INFLUENCE OF CERTAIN WILD PLANT EXTRACTS AGAINST TETRANYCHUS URTICAE AND PREDACIOUS MITE AMBLYSEIUS ZAHERI (ACARI: TETRANYCHIDAE: PHYTOSEIIDAE)

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INTRODUCTION

Vegetable crops in Egypt are highly infested by sucking pests as aphid, whitefly and mites. Synthetic pesticides are still the major agent currently used for controlling these pests. The two-spotted spider mite, *Tetranychus urticae* Koch. has been recognized as a factor limiting the production of crops and is controlled by biological means. Its satisfactory control depends on selective pesticides. Although there were some privileges in their use, other problems such as resistance, environmental pollution, residual toxicity in food and hazards to natural enemies and human health hindered the successful application of such control (Pimental *et al.*, 1992). In recent years botanical pesticides have played an important role in the control of spider mite and proved to be effective on target mites (Mansour *et al.*, 1986; Reda and El-Banhawy 1986; Amer *et al.*, 1989).

The effectiveness of the essential oils on mite has been reported by Amer *et al.*, (1993); Ibrahium *et al.*, (1993) and El-Gengaihi *et al.*, (1996). The toxic and deterrent effects of the essential oils were studied on insects and mites by Orphanidis and Kalmoukas (1970); Rodriguez and Levin (1975); Purohit *et al.*, (1983) and Don-Perdo (1985).

The toxicity of the crude extracts for certain plants against adult females and eggs of *T. urticae* Koch, have been reported by several investigators, Barakat, *et al.*, (1985) for acetone and diethyl ether extracts of onion, garlic, caraway, devils apple, lupin, fenugreek, blackpepper, turnip, canna and glonybower on adult females and eggs of *T. urticae* Koch., Osman *et al.*, (1986) for *Nerium oleander* and *Caesalpinia sepiaria* against predatory mite, *Amblyseius gossipi*, El-Halawany *et al.*, (1989) for *Lantana*

camara against *Tetranychus arabicus* and Dimetry *et al.*, (2000) for *Curcuma longa* L., *Nicandera physaloids* L. and *Dodonaea viscose* L. against adult females of *T. urticae*.

The aim of the present study is to evaluate the toxic and repellent effects of the crude extracts of three wild plant, *Lygos raetam*, *Silybum marianum* and *Fagonia indica* on the eggs and adult stages of *T. urticae* and the adult stages of *Amblysius zaheri*.

MATERIAL AND METHODS

Phytophagous mites were obtained from a laboratory colony of *T. urticae* Koch, reared on Lima bean (*Phaseolus vulgaris* L.) foliage at NRC Cairo under laboratory conditions $(27 \pm 2 \, ^{\circ}\text{C})$ and $70 - 75 \, ^{\circ}\text{C}$ r.h.).

The predactions mite A. zaheri Yousef and El-Borolossy was found on leaves of egg plant and was fed on T. urticae in the laboratory.

Three wild plants, *Lygos raetam* (Fam: Leguminosae), *Silybum marianum* (Fam. Compositae) and *Fagonia indica* (Fam. Zygophyllaceae) were collected from different areas in South Sinai. Samples of the collected plants were left to dry in the laboratory for one week. Dried plants were ground using an electric mill, sieved and kept for extraction.

The crude extract was prepared according to the method adopted by Su and Horvat (1981) with some modifications. Samples (1 kg of plant) were blended and soaked in 3 liters of chloroform: methanol (1:1 v/v) for 24 hours and kept in brown colored bottles, 5 liters, provided with tight stoppers with continuous shaking overnight. The extracted solute was separated from the insoluble plant material collected and the latter reextracted with another 2 liters of the same solvent system for another 12 hours and the solvent was also separated. The collected two extracts (5 liters) were filtered over anhydrous sodium sulphate and then the solvent was evaporated under reduced pressure using a Rotary evaporator. The remaining residues were subjected to successive extractions with 4 different solvents (250 ml each), namely n-hexane, diethyl ether, ethyl acetate and ethanol. The crude extract of each solvent was filtered over anhydrous sodium sulphate and the solvent was evaporated using rotary evaporator at 40–50 °C to dryness. The resulting crude extract of each solvent used was weighted and kept in the deep freezer until evaluation.

