesticide; insect pest control.

INTRODUCTION

The crop cowpea (*Vigna unguiculata* (L.) Walp.) is an important food and forage legume grown in the semi-arid tropics, which include parts of Asia, Africa, Southern United States, Central and South America and Southern Europe (Singh, 2005; Timko *et al.*, 2007a). It is a multifunctional crop, providing nutrition for both man and livestock and serving as a dependable and liable revenue-generating commodity for traders and farmers (Singh, 2002; Langyintuo *et al.*, 2003). The crop is attacked by many pests, which includes the cowpea weevil *Callosobruchus maculatus*.

The bruchid beetle, *Callosobruchus maculatus* (Fabricus) of the Chrysomelidae family, is a reddish-brown insect with black and gray elytra, marked with two central black spots (Fox and Reed, 2011). They are usually sexually dimorphic and the sexes are easily distinguished from each other. Females are sometimes larger and darker than males (Beck and Blumer, 2014) and they exist in two morphs; the flightless and the flying form. The flying form is more common in beetles that developed in conditions of high temperature like the tropics (Umar and Turaki, 2014). A female

C. maculatus has been reported to lay more than a hundred eggs on the surface of cowpea with most of the eggs viable and hatching. Larvae of the insect emerges after four to eight days and then burrow themselves into the seeds (Umar and Turaki, 2014). The larva proceeds to feed on the seed content usually feeding on the embryo and then leaving tiny holes through, which it will exit when it matures to adult after three to seven weeks depending on the environmental situations. The higher the temperature the quicker the rate of development. The adult period ranges from 9-14 days (Beck and Blummer, 2014).

Callosobruchus maculatus is a field-to-store insect pest. Infestation begins from the field and damage continues in storage (Ali *et al.*, 2004; Swella and Mushobozy, 2007). Estimates of storage losses have been reported from 4 - 90% due to punctures caused by the beetles, thus reducing the degree of usefulness and rendering seeds unfit for planting or consumption (Ali *et al.*, 2004; Umeozor, 2005). Kéïta *et al.* (2000) also reported that *C. maculatus* can cause weight loss by up to 60% when cowpea seeds are stored without any protection.

Plants are known to have evolved millions of years ago to defend themselves from external attacks by pests, in so doing, they develop natural repellent mechanisms that are antagonistic to a varying degree of pests. In addition, as a result of the problems associated with pesticide resistance, negative effects

on non-target organisms, man and the environment, the idea of using plant materials as natural (botanical) insecticides to protect crops from damages incurred by insect pest is a safer alternative in pest control, when compared to the hazardous effects of synthetic chemicals and this is gaining much prominence among researchers and household users (Lajide *et al.*, 2003). From ancient times, these plant materials have been noticed and used as natural insecticides in stored grains by mixing grains with leaf, bark, seed powder or oil extracts of plants to suppress insect infestation and developments in grains (Addor, 1995; Talukder *et al.*, 1995). Thus, these botanical insecticides are naturally occurring biochemicals that are broken down easily and readily unlike synthetic chemical pesticides (Egho, 2011).

In this study, *Trichilia heudelotii* (Meliaceae) leaves and stem bark were used against the bruchid beetle *Callosobruchus maculatus*. The plant has gained the attention of many phytochemists interested in bioproduction due to its diverse classes of compounds identified as well as insecticidal actions against several insect pests (Vieira *et al.*, 2013; Bankole *et al.*, 2016). Over the years, phytochemical investigation had revealed that the genus *Trichilia* is rich in various classes of compounds, which includes terpenoids, steroids, flavonoids and other terpenes by-products (Ramirez *et al.*, 2000; Rodrigues *et al.*, 2010; Vieira *et al.*, 2014). Furthermore, a wide range of

research had been reported on various extracts of the different species from the genus *Trichilia* revealing insecticidal activities on different groups of insect pests (Xie *et al.*, 1994; Ayo *et al.*, 2013; Freitas *et al.*, 2014). Previously, there has been various reports of other species of the genus *Trichilia* used in the control of various insect pests (López-Olegrín, 1998; Isman, 2001; Sengottayan, 2013; Curcino-Vieira *et al.*, 2014; Garcia-Gomez *et al.*, 2019); however the species *Trichilia heudelotii* is yet to be evaluated for its insecticidal potential. Thus, this prompted us to investigate the effectiveness of *Trichilia heudelotii* leaf and stem bark non-polar and polar extracts as natural insecticides in the control of *Callosobruchus maculatus* infesting cowpea, which to the best of our knowledge has not been studied previously and to carryout phytochemical screening on the extracts.

MATERIALS AND METHODS

Sample collection and identification

Fresh leaves and stem bark of *Trichilia heudelotii* were collected from a thick forest in Oyan town, Osun State, Nigeria. The plant was authenticated at the Ward Herbarium of the University of Ilorin, and was assigned a voucher number, UILH/001/1367.

Sample preparation and Extraction

The *Trichilia heudelotii* leaves and stem bark were air-dried at room temperature. The dried leaves and stem bark were ground using an electric grinder

NOVEL APPLICATION OF TRICHILIA HEUDELII ON THE CONTROL OF COWPEA BEETLE

and sieved using a 90-micron mesh sieve to obtain fine powder of the samples.

