# **Original Research Article**

# TOXICITY AQUEOUS EXTRACT OF THE CASTOR OIL (*Ricinus comunus*): TO *Maconellicoccus hirsutus* (GREEN) (HEMIPTERA: PSEUDOCOCCIDAE)

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# ABSTRACT

The objective of this study was to evaluate the insecticidal effect of aqueous extract of castor bean cakepie for the control of *Maconellicoccus hirsutus*. The toxicity testsexperiments were performed in the laboratory. The toxicity test was conducted. Subsequently it was estimated lethal concentrations 50 and 90% (LC<sub>50</sub>and LC<sub>90</sub>, respectively) were estimated. Tween80<sup>®</sup> + distilled water was used aAs surfactants and solvent and witness was used distilled water + respectively. Tween80<sup>®</sup> - The assay was performed in petri dishes containing an *Coffea canephora* leaf disc. The spraying was carried out in Torre de Potter spray tower. The extract was toxic to mealybug and to the probit model. Estimates of LC<sub>50</sub> and LC<sub>90</sub> were 5.32% and 29.30%, respectively. It was concluded found t that the aqueous extract of castor pie is promising for the management of *M. hirsutus*.

Keywords: Insect pests, pink mealybug, Ricinus communis.

#### 1. INTRODUCTION

InFor Brazil, *Maconellicoccus hirsutus*(Green)(Hemiptera: Pseudococcus), commonly known as the pink mealybug or the pink hibiscus mealybug, it is a quarantine pest type A2, may cause damage in more than 200 plant species, as was fact already registered recorded in other countries. Brazil offers idealadequate climatic conditions for its development, being its greater economic impact on the fruticulture. While feeding, the mealybugs introduce toxic substances in the plants, causing poor formation of leaves and fruits, apical growth, and may lead to premature senescence shriveled, dry and fall of flowers [1].

Because ilt is a recent pest in the country, <u>becausesince</u> the first signs of it were <u>only</u> observed <u>duringin the year\_2010</u>, <u>T</u>there is <u>therefore</u> no effective method of <u>controlling the</u> <u>pesthandling</u> and <u>its</u>the environmental <u>impactconsequences</u> that it can bring agre therefore <u>also</u> unknown. For the\_countries that already <u>have-live with</u> this pest, the biological control, using parasitoids and predators, has been an alternative. However, due to its rapid dispersalsemination, studies are needed to develop management programs to combat this pest [2].

Researches related to the use of extracts and substances obtained from plants are demonstrating a satisfactory efficiency in the control of some pests [3]. These substances

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act as a defense mechanism for plants to protect themselves from pests and diseases and to help with the adverse environmental conditions [4].

Castor bean [*Ricinus communis*\_L. (Euforbiaceae)] is an example of an insecticidal plant in which studies have demonstrated its effectiveness in pest control. Lima et al.[5] obtained excellent results by testing the oil extracted from castor <u>oil</u> seeds on *Diaphania nitidalis*(Stoll) (Lepidoptera: Pyralidae) caterpillars. The author reports that on the second day of evaluation, a mortality of 92% <u>was obtained of these caterpillars occurred</u>. Santiago et al. [6] studying the aqueous extract of 10% (v / v) *R. communis* green fruits on *Spodoptera frugiperda* (Smith) (Lepidoptera: Noctuidae) caterpillars observed <u>reduced ?</u> activity on biological parameters, such as larval and pupal duration and viability, reduction in pupal weight. Carvalho [7], on the other hand, verified reduced activity of aqueous extracts of the castor bean castor oil the results showed high toxicity of the oil for the species.

However, castor oil is widely used in the production of biodiesel. In theis process of production, a residue is generated, the pie, which, in the case of <u>castor beanR communis</u>, does not yet have a specific use [8]. However, studies have already proven that the insecticidal activity of this <u>byproductresidue may be</u> due to the presence of ricin, which is a <u>very</u> toxic substance [9]. Thereforeus, with the need to develop methods for the management of pink cochineal, the residue of the castor bean pie may be considered as an alternative to the management of this pest.

Theus, this project has the objective of this of studying is to quantify the insecticidal potential of the castor bean cake as an in the alternative control method of *M. hirsutus*.

#### 2. MATERIAL AND METHODS

The experiment was carried out at the Federal Institute of Education, Science and Technology of Espírito Santo — Campus Itapina (IFES-Campus Itapina), Colatina, ES, Brazil, in air-conditioned chambers at 25  $\pm$  1°C, relative humidity 70%  $\pm$  10 and 12h photophase. Seeds of *R. communis* were used to study the insecticidal activity.

