Control of locust Schistocerca gregaria (Orthoptera: Acrididae) by using imidaclorprid

Sabbour M.M1 and S.M. Singer2
1.Department of Pests and Plant Protection, 2. Vegetable research Dep. National Research Centre, Dokki, Cairo, Egypt
E-mail: sabbourm@yahoo.com Tel.: +202/01223305136 Fax: +202/33370931

Abstract— The effect of imidaclorprid IMI on the target insect pest Schistocerca gregaria and the C50 recorded, 270, 214, 223, 249 and 240 mg/L for newly hatched, nymphs, Last nymphal stage Adult ♀ and Adult ♂ respectively. Under semfield conditions, the corresponding LC50 obtained, 221, 243, 254, 256 mg/L for newly hatched, nymphs, Last nymphal stage Adult ♀ and Adult ♂ respectively of S. gregaria. Also, under semi field conditions, the number of S. gregaria were significantly decreased after the IMI first applications. The infestation number obtained 1.1±2.1, 9±2.1, 13±3.8 and 21±3.6 individuals after 20, 50, 90 and 120 days as compared to 12.2±3.4, 36±3.8, 58±6.6 and 98±8.7 individuals in the control.

Key words: locust, Schistocerca gregaria, imidaclorprid (IMI).

1. INTRODUCTION

Imidaclorprid, (1-[(6-chloro-3-pyridinyl)methyl]-4,5-dihydro-N-nitro-1H-imidazol-2-amine (IMI), is a systemic and chloronicotinyl insecticide, that specifically blocks the microtubulin neuronal pathway. It has recently been demonstrated to be highly effective as a systemic insecticide [Byrne et al. 2005, 2006]. Imidaclorprid is the first commercially available representative of a new chemical class, the chloronicotinoyl or neonicotinoid insecticides. It was synthesized in 1985 and the first registration was achieved in France (1991). It is a systemic broad-spectrum insecticide and acts as a contact and stomach poison against sucking and some biting insects (rice hoppers, aphids, thrips, whitefly, termites etc.). It can be applied for seed, soil or foliar treatment. The molecule exhibits a novel mode of action as it is an agonist of the nicotinic acetylcholine receptor leading to paralysis and death of pest organisms. The life history of the desert locust, S. gregaria (Forsk.), and the epidemiology of its outbreaks were obscure until the 20th Century. One of the discoveries of importance in this century was that of solitary and gregarious phases. The life history of the desert locust, Schistocerca gregaria (Forsk.), and the epidemiology of its outbreaks were obscure until the 20th Century. One of the discoveries of importance in this century was that of solitary and gregarious phases. The most prominent feature of the ‘arbeh’ is that it can exist as scattered individuals within the ‘recession areas’ or, when gregarious, as swarms throughout the ‘invasion areas’. This is because the locust exists in different phases. When breeding conditions lead to an increase in the number of locusts crowded together, the insects have the ability to change their color, behavior, shape and physiology, with color and behavior being the characteristics to change first. Aim of this work to evaluate the Imidaclorprid IMI against locust.

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2. MATERIALS AND METHODS

2.1. Insect rearing

Locust was reared under laboratory condition for several generations on semi-artificial diet as mentioned by Sharaby et al. (2010).

2.3. Preparation of nano-imidaclorprid

Imidaclorprid Nanoparticles were synthesized by hydrolyzing titanium tetra isopropoxide in a mixture of 1:1 anhydrous ethanol and water. 9 ml of titanium tetra isopropoxide is mixed with 41 ml of anhydrous ethanol (A), 1:1 ethanol and water mixture is prepared. (B) Solution A is added in drop wise to solute ion B and stirred vigorously for 2 hrs. At room temperature hydrolysis and condensation are performed, using 1M sulphuric acid and stirred for 2 hrs. Then the ageing was undertaken for 12 hrs. The gel was transferred into an autoclave and tightly closed, and the mixture was subjected to hydrothermal treatment at 353K for 24 hrs. After filtration the solid residue was washed thoroughly with water and ethanol mixture, dried at 373K in an oven and calcined at 773K.

2.4. Efficacy of imidaclorprid against the target insect pests

The insecticide imidaclorprid were tested at the 6 concentrations: 6 g, 5g , 4g, 3g, 2g, 1g. The insecticide, prepared 6 concentrations (prepared according Sameh et al., 2009) Percentages of mortality were calculated according to Abbott’s formula (Abbott, 1925), while the LC50 values was calculated throughout probit analysis (Finney, 1971). The experiment was carried out under laboratory conditions at 26±2°C and 60-70% RH.

2.5. Bioassays

The insecticidal efficacy of nano-imidaclorprid were tested at three dose rates, 0.25, 0.50 and 1 g/kg wheat against the 3rd instar larvae of S. gregaria (Orthoptera: Acrididae). For each case, four glass jars as replicates were used. Each replicate was treated individually with the respective nano-imidaclorprid quantity and then shaken manually for one minute to achieve equal distribution of the imidaclorprid. Subsequently, ten 3rd instar larvae of the two tested species were introduced into each glass jar and covered with muslin for sufficient ventilation. Twelve replicates glass jars containing untreated wheat served as control. Mortality was assessed after 7 d of exposure in the treated and untreated jars. Mortality was corrected according to Abbott (1925). All tests were conducted at 27 ± 2°C and 65 ± 5% relative humidity (RH). All the experiments were repeated three times.