The powdered leaves (650 g) and stem bark (1.1 kg) were extracted twice by maceration, sequentially with a non-polar and polar organic solvents, hexane (Hex) and ethanol (EtOH), respectively. Yields of 14.26 g (Hex) and 10.24 g (EtOH) of the leaves was obtained after evaporation of the organic solvents. For the stem bark, 9.26g (Hex) and 13.59 g (EtOH) was obtained.

The crude extracts obtained used for the experiment were then prepared using an aqueous solution of 1% v/v of acetic acid in distilled water and concentrated at 0.5 g per ml.

Qualitative phytochemical screening

Test for tannins. Few drops of 1% lead acetate was added to 0.2 g of the extract and observed for the formation of yellow precipitate (Trease and Evans, 1989).

Test for alkaloids. Exactly 0.2 g of the extract was stirred with 5 ml of 1% aqueous HCl on water bath and then filtered. One ml of the filtrate was taken individually into two separate test tubes. To the first portion, Mayer's reagent was added and appearance of buff-colored precipitate was an indication for the presence of alkaloids. To the second portion, few drops of Dragendorff's reagent was added to the filtrate and observed for the formation of an orange-red precipitate (Sofowora, 1993).

Test for terpenoids (Salkowski test). Exactly 2 ml of chloroform was added to 0.2 g of the extract, 3 ml of concentrated sulphuric acid was added carefully to form a layer. Formation of a reddish-brown colouration at the interface indicates the presence of terpenoids (Sofowora, 1993).

Test for phenolic compounds. The extract (0.5 g) was dissolved in 5 ml of

distilled water. To this, few drops of neutral 5% ferric chloride solution was added. A dark green colour indicates the presence of phenolic compounds (Sofowora, 1993).

Test for flavonoids. Exactly 4 ml of dilute ammonia solution was added to a portion of the extract, followed by addition of concentrated sulphuric acid. A yellow colouration indicates the presence of flavonoids (Sofowora, 1993).

Test for saponins. Exactly 1 g of the extract was boiled with 5 ml of distilled water and filtered. To the filtrate, about 3 ml of distilled water was further added and shaken vigorously for about 5 min. Frothing which persists on warming shows the presence of saponins (Trease and Evans, 1989).

Test for anthraquinones. Few drops of dilute sulphuric acid were added to 0.2 g of the extract. This was boiled and filtered. The filtrate was extracted with 2 ml of chloroform. The chloroform layer was then treated with 1 ml of ammonia. The formation of red colour on the ammoniacal layer shows the presence of anthraquinones (Sofowora, 1993; Harborne, 1973).

Test for steroids. Acetic anhydride (2 ml) was added to a portion of the extract with 2ml H₂SO₄. Colour change from violet to blue or green indicates the presence of steroids (Sofowora, 1993).

Test for anthocyanins. Exactly 3 ml of 2M HCl and ammonia solution were added to a portion of the extract. The appearance of pink-red colouration which turns blue-violet indicates the presence of anthocyanins (Savithramma *et al.*, 2011).

Insect culture

The method of Ousman *et al.* (2007) was adopted in rearing of *C. maculatus*. The insects were obtained from an existing culture from Nigerian Stored Products Research Institute (NSPRI), Ilorin,

Nigeria. Eight pairs of active male and female *C. maculatus* were introduced into transparent plastic containers (7 × 8 cm) containing untreated cowpea seeds. Next, they were allowed to mate and oviposit for seven days. The plastic containers were covered with muslin cloth held in place by a rubber band to prevent escape of the insect and allow for ventilation. After one week, the introduced *C. maculatus* were removed and the containers were left under laboratory conditions (temperature 24° - 28°C, relative humidity of 70%) till the emergence of F₁ progeny. The F₁ progenies from the cultures were used for the experiment.

Source and type of cowpea seeds used

The cowpea variety IT96D-610 was obtained from the International Institute for Tropical Agriculture (IITA), Ibadan, Nigeria. The variety is characterized by having a smooth seed coat, high root biomass and also drought-resistant. Clean and undamaged seeds were picked and used for the experiment. The cowpea seeds were stored in the freezer compartment of a refrigerator at -17°C for four days to incapacitate any existing insects before experimenting.

Experimental procedure and design

100 g of clean and undamaged cowpea seeds were placed in transparent plastic containers (7 × 8 cm). The seeds were properly mixed with the various levels of the botanical extracts (0.5 ml, 1.0 ml, and 1.5 ml) of each treatment using a wooden spatula to ensure uniform coating of the seeds. The seeds were air-dried for 5 min. before introducing four pairs of active male and female *C. maculatus* (24 hrs old) into the treatments. The containers were covered with muslin cloth tied with rubber band to allow for ventilation and prevent escape of *C. maculatus*.

The experiment was arranged in a Completely Randomized Design (CRD) with five treatments (hex leaf; hex stem bark extracts [non-polar extract] and EtOH leaf; EtOH bark extracts [polar extract] and control) at three levels (0.5 ml, 1.0 ml and 1.5 ml) and each level was replicated three times plus the control. The experiment was carried out in two batches, which was dependent on the period of introduction of *C. maculatus* into the experimental units. Introduction of *C. maculatus* into the experimental units was done immediately after seed treatment for the first batch, while the second batch was infested 30 days after seed treatment to test for the strength of shelf life of the plant extracts. All the experimental units were treated at the same time. The experiment was carried out under ambient laboratory conditions.