Direct effect on eggs of T. urticae:

The direct effect on the egg stage was studied by confining two females on leaf discs for 24 hr., and then removed. These discs with deposited eggs (24 hr old)

were sprayed with different concentrations from each extract using a glass atomizer. Each test contained 5 concentrations and each concentration had 5 replicates (20 eggs/ replicate). Hatchability of eggs on each concentration was recorded 7 days after treatment.

Direct effect on adult females of T. urticae:

Adult females of *T. urticae* were confined on the lower surfaces of detached raspberry leaves (3 cm in dia.) while the upper surfaces were placed on cotton saturated with water. Mites were sprayed with different extracts using a glass atomizer. Each test contained 5 concentrations and each concentration had 5 replicates (20 females/replicate). Mortality was recorded 48 hr. after application. In every test, a control was included.

Repellency and oviposition deterrence:

Raspberry leaf discs (5 cm in dia.) were placed with the lower surface upwards in a Petri-dish lined with moist cotton wool. One half of each disc was treated separately with LC₅₀ concentration of each extract, while the other served as a control. Ten adult females of *T. urticae* were then introduced into the middle of each leaf disc. Ten replicates of leaf discs were used per extract. Orientation of the females *T. urticae* on treated and control discs were recorded 3, 5, 24 and 48 hr. after treatment. The number of eggs laid on both sides were recorded after 48 hr. The repellency was calculated according to Lwand *et al.*, (1985).

Direct effect on predacious mite:

Adult females of the predator A. zaheri were sprayed with different concentrations from different extracts using glass atomizer. Females were confined on the lower surfaces of detached raspberry clean leaves (5 cm in dia.) while the upper surfaces were placed on cotton saturated with water. Each test contained 5 concentrations and each concentration had 5 replicates (20 females/replicate). In every test, a control was included.

Corrected mortality counts according to Abbott's formula (1925) were statistically analyzed by Finney (1971). The toxic index of each extract was determined according to Sun (1950).

RESULTS AND DISCUSSION

The data presented in Tables (1 and 2), show the relation between the percentage of mortality and concentrations of various extracts of L. raetam, S.

marianum and F. indica on eggs and adult females of T. urticae. The different plant extracts tested appeared to be more variable in their effects. The data obtained in tables 1 and 2 show that the toxicity effect of diethyl ether extract of L. raetam on both stages egg and adult were very close ($LC_{50}=0.0034$ and 0.0036 gm/ml), respectively.

TABLE (I)Toxicity of some plant extracts against egg stage of *T. urticae*.

	LC ₅₀	LC ₉₀	Clono	Toxicity index at:		
Extract	(gm/ml)	(gm/ml)	Slope	LC ₅₀	LC ₉₀	
	Lygos raetam					
Hexane	0.3702	1.1350	2.63	0.93	0.67	
Diethyl ether	0.0034	0.0111	2.51	100	68.83	
Ethyl acetate	0.0050	0.0076	6.81	69.15	100	
Ethanol	0.1160	0.2545	3.75	2.96	3.0	
	Silybum marianum					
Hexane	0.0113	0.0222	4.23	61.82	86.74	
Diethyl ether	0.0070	0.0192	2.91	100	100	
Ethyl acetate	0.0262	0.2572	1.29	26.67	7.48	
Ethanol	0.0824	0.2041	3.25	8.48	9.42	
	Fagonia indica					
Hexane	0.0018	0.0042	3.52	100	100	
Diethyl ether	0.0529	0.1035	4.39	3.42	4.05	
Ethyl acetate	0.0152	0.0314	4.05	11.92	13.33	
Ethanol	0.1976	0.7152	2.29	0.92	0.59	

Based on the LC₅₀ value it can be arranged for egg stage in the following descending order of effectiveness: Diethyl ether, ethyl acetate, ethanol and hexane; while in case of adult females, toxicity can be arranged in a descending order as follows: Ethyl acetate, diethyl ether, ethanol and hexane. The data obtained in tables 1 and 2 show that the various extracts of *S. marianum* were the most toxic to adult females than the egg stage, while the toxicity effect of the extracts hexane, diethyl ether and ethyl acetate of *F. indica* on both stages egg and adult were very close (LC₅₀= 0.0018, 0.0529 and 0.0152 gm/ml and LC₅₀= 0.0023, 0.0559 and 0.0109 gm/ml, respectively).

