# THERMAL AND PHOTODECOMPOSITION OF FENITROTHION AND MALATHION INSECTICIDES Mahmoud, A. A.

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## **ABSTRACT**

The effect of exposure to heat, UV-rays and sunlight on the degradation of fenitrothion and malathion were studied. It was found that fenitrothion insecticide residues was more persistence than malathion insecticide residues and both showed progressive loss as temperature and prolongation of exposure period increased, this indicates that the interval between successive sprays should be shorter at high temperature and vice versa. Photodegradation with UV-rays was positively correlated with exposure period and chemical structure of insecticide, malathion degraded faster than fenitrothion in this respect. Sunlight was more effective than UV-rays in accelerating the photodecomposition of fenitrothion and malathion residues. **Keywords:** Thermal – Photodecomposition – Fenitrothion- Malathion

#### INTRODUCTION

The decrease of deposits or residues of insecticides which occurred under field conditions, take place as a result of different environmental factors such as heat, light, air and moisture. The ultraviolet component of sunlight, which varied from 240 to 400 nm, is responsible for pesticide photolysis in the environment (Ohkawa *et al.*, 1974 and Abdel-Razik *et al.*, 1982-a). Therefore, it is an important prerequisite before recommending the use of any insecticide to study how far it is affected by these factors. Direct sunlight is more effective thane UV-rays in accelerating the photodecomposition (Soliman, 1994).

The present study was undertaken to investigate the effect of temperature, UV-rays (short wave 254 nm) and direct sunlight on the stability and degradation of the active ingredient of fenitrothion and malathion insecticides.

# **MATERIALS AND METHODS**

The role of heat, ultraviolet ray and sunlight exposure on the degradation of the active ingredients of fenitrothion and malathion were studied. One milliliter of 0.1% pure active ingredient purity source 98%, 99.5% from Dr.Ehrenstorfer of each of fenitrothion and malathion in n-hexane (1000 ug) was spread as thin film as uniformly as possible on the surface of uncovered Petri dish. The n-hexane solvent was left to dry at room temperature and the dishes were divided and subjected to different treatments.

The 1st set of treated petri-dishes were exposed to 20, 30, 40 and 50°C for 1, 4, 12, 24, 48, 96, and 192 hours at each temperature respectively, inside a dark electric oven provided with temperature regulating system. The second set was exposed to the short wave of an ultraviolet lamp (254 nm) at a distance of 12 cm for 1, 2, 4, 8, and 12 hours. The third set was exposed to

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normal and direct sunlight regime on July 2008 for 1, 4, 12, 24 and 48 hours, dominating temperature ranged between 35 and 37  $^{\circ}$ C.

Residues of tested insecticides which were remained on exposed surface were quantitatively transferred to standard glass stopper test tubes using n-hexane and the solvent was evaporated to dryness. An internal standard (ditalimphos 0.3 mg/L) was added to the test tubes before quantitative analysis and the residues were ready for determination by gas liquid chromatography (nitrogen-phosphorus detector – NPD).

Detection of residues and confirmation of the presence of fenitrothion and malathion was done by using gas chromatograph equipped with two chromatographic columns of different polarities and two nitrogen phosphorus detectors.

# GC (6890)- NPD instrument conditions

- Injector temp. = 225 °C
- Detector temp. A& B = 280 °C
- Flow rate of hydrogen 3.5 ± 0.1 ml/min
- Flow rate of air 110 ml/min
- Column head pressure 75 kpa
- Carrier gas nitrogen 2.5 ml/min
- Flow rate of carrier gas + detector auxiliary gas 25 ml/min
- Septum purge 5 ml/min, split vent 70 ml/min
- Splitless time 0.7 min

-Oven temperature program:

Level	Rate (°C / min)	Temp. ( °C)	Time (min.)
1	0	90	2
2	20	150	3
3	6	270	15

Total run time: 35 min.