Data collection

Data were collected on the following parameters:

Adult mortality. This was taken for five consecutive days. The insects were confirmed dead when there was no response to probing with an entomological needle at the abdomen, leg or antennae. The numbers of dead *C. maculatus* were counted daily and recorded.

Oviposition rate. This was taken at seven days after infestation. Ten seeds were randomly selected and seeds were viewed for the presence of eggs using a × 100 magnifying lens.

Number of larvae and pupae. For the presence of larvae and pupae, four seeds were selected at random and seeds were carefully dissected using a sharp surgical knife to expose larvae and pupae in seeds. The numbers of larvae were taken 13 days after treatment while the number of pupae was recorded 18 days after treatment.

NOVEL APPLICATION OF TRICHILIA HEUDELOTII ON THE CONTROL OF COWPEA BEETLE

Emergence of F_1 progeny. At 28 days after treatment, observation was taken for the emergence of new progeny of the weevil by counting newly emerged adults found in each of the treatments.

Statistical analysis

Data were subjected to two-way analysis of variance (ANOVA). Where treatment means were significant, multiple comparisons of treatments was done using the new Duncan's Multiple Range Test (n DMRT) at 5% level of significance. All statistical analysis was done using statistical package of social science (SPSS) version 20.

RESULTS

Effect of *Trichilia heudelotii* extracts on adult mortality of *Callosobruchus maculatus* after treatment

Tables 1 and 2 show the effects of the different extracts (non-polar and polar) of *Trichilia heudelotii* on the mortality of adults of *C. maculatus* from 1 and 31 days after treatment (DAT). Post hoc analysis revealed that there were significant differences between the treatment of the extracts of *T. heudelotii* and the control at 1 and 2 DAT (Table 1). The polar EtOH extract (Table 1) of *T. heudelotii* stem bark had a significantly ($p < 0.05$) higher rate of adult mortality of *C. maculatus* at the concentrations of 1.5 ($M = 7.00 \pm 1.00$) and 1.0 ml ($M = 6.33 \pm 1.528$), respectively, at 1 DAT. It was also observed that at 1 DAT, all the treatment extracts of *T. heudelotii* were effective in significantly ($p < 0.05$) increasing the number of adult mortality, when compared to the control, which only had a mortality

rate of 2.33 (± 0.577) (Table 1). At 2 DAT, the EtOH leaf extract (polar extract) was significantly ($p < 0.05$) the highest in causing more adult mortality of *C. maculatus* at 1.0 ml ($M = 5.33 \pm 0.577$), when compared to the other extracts and the control (Table 1).

In addition, observations on the shelf life of the treatments revealed that the EtOH stem bark (polar) extract of *T. heudelotii* at the concentration rate of 1.5 ml was significantly ($p < 0.05$) able to suppress the population of *C. maculatus* adults ($M = 7.00$), followed by 1.0 ml ($M = 6.33$) and 0.5 ml ($M = 6.00$), *i.e.* in a dose dependent manner, respectively, when compared to the other treatment extracts, which were lower at 31 DAT, as shown in Table 2.

Effect of *Trichilia heudelotii* extracts on oviposition rate of *Callosobruchus maculatus* 7 and 37 days after treatment

The effects of the extracts at varying concentrations of *T. heudelotii* on the oviposition rate of *C. maculatus* from 7 and 37 days after treatment (DAT) is shown in Table 3. There were significant differences between the non-polar and polar leaf and stem bark extracts and the control.

All the extracts were significantly able to control the oviposition rate of *C. maculatus* at 7 and 37 DAT however, the EtOH stem bark (polar) extract had the lowest mean number of oviposition ($M = 1.33$) at the concentration rate of 1.5 ml,

which was significantly ($p < 0.5$) able to suppress the oviposition rate of *C. maculatus*. This was also observed at 37 DAT, showing the least mean number (2.00) of oviposition, when

compared to the control (Table 3). During the cause of the experiment, the control did not affect the rate of oviposition of *C. maculatus*.

Table 1 - Effect of non-polar and polar leaf and stem bark extracts of *Trichilia heudelotii* on adult mortality in the first 5 days after treatment against *Callosobruchus maculatus*