The ovipositional deterrent effects of nano-imidaclorprid were also tested. The nano-imidaclorprid were used at the rate of 0.5 g/kg wheat. Four replicates of 100 g wheat for each treatment were used. Each replicate was treated individually with the formulations for 1 min and put inside glass jars. Four replicates in jars containing untreated wheat served as control. Subsequently, one pair of newly emerged adults were introduced into each jar. The number of deposited eggs on treated or untreated wheat/female was counted and the
percent repellency values were calculated according to the equation of Lwande et al. (1985), D = (1 - T/C) x 100, where: T and C represent the mean number of deposited eggs per female of the treated and check set, respectively.

Efficacy of tested nano-imidaclorprid applied alone on the mean number of deposited eggs of target insects for conducting the combination tests with imidaclorprid formulations (0.5 g/kg of grains). The imidaclorprid alone were used at rate (1.0 g/kg) of grains. Four replicates of 100 g grains for each treatment were used. Each replicate was treated individually with treatments and then shaken manually for 1 min to achieve equal distribution of the dust in the entire formulation quantity and was placed in glass jar. Four replicates jar containing untreated grain served as control. Subsequently, one pair of newly emerged adults were introduced into each jar. The number of deposited eggs on treated or untreated grains/female was counted. The data was analyzed using analysis of variance (ANOVA), where significant differences between the treatments were observed. Mean values were significantly separated by using the least significant difference (LSD) test at 5% level (Sokal and Rohlf 1981).

2.6. Efficacy of Imidaclorprid against the target insect pests.

The insecticide Imidaclorprid were tested at the 6 concentrations: 6 g, 5 g, 4 g, 3 g, 2 g, 1 g. The insecticide, prepared 6 concentrations (prepared according Sameh et al., 2009) Percentages of mortality were calculated according to Abbott’s formula (Abbott, 1925), while the LC50 values was calculated throughout probit analysis (Finney, 1971). The experiment was carried out under laboratory conditions at 26±2°C and 60-70% RH. Nano-Imidaclorprid, were prepared by the National research center team microbiological team according to Leiderer et al. (2008). The tested pathogens was considered the standard for comparison with the other ones.

3. RESULTS AND DISCUSSION

Table 1 show the effect of IMI on the target insect pest Schistocerca gregaria and the LC50 recorded, 278, 214, 223, 249 and 240 mg/L for newly hatched, nymphs, Last nymphal stage Adult ♀ and Adult ♂ respectively. Under semifield conditions, the corresponding C50 obtained, 221, 243, 254, 256 mg/L for newly hatched, nymphs, Last nymphal stage Adult ♀ and Adult ♂ respectively of Schistocerca gregaria (Table 2).

Also, under semi field conditions, the number of Schistocerca gregaria were significantly decreased after the IMI first applications. The infestation number obtained 1.1±2.1, 9±2.1, 13±3.8 and 21±3.6 individuals after 20, 50, 90 and 120 days as compared to 12.2±2.2 individuals after treated with destruxin in the control. Sabbour, 2013 a,b reported that. Desert locust Schistocerca gregaria bioassayed by using the leaves containing early stages larvae and the data were recorded after 1, 2, 3 and 4 days after treatment. Results showed that range of mortality was between 84-65% based on the end point data. The minimum of three days to achieve 60% mortality was proved by probit analysis of time-mortality responses. They found that, the range of mortality was between 88-65% based on the end point data. The minimum of three days to achieve 50% mortality was proved by probit analysis of time-mortality responses. The same results obtained by Sabbour and Singer 2015. Sabbour a, b, and b 2015. Sahab et al 2015 found the insecticidal activity the nano-chitosan (CS-g-PAA) showed highest effect against the three insect of soybean. as the means number of eggs deposited /female were significantly decreased. Under laboratory and semifield conditions, Aphis gossypii were significantly decreased to 20.9±9.1 and 28.9±9.2 eggs/female respectively as compared to 97.3±4.9 and 90.3±4.9 eggs/female in the control, respectively. The same trends were also observed against Callosobruchus maculatus. Sabbour 2015, a, b, c found that the nano insecticides of Imidaclorprid and fungi strains cases a higher mortality for insect infestations. Our results agree with Sabbour and Abd Raheem 2015, a &b, Sabbour and Singer 2015 a&b and Sabbour and shadia 2015 who find that the nano pesticide decrease the infestation percentage of different pests.

4. Acknowledgments

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Fig 1. Percentage of infestations of the target pests under semifield conditions

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<td>Effect of IMI against Schistocerca gregaria under laboratory</td>
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5. Conclusion

The nature product IMI is effective against the locust \textit{S. gregaria} under laboratory semi field and field conditions.

6. References


Sabbour, M.M. 2013b. Evaluating toxicity of extracted destructin from...


Sabbour, Magda and MA Abdel-Raheem.2015 a. TOXICITY OF THE FUNGUS BEAUVERIA BASSIANA AND THREE OILS EXTRACT AGAINST SITOPHILUS GRANARIES UNDER LABORATORY AND STORE CONDITIONS. American J. of innovative research and applied sci. 251-256.