Results indicated also, that the various extracts of 3 plants were more toxic to adult stage than the egg stage. These results agree with that of Dimetry et al., (2000) who reported that the petroleum ether extracts of C. longa L., N. physaloides L. and alcohol extract of D. viscose L. were the most potent tested against adult females of T. urticae. The egg stage was less susceptible to the different extracts tested. Amer et al., (2005) reported that the adult females of T. urticae were

sensitive to crude ethyl acetate extract of *Francoeria crispa* (Forssk) than the egg stage, as well as the egg stage of *T. urticae* was less susceptible to the different extracts of *Capparis aegyptia* leaves and fruits (Hussein *et al.* 2005).

TABLE (II)

Toxicity of some plant extracts against adult females of T. urticae

Extract	LC50	LC90	Slone	Toxicity index at:			
	(gm/ml)	(gm/ml)	Slope	LC ₅₀	LC_{90}		
	Lygos raetam						
Hexane	0.0687	0.1751	3.15	1.09	0.81		
Diethyl ether	0.0036	0.0073	4.24	20.60	19.32		
Ethyl acetate	0.00075	0.00141	4.70	100	100		
Ethanol	0.0295	0.0574	4.43	2.54	2.46		
	Silybum marianum						
Hexane	0.0022	0.0036	5.99	100	100		
Diethyl ether	0.0037	0.0080	3.88	58.45	44.72		
Ethyl acetate	0.0056	0.0100	5.02	39.28	35.67		
Ethanol	0.0155	0.0330	3.92	14.03	10.80		
	Fagonia indica						
Hexane	0.0023	0.0040	5.51	100	100		
Diethyl ether	0.0559	0.1633	2.75	4.19	2.44		
Ethyl acetate	0.0109	0.0201	4.83	21.41	19.83		
Ethanol	0.0145	0.1577	1.24	16.10	2.53		

Repellency and oviposition deterrence:

Table (3) show that females of *T. urticae* preferred the untreated part of the leaves to feed and deposit eggs. After 48 hours of exposure the females deposited eggs on the treated part, the average number of eggs varied according to various extracts. A higher percentage of repellency was recorded in various extracts of *S. marianum* and *F. indica* (86.60 – 98.23 % and 85.53 – 97.0 %), respectively. While percent repellency decreased with extracts of *L. raetam* (55.81 – 80.84 %). Mansour and Ascher (1983) revealed that neem seed kernels prepared from various solvents strongly repelled the females of *Tetranychusa cinnabarines* (Boisd). Other plants such as *F. crispa* showed similar remarkable repellency against *T. urticae*. (Amer et al., 2005)

Toxicity effect on adult females of predacious mite A. zaheri.

The data obtained in Table (4) show that adult females of A. zaheri were more sensitive to ethyl acetate extract of L. raetam (LC₅₀ = 0.00079 gm/ml) while less susceptible to ethanol extract of the same plant (LC₅₀ = 0.2944 gm/ml). From tables (2 and 4) the toxicity effect of ethyl acetate extract of L. raetam on both adult

females of *T. urticae* and predacious mite *A. zaheri* were very close ($LC_{50} = 0.00075$ and 0.00079 gm/ml, respectively).

TABLE (III)

Relative percentage distribution and oviposition of *T. urticae* on treated leaf discs with different extract of *L. raetam*, *S. marianum* and *F. indica* at LC₅₀ conc.

[% Di	Distribution of mites on No. of eggs/fer			gs/female	%	
Extract	treated leaf part after:			after 48 hr.		Repellency	
	3 h	5 h	24 h.	48 h.	Ť	C	
	Lygos raetam						
Hexane	2	4	12	30	1.14	5.95	80.84
Diethyl ether	5.56	5.56	25.93	44.44	1.9	4.31	55.81
Ethyl acetate	1.82	1.82	22.22	29.10	1.5	5.4	72.04
Ethanol	3.77	3.77	7.55	43.40	1.04	4.09	74.57
	Silybum marianum						
Hexane	3.23	4.84	5	5	0.12	6.78	98.23
Diethyl ether	0	2.08	2.13	10.64	0.91	6.79	86.60
Ethyl acetate	4.76	4.92	3.39	6.78	0.14	3.69	96.21
Ethanol	0	0	5.56	17.65	0.33	5.15	93.59
	Fagonia indica						
Hexane	2	4	6	10	0.6	5.1	88.24
Diethyl ether	1.54	0	13.85	37.25	0.17	5.66	97.0
Ethyl acetate	2.82	1.41	18.31	34.29	3.11	3.11	85.53
Ethanol	3.08	3.08	18.46	32.81	3.98	3.98	92.71