Treatment	Conc. (ml) /100 g cowpea	Days after treatment (DAT)				
		1	2	3	4	5
Non-polar extracts						
HTL	0.5	4.00±2.00 ^{abcdef}	1.67±0.577 ^{abcd}	1.67±1.528 ^a	0.67±0.577 ^{ab}	1.33±0.577 ^{abcd}
	1.0	5.00±0.000 ^{cdefgh}	1.33±0.577 ^{abcd}	1.33±0.577 ^{ab}	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.5	5.33±0.577 ^{defgh}	1.67±0.577 ^{abcd}	1.00±1.000 ^{ab}	0.00±0.000 ^a	0.33±0.577 ^a
HTB	0.5	5.33±0.577 ^{defgh}	1.33±0.577 ^{abcd}	0.67±0.577 ^a	0.67±1.155 ^{ab}	0.33±0.577 ^a
	1.0	5.67±1.155 ^{efgh}	1.33±1.155 ^{abcd}	1.00±0.000 ^a	0.00±0.000 ^a	0.33±0.577 ^a
	1.5	6.00±1.000 ^{gh}	1.33±0.577 ^{abcd}	0.67±0.577 ^a	0.00±0.000 ^a	0.67±0.577 ^{ab}
Polar extracts						
ETL	0.5	4.67±1.155 ^{bcddefg}	1.67±1.155 ^{abc}	0.67±0.577 ^a	0.67±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.0	5.33±0.577 ^{defgh}	5.33±0.577 ^e	0.67±0.577 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.5	6.00±1.000 ^{gh}	0.67±0.577 ^{ab}	0.67±0.577 ^a	0.67±0.577 ^{ab}	0.33±0.577 ^a
ETB	0.5	6.00±1.000 ^{gh}	1.00±1.000 ^{abc}	0.33±0.577 ^a	0.67±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.0	6.33±1.528 ^{gh}	1.00±1.732 ^{abc}	0.33±0.577 ^a	0.67±0.577 ^{ab}	0.33±0.577 ^a
	1.5	7.00±1.000 ^h	0.33±0.577 ^a	0.33±0.577 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
Control	0.0	2.33±0.577 ^a	1.00±1.000 ^{abc}	1.00±1.000 ^{ab}	1.67±2.082 ^b	2.33±0.577 ^d

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stem bark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stem bark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Table 2 - Effect of non-polar and polar leaf and stem bark extracts of *Trichilia heudelotii* on adult mortality 30 days after treatment

Treatment	Conc. (ml)/ 100 g cowpea	Days after treatment (DAT)				
		31	32	33	34	35
Non-polar extracts						
HTL	0.5	3.00±1.732 ^{abc}	2.33±0.577 ^{bcd}	1.33±0.577 ^a	0.67±0.577 ^{ab}	1.33±0.577 ^{abcd}
	1.0	3.33±0.577 ^{abcd}	2.33±0.577 ^{bcd}	1.33±0.577 ^a	0.67±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.5	3.00±1.000 ^{abc}	3.00±1.000 ^d	1.00±1.000 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
HTB	0.5	3.33±1.528 ^{abcd}	2.67±0.577 ^{cd}	1.33±1.155 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.0	3.67±1.155 ^{abcde}	3.00±1.000 ^d	0.67±0.577 ^a	0.00±0.000 ^a	0.33±0.577 ^a
	1.5	4.33±1.528 ^{abcdefg}	3.00±1.732 ^d	0.33±0.577 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
Polar extracts						
ETL	0.5	2.33±0.577 ^a	2.33±1.155 ^{bcd}	1.33±0.577 ^a	0.33±0.577 ^{ab}	1.33±0.577 ^{abcd}
	1.0	3.33±0.577 ^{abcd}	2.33±0.577 ^{bcd}	1.33±0.577 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.5	3.00±0.000 ^{abc}	3.00±1.732 ^d	1.00±1.000 ^a	0.67±0.577 ^{ab}	1.00±0.000 ^{abc}
ETB	0.5	6.00±1.000 ^{fgh}	2.67±1.155 ^{cd}	1.00±0.000 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.0	6.33±1.528 ^{gh}	3.00±1.000 ^d	0.67±0.577 ^a	0.67±0.577 ^{ab}	0.33±0.577 ^a
	1.5ml	7.00±1.000 ^h	3.00±1.000 ^d	0.67±0.577 ^a	0.33±0.577 ^{ab}	0.33±0.577 ^a
Control	0.0	2.33±0.577 ^a	1.00±1.000 ^{abc}	1.00±1.000 ^{ab}	1.67±2.082 ^b	2.33±0.577 ^d

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stem bark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stem bark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Table 3 - Effect of non-polar and polar leaf and stem bark extracts of *Trichilia heudelotii* on oviposition rate

Treatment	Conc. (ml)/100 g cowpea	Days after treatment (DAT)	
		7	37
Non-polar extracts			
HTL	0.5	4.00±2.00 ^{bcd}	4.33±1.528 ^{cde}
	1.0	3.00±2.646 ^{abcde}	4.00±1.000 ^{bcd}
	1.5	3.33±1.528 ^{abcde}	3.33±1.155 ^{abcde}
HTB	0.5	2.33±0.577 ^{abcd}	2.67±0.577 ^{abcde}
	1.0	1.67±0.577 ^{ab}	2.33±0.577 ^{abcd}
	1.5	2.33±0.577 ^{abcd}	2.67±0.577 ^{abcde}
Polar extracts			
ETL	0.5	4.00±1.000 ^{bcd}	4.33±1.155 ^{cde}
	1.0	3.33±1.000 ^{bcd}	3.33±0.577 ^{abcde}
	1.5	3.00±1.000 ^{abcde}	3.00±1.000 ^{abcde}
ETB	0.5	2.33±0.577 ^{abcd}	2.67±0.577 ^{abcde}
	1.0	1.67±1.155 ^{ab}	2.67±1.155 ^{abcde}
	1.5	1.33±0.577 ^a	2.00±0.000 ^{abc}
Control	0.0	8.33±1.155 ^f	8.33±1.528 ^f

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test.