#### 2.1 Rearing of pink mealybug M. hirsutus

Samples of *M. hirsutus* were collected in commercial <u>cocca</u> plantations<u>of</u> <u>cocca</u> [*Theobroma cacao* (Malvaceae)] in the municipality of Colatina, ES, Brazil and taken to the laboratory. The breeding technique <u>usedadopted</u> was an adaptated from ion of Sanches & Carvalho [10]. Pumpkins were used in the initial state of maturation to feed the *M. hirsutus*. In the initial infestation of the pumpkins, the individuals were collected in the field on infested host plants. After establishment of the colony, the process of <u>multiplication of the</u> mealybug <u>breeding</u> began. If there was a need to change the pumpkins, new fruits were placed in contact with those already infested for approximately two hours. The approximation of these favored the transfer of newly hatching nymphs from the cochineal to the new fruit due to its mobility in that phase.

#### 2.2 Obtaining aqueous extract from castor pie

The preparation of the pie was through the extraction of the oil from the castor seeds collected at the IFES-*Campus* Itapina. After this procedure, the pie generated from the extraction was submitted to grinding, in knives mills, to obtain a fine powder.

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To extract the *R*. communis pie extractactive ingredient, the powder of the ground plantvegetable material (30 g) was transferred to an Erlenmeyer flask (100 mL) containing distilled water and Tween<sup>®</sup> 80 (0.05% v / v) adhesive spreader, to obtain 100 mL of the initial solution at 30% (w / v). Subsequently, it was maintained under homogenization in a cross stirrer (240 rpm) for a period of 24h. After this time, the mixture was filtered with fine sterile tissue, and transferred to a volumetric flask and the volume checked to 100 mL.

## 2.3 Bioassays

The direct application tests were carried out on the individuals of the cochineal to evaluate the insecticidal action and to estimate the lethal concentration for a medium population  $(LC_{50})$ .

#### 2.3.1 Toxicity test of extract to M. hirsutus

The treatment consisted of 6 replicates with 12 individuals of the *M. hirsutus* by repetition, totaling 72 insects. The replicates were maintained in Petri dishes (10.0 x 1.2 cm) on coffee leaf discs <u>of</u> about 4 cm in diameter. The discs were fixed to the petri dish with a 0.5 cm layer of agar-agar solution and solid <u>petroleum jelly (V</u>+aseline) around the disc to prevent insect escape.

The spray was <u>applied with acarried out by</u> Potter Tower<sup>®</sup> <u>atwith</u> a pressure of 15 lib/in<sup>2</sup> and 6 mL of solution for each repetition, corresponding to an average volume of 1.62 mg/cm<sup>2</sup>. Distilled water and Tween<sup>®</sup> 80 adhesive spreader (0.05% v / v) were used as controls. The insecticidal effect was evaluated 24, 48 and 72 hours after spraying. The experiment was conducted in an air-conditioned room (25 ± 10 ° C, 70 ± 10% RH and 12-hour photophase). The design was completely randomized, <u>andwith</u> mortality <u>assessments were conducted</u> after 72 hours. <u>of treatment and control compared by unilateral t test (p ≤ 0.05) [11].</u>

## 2.3.2 Estimation of lethal concentration (LC)

Each treatment consisted of 6 replicates and 12 insects per replicate, totaling 72 individuals.
The sprays were <u>appliedfollowed</u> as described above for the toxicity test. The treatments were composed of 4 concentrations of aqueous extract of castor bean cake, spaced on a logarithmic scale (limits of 0.01% to 30% m / v). Distilled water plus Tween 80<sup>e</sup> (0.05% v / v) was used as the solvent <u>and was used i</u>. In the control treatment<u>it was only the solvent</u>. Mortality was corrected by the formula proposed by Abbott [12]. The experiment was conducted in an air-conditioned room (25 ± 10 ° C, 70 ± 10% RH and 12h photophase) and evaluated 24, 48 and 72 h after spraying the solutions. With mortality corrected data of 72 h, lethal concentrations were estimated using Probit analysis [11-13].

#### 2.3.3 Data analysis

All data processing and graphical presentation of results were performed in the computational environment R, as proposed by Carvalho et al. [11]. <u>ef-t</u>Treatment means were and control compared by a unilateral t test ( $p \le 0.05$ ) [11].

# 3. RESULTS AND DISCUSSION

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The toxicity test indicated that the aqueous extract of the <u>castor bean pie</u> <u>differed</u> significantly from the control (t = 14.02, P< .0001) (**Figure 1**), with an average mortality of 83.46%. Based on this result, <u>it waswe</u> proceeded to obtain the estimated lethal concentration.



Treatments

Figure 1. Mortality of *Maconellicoccus hirsutus* (Green) (Hemiptera: Pseudococcidae) treated with aqueous extract of *Ricinus communis* (Euphorbiaceae) at 30% (m / v).