T = treated C = control

TABLE (IV)
Toxicity of some plant extracts against adult stage of the predactious mite A. zaheri

	LC ₅₀	LC90	Slope	Toxicit	Toxicity index at:	
Extract	(gm/ml)	(gm/ml)	Stope	LC ₅₀	LC ₉₀	
		Lyg	zos raetan	η		
Hexane	0.0824	0.1711	4.04	0.96	0.81	
Diethyl ether	0.0076	0.0121	6.34	10.37	11.38	
Ethyl acetate	0.00079	0.00138	5.27	100	100	
Ethanol	0.2944	0.9890	2.43	0.27	0.14	
	Silybum marianum					
Hexane	0.0036	0.0053	7.69	100	100	
Diethyl ether	0.0062	0.0105	5.50	58.70	50.38	
Ethyl acetate	0.0091	0.0179	4.36	39.54	29.50	
Ethanol	0.0356	0.1065	2.69	10.13	4.97	
	Fagonia indica					
Hexane	0.0035	0.0123	2.34	100	100	
Diethyl ether	0.0840	0.1703	4.17	4.17	7.22	
Ethyl acetate	0.0204	0.0265	11.21	17.19	46.45	
Ethanol	0.0420	1.4705	0.83	8.33	0.84	

Generally the data from tables (2 and 4) indicated that the all extracts of the 3 plants were more toxic to adult female of *T. urticae* than the female of *A. zaheri*. Momen and Amer (1994) reported that the Lupin extract was slightly toxic to eggs and females of the predator *A. barkeri*. Essential oil of mint, *Mentha virdis* was more toxic to female of *T. urticae* than to female of Phytoseiid predators, *Amblyseius yosefi*, *A. zaheri*, *P. finitimes*, *A. barkeri*, *A. deleomi* and *T. athiasae* (Momen *et al.*, 2001).

Take into consideration the combined effects of these extracts on the phytophagous mite *T. urticae* and the predator *A. zaheri* (Table 5), the ethanol extracts of *L. raetam, S. marianum* and *F. indica* were 9.98, 2.30 and 2.90 times more toxic to the female of *T. urticae* than to the female of the predator *A. zaheri*. Thus in a habitat where the predator is associated with phytophagous mites it is necessary to apply the least toxic material to the predator and the most efficient to the prey.

TABLE (V)

Evaluation of the acaricidal properties of some plant extracts against adult stages of

Turticae and the predactions mite A zaheri

	T. urticae	A. zaheri	No. of folds compared				
Extract	LC ₅₀ (gm/ml)	LC ₅₀ (gm/ml)	with the predator A. zaheri				
	Lygos raetam						
Hexane	0.0687	0.0824	1.20				
Diethyl ether	0.0036	0.0076	2.11				
Ethyl acetate	0.00075	0.00079	1.05				
Ethanol	0.0295	0.2944	9.98				
	Silybum marianum						
Hexane	0.0022	0.0036	1.64				
Diethyl ether	0.0037	0.0062	1.68				
Ethyl acetate	0.0056	0.0091	1.63				
Ethanol	0.0155	0.0356	2.30				
	Fagonia indica						
Hexane	0.0023	0.0035	1.52				
Diethyl ether	0.0559	0.0840	1.50				
Ethyl acetate	0.0109	0.0204	1.87				
Ethanol	0.0145	0.0420	2.90				

SUMMARY

Three wild plants, Lygos raetam, Silybum marianum and Fagonia indica were successively extracted with four different solvents. These extracts were tested for their toxicity and repellency against the two-spotted spider mite, Tetranychus

urticae Koch and their direct toxicity to adult female of the predacious mite Amblyseius zaheri Yousef and El-Borolossy. The adult females of T. urticae were sensitive to various extracts of three plants than the egg stage. A higher repellency was recorded in various extracts of S. marianum and F. indica and decreased with extracts of L. raetam.

Laboratory studies indicated that the adult female of the predacious mite A. zaheri was more sensitive to ethyl acetate extract of L. raetam while less susceptible to ethanol extract of the same plant. All various extracts of the three plants were more toxic to adult female, T. urticae than the female of A. zaheri.

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