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stem bark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stem bark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Effect of *Trichilia heudelotii* extracts on larva emergence of *Callosobruchus maculatus* at 1 and 30 days after treatment

The larva emergence of *Callosobruchus maculatus* reared on *Trichilia heudelotii* non-polar and polar leaf and stem bark extracts at 1 and 30 days after treatment (DAT) is shown in *Table 4*. There was a significant difference between the treatments and control with all the treatments showing a significant level of reduction in the population of *C. maculatus* larvae, when compared to the control. At the highest concentration (1.5 ml) and at 1 DAT, the non-polar (hex) leaf extract, EtOH

leaf and stem bark (polar) extract of *T. heudelotii* had the least mean number (0.67) of larva emergence of *C. maculatus*.

While at 30 DAT, only the non-polar (hex) stem bark extract had the least mean number (0.67) of larva emergence at the highest concentration (1.5 ml) used and was significantly ($p < 0.05$) able to halt the emergence of *C. maculatus*, when compared to the control. The control ($M = 6.00$) did not affect the emergence of *C. maculatus* larvae throughout the experiment (*Table 4*).

NOVEL APPLICATION OF *TRICHILIA HEUDELOTII* ON THE CONTROL OF COWPEA BEETLE

Table 4 - Effect of *Trichilia heudelotii* non-polar and polar leaf and stem bark extracts on larva emergence of *Callosobruchus maculatus*

Treatment	Conc. (ml)/100g cowpea	Day after treatment (DAT)	
		1	30
Non-polar extracts			
HTL	0.5	2.00±1.00 ^{ab}	2.67±1.155 ^{ab}
	1.0	1.33±1.528 ^{ab}	1.67±0.577 ^{ab}
	1.5	0.67±1.155 ^a	1.67±1.155 ^{ab}
HTB	0.5	1.33±1.155 ^{ab}	2.00±1.00 ^{ab}
	1.0	1.33±0.577 ^{ab}	1.33±0.577 ^{ab}
	1.5	1.00±1.000 ^{ab}	0.67±0.577 ^a
Polar extracts			
ETL	0.5	1.67±0.677 ^{ab}	2.00±0.000 ^{ab}
	1.0	1.33±0.577 ^{ab}	1.33±0.577 ^{ab}
	1.5	0.67±0.577 ^a	1.00±0.000 ^{ab}
ETB	0.5	1.33±0.577 ^{ab}	2.00±1.000 ^{ab}
	1.0	1.00±1.00 ^{ab}	1.67±1.15 ^{ab}
	1.5	0.67±0.577 ^a	1.67±0.577 ^{ab}
Control	0.0	6.00±1.000 ^c	6.00±1.73 ^c

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test.

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stembark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stembark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Effect of *Trichilia heudelotii* non-polar and polar extracts on pupae emergence of *Callosobruchus maculatus* at 1 and 30 days after treatment

Table 5 shows the effects of the treatment extracts on the emergence of *C. maculatus* pupae. A significant difference was observed between the treatment extracts and control. At concentration of 1.0 and 1.5 ml and at 1 DAT, the non-polar (hex) stem bark extract and polar (EtOH) leaf extract of *T. heudelotii* had the lowest mean number (M = 0.67) of *C. maculatus* pupae while the lowest mean number (M = 0.67) of *C. maculatus* pupae was also observed for the polar (EtOH)

stem bark extract at only 1.0 ml concentration on the same 1 DAT which was significantly ($p < 0.05$) lower, when compared to the control (M = 4.67), which had no reduction in the number of pupae emerging.

The various treatments showed a significant ($p < 0.05$) level of control of *C. maculatus* pupae at 30 DAT (Table 5). However, the non-polar (hex) bark extract and EtOH (polar) leaf extract of *T. heudelotii* gave the lowest mean numbers (M = 0.67) of pupae emergence at the highest concentration level (1.5 ml) used. The control showed no reduction in the number of pupae emergence of *C. maculatus*.

Table 5 - Effect of *T. heudelotii* leaf and stem bark (non-polar and polar) extracts on pupae emergence of *Callosobruchus maculatus* at different concentrations

Treatment	Conc. (ml)/100g cowpea	Day after treatment (DAT)	
		1	30
Non-polar extracts			
HTL	0.5	1.67±1.155 ^{ab}	2.00±1.00 ^{ab}
	1.0	1.00±1.000 ^{ab}	1.67±1.155 ^{ab}
	1.5	1.00±1.000 ^{ab}	1.00±1.000 ^{ab}
HTB	0.5	1.33±0.577 ^{ab}	1.67±0.577 ^{ab}
	1.0	0.67±0.577 ^a	1.33±0.577 ^{ab}
	1.5	0.67±0.577 ^a	0.67±1.155 ^a
Polar extracts			
ETL	0.5	1.33±0.577 ^{ab}	1.67±0.577 ^{ab}
	1.0	0.67±0.577 ^a	1.00±0.000 ^{ab}
	1.5	0.67±0.577 ^a	0.67±0.577 ^a
ETB	0.5	1.33±0.577 ^{ab}	1.33±0.577 ^{ab}
	1.0	0.67±0.577 ^a	1.67±0.577 ^{ab}
	1.5	1.00±1.000 ^{ab}	1.00±0.000 ^{ab}
Control	0.0	4.67±1.53 ^c	4.67±1.155 ^c

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test.