In the LC<sub>50</sub> bioassay there was an increase in the percentage of mortality of the **Dink cochinea** proportional to the increase of the concentration of the aqueous extract of castor bean cake. The data <u>was</u> adapted to the <u>Probit</u> model of <u>Probit</u> (**Table 1** and **Figure 2**). The slope of the concentration-mortality curve was 1.73. The lethal concentration required to cause mortality of 50% of the *M. hirsutus* (LC<sub>50</sub>) population was estimated to be 5.32% and to kill 90% of the population (LC<sub>90</sub>) was estimated to be 29.30%.

Table 1. Concentration-mortality curve parameters and respective LC50 and CL9	0 of
aqueous extract of <mark>castor bean pie</mark> on <i>Maconellicoccus hirsutus</i> (Green) (Hemipt	era:
Pseudococcidae) (Temp: $25 \pm 1^{\circ}$ C, RH 70 $\pm 10\%$ and 12h of photophase)	

n <sup>1</sup>	Slope ± SE <sup>2</sup>	LC <sub>50</sub> <sup>3</sup> (Cl <sup>4</sup> a 95%) (g/100ml)	LC <sub>90</sub> <sup>3</sup> (Cl <sup>4</sup> a 95%) (g/100ml)	DF⁵	χ <sup>2(6)</sup>	P <sup>7</sup>
200	1.73±0.52	5.32 (1.00-8.82)	29.30 (21.40- 73.43)	2	0.5769	> .05

<sup>1</sup> Number of insects used in the test; <sup>2</sup>Slope ± standard error; <sup>3</sup> Lethal concentration; <sup>4</sup> CI confidence interval at 95% probability; <sup>5</sup> Degrees of freedom; <sup>6</sup> Chi-square test; and <sup>7</sup> p-value.

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# Figure 2.Concentration-mortality curve and respective LC50 and CL90 of aqueous extract of castor bean pie on *Maconellicoccus hirsutus* (Green) (Hemiptera: Pseudococcidae) (Temp.: $25 \pm 1$ ° C, RH 70 $\pm 10\%$ and 12h of photophase).

The toxicity of castor bean to insects and mites is due to different compounds, <u>such asamong them we</u> can mention ricin, <u>which is a protein in that is a protein</u> found exclusively in the endosperm of the seed, and is not detected elsewhere in the plant [14]. Another substance found in castor oil is ricinin, an alkaloid found in all parts of the plant. The insecticidal effect occurs mainly due to this toxin [15]. However, this is in low concentrations and, consequently, <u>has a</u> low toxic activity. In castor bean fruits, mainly in seeds, the ricinin content is higher in the outer capsule, medium in the seed shell and small in the endosperm [14]. This toxin has two subunits with different functions, but acting together. One of them has an inhibitory effect on the digestion processibility by the action of  $\alpha$ -amylase enzyme inhibitors, preventing the digestion and absorption of the starch and the other has an insecticidal effect due to the action of ribosome inactivating proteins (RIPs), which promotes cell death by inhibition of protein synthesis [16].

However, as the insect under study feeds directly from the sap of the plant, because it is an Auchenorrhyncha hemipterane, these compounds hardly act through the gastrointestinal tract. Due to the spraying process, the insects were subjectedmitted to dermal and possibly respiratory contact. In this context, the insect could be absorbing these compounds via integument or respiratory, through the spiracles.

Different from other mealybugs. *M. hirsutus* is different from other mealybugs becausedoes not have the its body is protected by a pulverulent carapace [17], which could facilitate the absorption of toxins by the dermal route.

However, <u>no</u> studies on the action or ricin absorption and, or ricinine in cutaneous and respiratory systems of the insects were not reported.

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Another factor that should be considered is the amount of toxins presents in castor bean cake and if there is seasonal variability due to the harvest and storage time, and the effect on the toxicity of these substances. A study carried out with *Moringa oleifera* (Moringaceae) demonstrated increased toxicity over time of storage on the mite *Tetranychus urticae* (Acari: Tetranychidae) [18]. These results justify the need for further chromatographic and storage studies in order to better understand the toxicity mechanisms of the castor bean extract and possibly produce a commercial product that is safe for farmers and the environment.

# 4. CONCLUSION

It is concluded that the extract of castor bean cake is promising for the management of *M. hirsutus*. However, studiesy in semi-field and field conditions are should be conductedstimulated seeking to optimize and validate the results of the present study.

In addition, stud<u>iesy</u> to elucidate the toxicological mechanisms of ricin and ricinine in <u>h</u>Hemipteran insects <u>may provide us with seen great truth of these would appear as</u> promising molecules for the control of insect pests.

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