#### Columns used:

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Column type	Length (meters)	I.D.* (mm)	Film thickness (um)
	(ineters)	(111111)	(uiii)
PAS-5	25	0.320	0.52
DP-1701P	25	0.320	0.25

<sup>\*</sup> Internal Diameter

# - Relative Retention time of the analyzed insecticides on the two used columns:

	Columns type				
Pesticides	PA	S-5	DP-1701P		
	RT* (min.)	RRT**	RT (min.)	RRT	
Fenitrothion	16.980	0.853	17.087	0.869	
Malathion	17.255	0.866	16.889	0.859	

<sup>\*</sup> Retention Time in minutes.

# **RESULTS AND DISCUSSION**

Data in tables (1) and (2), show that the persistence of fenitrothion and malathion residues, respectively, on glass surface were influenced by temperature degrees and period of exposure. The percentage of loss of

<sup>\*\*</sup> Relative Retention Time, relative to internal standard.

fenitrothion was 96.4, 60.1, 24.9 and 1.5 after 192 hours exposure at 50°C, 40°C, 30°C and 20°C, respectively. No loss of fenitrothion occurred till 96 hours, 1 hour exposure at 20°C and 30°C respectively while fenitrothion started to decompose only after 1 hour exposure at 40°C and 50°C. For malathion the data indicated that the percentage of loss was 97.5, 64, 28.5 and 8.9 after 192 hours exposure at 50°C, 40°C, 30°C and 20°C respectively. No loss of malathion occurred till 24 hours exposure at 200C, while malathion started to decompose only after 1 hour exposure at 30°C, 40°C and 50°C. In general, fenitrothion was more persistent than malathion and they suffered progressive loss as temperature and prolongation of the exposure period increased.

From the practical point of view in the pest control program, it can be recommended to use the two tested insecticides in area of dominant low temperature. Moreover, this clearly showed that the interval between successive sprays should be shorter at high temperature and vice versa.

The role of temperature in increasing degradation of insecticide residues were studied and confirmed by several investigators i.e. Abu-Zahw et al. (1988), Abdel-Baki et al. (1999) and Hegazy et al. (2001).

The effect of UV-rays on fenitrothion and malathion residues were tabulated in table (3). The results indicate that the rate of fenitrothion and malathion degradation insecticides was varied according to their chemical structure and time of exposure to UV-rays. malathion was degraded faster than fenitrthion in this respect.

The most important metabolites for fenitrothion are dimethylfenitrooxon and 3-methyl-4-nitrophenol. The degradation for malathion is by oxidative desulfuration, leading to the formation of malaoxon; malathion and malaoxon are hyrolysed and thus detoxified by caboxylesrases.as referenced by (e-Pesticide Manual) (2003).

Table (1): Effect of temperature on the persistence of fenitrothion insecticide

Time of	20°C		30°C		40°	C 2°	50°C	
exposure (hour)	ug.	% Loss	ug.	% Loss	ug.	% Loss	ug.	% Loss
0	1000.00	00.00	1000.00	00.00	1000.00	00.00	1000	00.00
1	1000.00	00.00	1000.00	00.00	950.7	4.9	926.3	7.4
4	1000.00	00.00	975.1	2	916.2	8.4	896.4	10.4
12	1000.00	00.00	950.2	4.9	865.2	13.5	851.3	14.9
24	1000.00	00.00	930.6	6.9	815.2	18.5	720.2	27.9
48	1000.00	00.00	890.3	10.9	725.3	27.5	610.2	38.9
96	1000.00	00.00	835.1	16.5	586.9	41.3	320.3	67.9
192	984.6	1.5	750.1	24.9	399.1	60.1	35.8	96.4

Table (2): Effect of temperature on the persistence of malathion insecticide

Time of exposure	20°C		30°C		40°C		50°C	
(hour)	ug.	% Loss	ug.	% Loss	ug.	% Loss	ug.	% Loss
0	1000.00	00.00	1000.00	00.00	1000.00	00.00	1000.00	00.00
1	1000.00	00.00	990	1.000	930	7	903	9.7
4	1000.00	00.00	961	3.9	902	9.8	815.3	18.5
12	1000.00	00.00	950	5	850	15	710.4	28.9
24	1000.00	00.00	910	9	799	20.1	620.1	37.9
48	991	0.9	860	14	660	34	450	55
96	960	4	821	17.9	520	48	90	91
192	910.1	8.9	715	28.5	360	64	25.3	97.5