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stem bark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stem bark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Effect of *Trichilia heudelotii* extracts on the emergence of *Callosobruchus maculatus* first filial progeny after treatment

Tables 6 and 7 shows the emergence of first filial generation of *C. maculatus* after treatment with *T. heudelotii* non-polar (hex) and polar (EtOH) extracts from 28 and 55 DAT. There was a significant difference between the different treatments and the control at 28, 29, 30 and 32 DAT (Table 6). All the various treatments had a significantly low number of progeny emergence whereas the control did not show any reduction in the emergence of *C. maculatus* progeny. However, at

28 DAT, both the non-polar and polar stem bark extracts (treatments) had the least mean number ($M = 0.00$) of progeny emergence of *C. maculatus* at a concentration level of 1.5 ml, which is significantly lower ($p < 0.05$), when compared to the control, which had the highest mean number ($M = 2.67$) of emergence, as shown in Table 6.

In consideration of the shelf life of the various extracts, Table 7 indicated that the various extracts were significantly ($p < 0.05$) effective from 55 to 59 DAT in the suppression of *Callosobruchus maculatus* first filial progeny in dose dependent manner, when compared to the control. Further observation revealed that the

NOVEL APPLICATION OF TRICHILIA HEUDELOTII ON THE CONTROL OF COWPEA BEETLE

non-polar (hex) and EtOH (polar) stem bark extract of *T. heudelotii* gave the least numbers of F1 progeny emergence at the highest concentration (1.5 ml) at 55 (0.33), 57 (0.33) and

59 (0.33) DAT, respectively, and were the best extracts in restricting the emergence of new generation of *Callosobruchus maculatus* from 55 DAT as shown in Table 7.

Table 6 - Effect of *T. heudelotii* extracts on the emergence of F1 Progeny of *Callosobruchus maculatus* 28 to 32 days after treatment

Treatment	Conc. (ml)/100 g cowpea	Days after treatment (DAT)				
		28	29	30	31	32
Non-polar extracts						
HTL	0.5	0.667±0.577 ^{ab}	1.00±1.000 ^a	1.33±1.533 ^{abcd}	1.33±0.577 ^{ab}	1.33±0.577 ^{bcd}
	1.0	1.33±0.577 ^{abc}	1.33±0.577 ^a	1.00±1.000 ^{abc}	1.00±1.000 ^{ab}	0.67±0.577 ^{ab}
	1.5	1.00±0.000 ^{ab}	0.67±0.577 ^a	1.33±1.155 ^{abcd}	0.67±0.577 ^{ab}	0.33±0.577 ^{ab}
HTB	0.5	0.33±0.577 ^{ab}	1.33±0.577 ^a	0.67±0.577 ^{abc}	0.33±0.577 ^a	0.33±0.577 ^{ab}
	1.0	0.33±0.577 ^{ab}	1.00±1.000 ^a	0.67±0.577 ^{abc}	0.00±0.000 ^a	0.33±0.577 ^a
	1.5	0.00±0.000 ^a	0.67±0.577 ^a	0.33±0.577 ^{ab}	0.00±0.000 ^a	0.67±0.577 ^a
Polar extracts						
ETL	0.5	0.67±0.577 ^{ab}	1.00±1.000 ^a	1.33±1.155 ^{abcd}	1.00±0.000 ^{ab}	1.00±0.000 ^{abc}
	1.0	1.00±1.000 ^{ab}	1.00±0.000 ^a	1.00±1.000 ^{abc}	1.00±1.000 ^{ab}	0.33±0.577 ^a
	1.5	0.67±0.577 ^{ab}	0.67±0.577 ^a	0.67±0.58 ^{abc}	0.33±0.577 ^{ab}	0.33±0.577 ^a
ETB	0.5	0.33±0.577 ^{ab}	1.33±0.577 ^a	0.67±0.577 ^{abc}	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.0	0.33±0.577 ^{ab}	1.00±1.000 ^a	0.67±1.155 ^{abc}	1.00±1.000 ^{ab}	0.33±0.577 ^a
	1.5	0.00±0.000 ^a	0.67±0.577 ^a	0.00±0.000 ^a	0.00±0.000 ^a	0.67±0.577 ^{ab}
Control	0.0	2.67±1.155 ^c	3.00±1.73 ^c	2.67±1.155 ^d	3.00±1.000 ^{ab}	2.33±0.577 ^a

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test.

