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Cylogenitic Effects of the Insecticide Cypermethrin on the Plants Vicia Vaba and Altium Cepa.

### Thesis

Submitted to
Botany Department
University College for Girls
Ain Shams University
in Partial Fulfilment of the Requirements
for the Degree of M.Sc. in
Botany (Cytology)

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National Research Centre

Cairo A.K.E.

1986

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# Cytogenetic Effects of the insecticide Cypermethrin on the plants Vicia faba and Allium cepa

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#### CONTENTS

		Page			
I-	AIM OF THE WORK	1			
II-	INTRODUCTION	3			
III-	MATERIALS AND METHODS	8			
	1- Preparation of the experimental agent	8			
	2- Seed-germination	8			
	3- Root-and shoot-growth	9			
	A- <u>Vicia</u> <u>faba</u>	9			
	B- Allium cepa	10			
	4- Effect on mitosis	11			
	A- Treatment	11			
	a- <u>Vicia</u> <u>faba</u>	11			
	b- <u>Allium</u> <u>cepa</u>	12			
	B- Technique	13			
	a- Feulgen squash method	13			
	C- Scoring	15			
	D- Statistical analysis	15			
	5- Effect on meiosis	15			
	A- Treatment	15			
	B- Technique	16			
	a- Aceto-carmine smear method	16			
	C- Scoring and statistical analysis	17			
IV-	RESULTS				
	1- Seed-germination				
	2- Root-and shoot-growth Central Library - Ain Shams University				

7

	A-	Vic	cia faba	18						
		a-	Seed-soaking-treatment	18						
		b-	Root-treatment	20						
	B-	Al	lium cepa	20						
		a- Root-treatment								
3-	Ef	Effect on mitosis								
	A-	- <u>Vicia faba</u>								
		a- Mitotic index and percentage of								
			abnormal mitoses	20						
			i- Seed-soaking-treatment	20						
			ii- Root-treatment	23						
		b-	Effect on the mitotic stages	29						
			i- Seed-soaking-treatment	29						
			ii- Root-treatment	29						
		c-	Types of the induced abnormalities	33						
			i- Seed-soaking-treatment	33						
			ii- Root-treatment	38						
	B-	All	lium cepa	43						
		a-	Root-treatment	43						
			i- Mitotic index and percentage of							
			abnormal mitoses	43						
			ii- Effect on the mitotic stages	43						
			iii-Types of the induced abnormalities	54						
4-	Efi	fect	on meiosis	66						
	1	Per	centage of abnormal pollen mother							
		cel	lls	66						
	2-	Per	ccentage of abnormalities in the							
		mei	iotic stages	66						

	3-	Types	of	the	induced	abnormalities	69	)
V-	DISCUSSION							ŧ
VI-	SUMMARY							3
VII-	REFERENCES							ŀ
	א כו ג פ ג דו ג פ ג	e ciimm	λOV					

## AIM OF THE WORK

### I- Aim of the Work

The aim of the present investigation is to study the effect of the synthetic pyrethroid insecticide Cypermethrin (I) on:

- 1- Seed germination, root- and shoot- growth of <u>Vicia faba</u> seedlings and root-growth of <u>Allium cepa</u>.
- 2- Root-mitosis of Vicia faba and Allium cepa.
- 3- Meiosis and pollen viability of Vicia faba.

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"Cypermethrin"

### INTRODUCTION

#### II-INTRODUCTION

Since the first successful development of the synthetic Pyrethrin, Allethrin by Dr. La Forge in 1947, a number of workers has started the modification of the natural pyrethrin which resulted in a remarkable change in its property. Particularly, the potent and photostable pyrethroids have been disclosed by the workers in Great Britain and Japan. Thus these pyrethroids are able to use for agricultural pest control in a large scale. (Nakajima, 1982).

The synthetic pyrethroids such as permethrin, Cypermethrin, and Deltamethrin are extremely potent and broad-spectrum insecticides. Although their toxicity to mammals and birds is relatively low, they are extremely toxic to fish and a variety of aquatic invertebrates. These chemicals have low water solubilities and are photostable. In soils, they are degradable via microbial hydrolysis at the ester linkage (Khan, 1982).

Miyamoto and Mikami (1982) discussed the degradation of the synthetic pyrethroids (Permethrin, Cypermethrin, Deltamethrin, Fenvalerate and Fenpropathrin) in plant, soil and the aqueous environment. The pyrethroids undergo a variety of photochemical reactions including trans/Cis and XR /XS interconversions except Fenvalerate.

When applied to the leaves of plants, they were penetrated to the tissues to almost the same degree, but little translocation to the untreated area occurred. When plants are grown in the treated soil, again little translocation of chemical to aerial parts of the plant occurs.

The pyrethroids are degraded in soil mainly by micro-organisms. Hydrolysis of the ester linkage together with oxidation are the major degradation routes. The products are further decomposed yielding  ${\rm CO_2}$  from radiolabeles at various positions of the molecules. The pyrethroids are immobile and can not be leached with water through soil. In the aqueous environment, pyrethroids were rapidly removed from the aqueous phase by adsorption on to sediments. (Miyamoto and Mikami, 1982).

Synthetic pyrethroids are toxic to the house flies, Musca domestica L. (Taylor et al., 1981), with no irritation to nasal passage (Adkins et al., 1971) and with no symptoms of irritation to animals (Faraone et al., 1978).

Synthetic pyrethroids are effective against larvae of various species of mosquitoes (Mulla et al, 1980, and Priester et al., 1981), and to larvae and adults of the Egyptian mosquito <u>C. pipiens</u> <u>L.</u> (Gaaboub, and Abu-Hashish, 1981).

Cypermethrin is also toxic to: larvae of the fall armyworm (Mcclanahan, 1978), house flies and German cockroaches (Scirocchia and Cesaroni, 1979); onion, seed-corn maggot flies and dark sided cut worm (Harris and Turnbull, 1980).

In the fields of wheat, shoot fly, Atheriqona nagvii Steyskal damage was effectively controlled by foliar sprays of Cypermethrin during the early growth phases of wheat crop. (Singh and Saxena, 1981).

Cypermethrin used as field treatments of silage significantly reduced corn borer damage (Ostrinia nubilalis) and increased dry matter yield than untreated plots (Raemisch and Walgenbach, 1983 and 1984).

In mamals, Cypermethrin is readily absorbed, distributed, and metabolised. It is readily cleaved at the ester linkage and subjected to oxidative degradation and conjugation of the metabolic products. Elimination from the body following acute and subecute administration is rapid. (JMPR, 1979).

In a wide variety of studies, there was no carcinogenic or mutagenic potential as evidenced by short-term bioassays or long-term chronic study (JMPR, 1979). However, Cypermethrin showed mutagenic potential in our studies. In

one study, oral administration of the insecticide at a dietary level of 900 ppm for 7 and 14 consecutive days as well as double and multiple (Total 4) dermal treatments (360 mg/kg body wt.) induced a statistically significant increase in the frequency of  $PE_S$  with micronuclei in mouse bone marrow (Amer and Aboul-Ela, 1985).