In general, photodegradation with UV-rays was positively correlated with the exposure period. These results are in accordance with those obtained by El-Sayed *et al.* (1980), Abdel-Razik *et al.* (1982-a and b), Hegazy *et al.* (1982), Abu-Zahw *et al.* (1988), Abdel-Baki *et al.* (1999), Hegazy *et al.* (2001), V Kannan *et al* (2005), Kaylynn Newhart. (2006) and Rusmidah Ali *et al* (2008).

Table (3): Effect of UV-rays (254 nm) on the persistence of fenitrothion and malathion insecticides

Time of exposure	Fenitrothion		Malathion		
(hour)	ug.	% Loss	ug.	% Loss	
0	1000.00	00.00	1000.00	00.00	
1	960	4	914	8.6	
2	881.0	11.9	825.3	17.5	
4	750.1	24.9	712.7	28.7	
8	681.3	31.9	610.1	38.9	
12	620.2	37.9	550.1	44.9	

Data presented in table (4) show that the percentage of loss for fenitrothion and malathion were 0.2 and 28.3% respectively after one hour of exposure to direct sunlight. The decomposition percentages of both insecticides rapidly increased to 97.9 and 99.1% respectively after 24 hours of exposure. The results show that malathion residues was degraded faster than fenitrothion.

These results are similar to those obtained by El-Sayed *et al.* (1980), Abdel-Razik *et al.* (1982 - a and b), Hegazy *et al.* (1982), Abu-Zahw *et al.* (1988), Abdel-Baki *et al.* (1999), Hegazy *et al.* (2001) and Balakrishnan *et al.* (2002).

Considering the time of exposure to sunlight, it was obvious that 24 hours exposure seemed to be nearly decompose completely the two insecticides. Also, sunlight was more effective than UV-rays in accelerating the photodecomposition of fenitrothion and malathion. This could be due to thermal, evaporation and light intensity considerations (Hegazy *et al.*, 2001).

Table (4): Effect of sunlight on the persistence of fenitrothion and malathion insecticides

Time of exposure	Fenitr	othion	Malathion			
(hour)	ug.	% Loss	ug.	% Loss		
0	1000.00	00.00	1000.00	00.00		
1	998	0.2	717	28.3		
4	611	38.9	340	66.0		
12	210.2	78.9	30.9	96.9		
24	20.2	97.9	8.8	99.1		
48	10.2	99	N.D.*	100		