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stem bark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stem bark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Table 7 - Effect of *T. heudelotii* extracts on the emergence of F1 Progeny of *Callosobruchus maculatus* 55 to 59 days after treatment

Treatment	Conc. (ml)/100 g cowpea	Days after treatment (DAT)				
		55	56	57	58	59
Non-polar extracts						
HTL	0.5	1.67±0.577 ^{bc}	1.67±0.577 ^a	1.67±0.577 ^{bcd}	1.33±1.533 ^{ab}	0.67±0.577 ^{ab}
	1.0	1.00±0.000 ^{ab}	1.33±0.577 ^a	1.00±0.000 ^{abc}	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.5	1.00±0.000 ^{ab}	0.33±0.577 ^a	1.00±1.000 ^{abc}	0.67±0.577 ^{ab}	0.67±0.577 ^{ab}
HTB	0.5	0.67±0.577 ^{ab}	1.33±0.577 ^a	0.67±0.577 ^{abc}	0.67±0.577 ^{ab}	0.33±0.577 ^a
	1.0	0.67±0.577 ^{ab}	0.67±0.577 ^a	0.67±1.155 ^{abc}	0.67±0.577 ^{ab}	0.67±0.577 ^{ab}
	1.5	0.33±0.577 ^{ab}	0.67±1.155 ^a	0.33±0.577 ^{ab}	0.00±0.000 ^a	0.33±0.577 ^a
Polar extracts						
ETL	0.5	1.33±1.577 ^{abc}	1.67±1.155 ^a	1.67±0.58 ^{bcd}	1.67±1.155 ^{bc}	1.33±0.577 ^{abcd}
	1.0	1.00±0.000 ^{ab}	1.33±0.577 ^a	1.33±0.577 ^{abc}	1.00±1.000 ^{ab}	0.67±0.577 ^{ab}
	1.5	1.00±1.000 ^{ab}	1.00±0.000 ^a	1.00±1.000 ^{abc}	0.67±0.577 ^{ab}	1.00±0.000 ^{abc}
ETB	0.5	0.67±0.577 ^{ab}	1.33±0.577 ^a	1.00±0.000 ^{abc}	0.67±0.577 ^{ab}	0.67±0.58 ^{ab}
	1.0	0.67±0.577 ^{ab}	0.67±0.577 ^a	1.00±0.000 ^{abc}	0.67±0.577 ^{ab}	0.33±0.58 ^a
	1.5	0.33±0.577 ^{ab}	0.67±0.577 ^a	0.33±0.577 ^{ab}	0.67±0.577 ^{ab}	0.33±0.58 ^a
Control	0.0	2.67±1.155 ^c	3.00±1.00 ^b	2.67±1.155 ^d	3.00±1.000 ^c	2.33±0.577 ^d

Means followed by the same letter (s) in a column are not significantly different from each other at $p < 0.05$ according to New Duncan's Multiple Range Test.

Keys: Conc. = Concentration; HTL & HTB = Hexane leaf and stem bark extract of *T. heudelotii*, respectively; ETL & ETB = Ethanol leaf and stem bark extract of *T. heudelotii*, respectively; DAT = Day after treatment.

Qualitative phytochemical analysis of *T. heudelotii* leaf and bark

Phytochemical analysis of both the non-polar (hex) and polar (EtOH) leaf and stem bark extracts of

T. heudelotii is shown in Table 8. The result revealed the presence of flavonoids, terpenoids, alkaloids, saponins in both the non-polar and polar extracts of the plant.

Table 8 - Qualitative phytochemical analysis of *Trichilia heudelotii* leaf and stem bark extracts

Phytochemicals	Leaf extract		Stembark extract	
	HTL	ETL	HTB	ETB
Flavonoids	-	+	+	+
Terpenoids	+	+	+	+
Saponins	+	+	+	+
Alkaloids	+	+	+	+
Tannins	-	-	-	+
Anthraquinones	-	-	-	-
Steroids	+	-	+	-
Phenolic compounds	-	+	-	+
Anthocyanins	-	-	-	-

Keys: + = present; - = absent; HTL & ETL = Hexane and ethanol leaf extract of *Trichilia heudelotii*, respectively; HTB & ETB = Hexane and ethanol stembark extract *Trichilia heudelotii*, respectively.

DISCUSSIONS

The genus *Trichilia* has been reported to be a good source of different classes of compounds with insecticidal potentials in the control of insect pests (Sengottayan, 2013; Curcino-Vieira *et al.*, 2014; García-Gómez *et al.*, 2019). The use of different polarity of organic solvent extracts of both leaf and stem bark of *T. heudelotii* gave strong insecticidal activities against the various life stages of *C. maculatus*. To the best of our knowledge, this is the first report of the insecticidal potential of extracts from leaf and stem bark of *T. heudelotii*. In the ability to cause adult mortality, the polar EtOH extract of *T. heudelotii*

stem bark was observed to be the most effective at increasing concentration and therefore served as the best extract in the suppression of *C. maculatus* adults. This findings was in accordance with the reports of some scientists stating the use polar extracts of various plant parts of different *Trichilia* species (from genus *Trichilia*) with antifeedant effects on *Spodoptera litura* and the last instar of *Spodoptera exigua* (López-Oleguín, 1998; Wheeler and Isman, 2001). Similarly, in 2019, García-Gómez and co-workers reported the reduction in the mean survival time of the larvae of *Copitarsia decolora* using methanol (polar) stem bark extract of another *Trichilia* species (*Trichilia hirta*) from the same genus.