\*N.D. not detected

## **REFERENCES**

- Abdel-Baki, M. A.; Hegazy, M.E.A.; Shady, M.F.A. and Shokr, S.A. (1999).Thermal and photodecomposition of some organophosphorus pesticides. Egypt J. Agric. Res., 66(4): 760-771.
- Abdel-Razik, M.; Hegazi, M.E.A. and El-Sayed, M.M. (1982-a). Thermal and photo-decomposition of the pyrethroid permethrin. Proc. Egypt's Nat. Conf. Ent., 11:753-761.
- Abdel-Razik, M.; Hegazi, M.E.A. and El-Sayed, M.M. (1982-b). Persistence of some organophosphorus pesticides under different degrees of temperature, ultra violet rays (2357 A0) and direct sunlight. Proc. Egypt's Nat. Conf. Ent., 11: 762-77
- Abu Zahw, M.M.; Hegazy, M.E.A.; Dogheim, S.M.; Almaz, M.M. and Shahin, A. (1988). Thermal and photodecomposition of fenitrothion and esfenvalerate insecticides.Bull. Ent. Soc.Egypt, Econ. Ser., 17: 193-198.
- Balakrishnan, V. K., Buncel, E., Loon, G. W. van (2002) .Micellar catalyzed degradation of fenitrothion, an organophosphorus pesticide, in solution and soils. Department of Chemistry, Queen's University, Kingston, Ontario K7L 3N6, Canada. Environmental Science & Technology.,21: 152-159.
- El-Sayed, M.M.; Shaaban, A. M.; Zidan, H. and Abu-Zahw, M. M. (1980). Persistence and biological transformation of Curacron, Cyolane and Reldan insecticides as affected by temperature, UV-raus and sunlight. Proc. 1<sup>st</sup> Conf. P1. Prot. Res. Ins. Dokki, Cairo, Egypt.PP. 361-380.
- Hegazy, M. E. A.; Abdel-Razik, M. and El-Sayed, M. M. (1982). Thermal and photo-decomposition of monitor and pirimiphos-methyl organophosphorus pesticides. Proc. Egypt's National Conf. Ent. Vol II: 743-751.
- Hegazy, M.E.A.; Youssef, M.M.M.; Abdel-Fattah, A.M.; Shokr, S.A. and Ahmed, N.S. (2001). Photosenitizing or stabilizing influence on two organophosphorus insecticides under ultraviolet rays and direct sunlight exposures. Egypt. J. Agric. Res. 79(2): 499-509.
- Ohkawa, H.; Mikant, N. and Miyamoto, J. (1974). Photodecomposition of Sumithion (o,o-dimethyl-o-(3-methyl-4-nitrophenyl)-phosphoro- thioate). Agr. Biol. Chem., 38(11): 2247-2255.

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- Soliman, M.M. (1994). Efficiency of some insecticides against leguminous pod borer Etiella zinkenella treitschke on cowpea with special reference to pesticide residues. M.Sc. Thesis, Fac. Of Agric., Cairo Univ.
- The British Crop Protection Council(2003). The e-Pesticide Manual (Twelfth Edition) Version 2.2.
- V Kannan and V Vanitha (2005) influence of assay medium on degradathion of malathion by serratia marcescens. Indian journal of biotechnology 4: 277-283.
- KayLynn Newhart. ( 2006) Environmental Fate of Malathion. California Environmental Protection Agency Department of Pesticide Regulation Environmental Monitoring Branch 67(4):1255-1266,
- Rusmidah Ali and Siti Habsah Hassan (2008) Degradation studies on paraquat and malation using TiO2/ZnO based photocatalyst. The Malaysian Journal of Analytical Sciences 12(1):1544-1556.

التحلل الحرارى والضوئى لمتبقيات مبيدى الفنتروثيون والملأثيون على على محمود المعمل المركزى لتحليل متبقيات المبيدات والعناصر الثقيلة في الأغذية

تم در اسة تأثير درجات الحرارة والأشعة فوق البنفسجية والتعرض لضوء الشمس لمدد مختلفة على متبقيات مبيدى الفنتروثيون والملأثيون. وقد دلت النتائج على أن مبيد الفنتروثيون أكثر ثباتاً من مبيد الملأثيون كما أنه كلما زادت درجة الحرارة أو فترة التعرض لضوء الشمس كلما زادت نسبة تحطم مبيدى الفنتروثيون والملأثيون مما يجعل من الضروري تقليل الفترة الزمنية بين تكرار الرش بهذين المبيدين على المحاصيل المختلفة. كما اتضح تأثير هذين المبيدين بالأشعة فوق البنفسجية حيث يزداد بزيادة فترات التعرض. وقد وجد أن مبيد الملأثيون يتحطم بصورة أكثر من مبيد الفنتروثيون نتيجة التعرض للأشعة فوق البنفسجية. وقد ثبت أن التعرض لضوء الشمس يؤدى إلى اسراع معدل التحطم الصوئلهذين المبيدين بشكل أكبر من تعرضهما للأشعة فوق البنفسجية. الكلمات الدالة: التحلل الحراري والضوئي الفنتروثيون — الملأثيون.

كلية الزراعة – جامعة المنصورة مركز البحوث الزراعية قام بتحكيم البحث أ.د / على على عبد الهادى أ.د / اشرف محمود حسن المرصفي