Furthermore, the presence of these classes of compounds; saponins, terpenoids, alkaloids, steroids and tanins lying more in the stem bark EtOH polar extract of the plant could probably have caused induction of the quick mortality of *C. maculatus* used in this study as a result of the polar extract incapacitating more adults of the insect (López-Olegrín, 1998; Wheeler and Isman, 2001). The adult mortality effect could also be attributed to the different phytochemicals extracted from the leaf and stem bark of the plant (Sengottayan, 2013; Curcino-Vieira *et al.*, 2014; García-Gómez *et al.*, 2019).

In addition, the shelf life of *T. heudelotii* extracts against *C. maculatus* adults investigated in this study revealed that the organic extracts (both non-polar and polar) used was effective at an extended time of 30 days when applied and thus could provide long term protection on stored cowpea against the insect pest and as such could possibly serve as an excellent alternative to synthetic insecticides. At the highest concentration (1.5 ml) used, all the extracts were significantly able to suppress the oviposition rate of *C. maculatus*, however the EtOH (polar) extract of stem bark of *T. heudelotii* gave the least oviposition also suggesting the use of the polar extract as an alternative crop protectant. Previous reports has shown the use of various polar organic extracts (aqueous, methanol, acetone, ethyl acetate) of *Trichilia Americana*, *Trichilia hirta* and

Trichilia havanensis had reduced fecundity of *Copitarsia decolora* at least by one-third and fertility by almost 50% at 1,500 ppm (García-Gómez *et al.*, 2019). Thus, the results in this study was not far-fetch.

All the plant treatments (extracts) at the larvae and pupae stage of *C. maculatus* showed significant levels of restriction of the larvae emergence and pupation of the insect pest. However, the non-polar (hex) leaf extract, both polar (EtOH) leaf and stem bark extracts of *T. heudelotii* had the least mean number of *C. maculatus* larvae emergence, while on the pupae emergence, the polar (EtOH) leaf, both the non-polar (hex) and polar (EtOH) stem bark extracts of the plant gave the lowest emergence of pupae of the insect pest all at an increased concentration rate of 1.5 ml.

At 30 DAT, the shelf life of the organic extracts indicated that the non-polar (hex) stem bark of *T. heudelotii* at 1.5 ml had the least mean larvae and pupae emergence as well as the polar (EtOH) leaf extract giving fewer pupae emergence, though, all the treatment extracts were sufficiently effective in restricting the larvae and pupae emergence of the insect pest at an extended period of 30 days, compared to the control which did not show significant reduction in the emergence of *C. maculatus*. Therefore, these effects on larval and pupae emergence as a result of the use of the extracts of *T. heudelotii* may be due to hormone alterations interfering with the process

NOVEL APPLICATION OF TRICHILIA HEUDELOTII ON THE CONTROL OF COWPEA BEETLE

(Mordue and Blackwell, 1993). Our findings were in accordance with those reported in literature where the polar extract (methanol-MeOH) of *Trichilia hirta* stem bark significantly reduced the mean survival time of the larvae of *Copitarsia decolora* (García-Gómez *et al.*, 2019). Similarly, another specie from the genus *Trichilia* (*Trichilia sylvatica*) was previously reported to have decrease in fecundity and fertility by about 87 to 100% of *Spodoptera frugiperda* when its larvae were fed with polar (MeOH) extract (Freitas *et al.*, 2014).

The treatment (extracts) of *T. heudelotii* used in this study also showed a significant reduction in the emergence of *C. maculatus* filial progeny. Close observations showed that both the non-polar (hex) and polar (EtOH) stem bark extracts of the plant at the highest concentration rate had the least number of *C. maculatus* first filial progeny emergence and performed better in its ability to suppress the emergence of the new generation of the insect pest. Aba-Toumou *et al.* (2016) reported the use of both non-polar (hex) and polar (ethyl acetate- EtOAc) extracts of *Trichilia gilgiana* being effective against different storage pests; *S. zeamais*, *T. castaneum* and *R. dominica*. The insecticidal activity of *T. gilgiana* against *S. zeamais* and *T. castaneum* were also found in literature (Aba-Toumou *et al.*, 2012b). Previous reports cited in this study further corroborates our findings, suggesting the use of different polarity of organic solvents

extracts of *Trichilia heudelotii* as a natural alternative remedy for crop protectants against insect pests.

CONCLUSION

Non-polar (hex) and polar (EtOH) extracts of *Trichilia heudelotii* were effective in controlling the various life stages of bruchid population in stored cowpea. Strong insecticidal activities against *C. maculatus* with a high rate of mortality were reported within 96 hours using both non-polar and polar extracts of *Trichilia heudelotii* leaf and stem bark at the highest concentration (1.5 ml). In addition, the extracts were able to protect the cowpea against *C. maculatus* at an extended period of 30 days suggesting the extracts to possess a long term insecticidal effect against insects' pest infesting cowpea. The potential of *T. heudelotii* extracts as a plant derived insecticides in the control of *C. maculatus* could further be studied to investigate it's usage as an eco-friendly biopesticides alternative to the use of conventional synthetic pesticides. Further research is suggested to possibly isolate the actual compound(s) responsible for the insecticidal activities and further testing carried out. Thus, *T. heudelotii* could possibly serve as a sustainable means of pest control where the plant is readily available in high quantity.

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