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EMBRYONIC TOXICITY OF INSECTICIDE SUMITHION 50 EC AND HERBICIDE FUSILADE S IN PHEASANTS AFTER INDIVIDUAL OR COMBINED ADMINISTRATION

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The purpose of this work was to determine the individual and combined effects of insecticide Sumithion 50 EC (50% fenitrothion) and herbicide Fusilade S (12.5% fluazifop-P-butyl) on the development of pheasant embryos. Eggs were treated by injection of various concentrations of pesticides into the air space on day 12 of incubation. Pathological examination of embryos was carried out on day 23 of the hatching period. Mortality rate, body weight data and morphological alterations were evaluated after the macroscopic examination. The skeletal staining method was used to detect deformities. The two pesticides used in combination moderated the toxic/teratogenic effects of individual treatment.

Key words: Teratology, pesticide, interaction, pheasant, fenitrothion, fluazifop-P-butyl

The avian embryo is sensitive to environmental contaminants. The toxic and teratogenic potential of various chemicals in avian (quail, pheasant, chicken, etc.) embryos has been studied by many authors (Dareste, 1891; Féré, 1901; Deakin and Robertson, 1933; Bowman, 1967; Hoffman and Albers, 1984; Somlyay et al., 1992). Researchers have demonstrated the sensitivity of avian embryos to pesticides, and emphasised the potential use of chick and pheasant embryos as a screening model (Hoffman, 1990).

Since 1988, in order to obtain an authorisation for use in Hungary, a newly developed pesticide formulation has to be tested on chick or pheasant embryos to detect its ecotoxic, embryotoxic and teratogenic effects (Várnagy, 1989).

Teratogenicity studies generally involve individual chemical exposure, but in the plant protection practice various pesticides can be used simultaneously or consecutively within a short period of time on cultivated land (Várnagy et al., 1996).

The aim of this study was to determine the individual and combined effects of insecticide Sumithion 50 EC (50% fenitrothion) and herbicide Fusilade S (12.5% fluazifop-P-butyl) on late embryonic development of the pheasant.

On the day of injection (day 12 of incubation) the developmental stage of the pheasant embryo was characteristic: the embryo had already passed the histodifferentiation period, the main organs had been formed, and growth had been finished (Nagy and Ernhaft, 1962).

The pheasant was chosen as a model animal because the above-mentioned two pesticides can easily contaminate pheasant eggs on lands.

Materials and methods

Materials

Fusilade S (12.5% fluazifop-P-butyl): liquid herbicide, slightly toxic [acute oral LD₅₀ for rats: 2000 mg/kg, Sumithion 50 EC (50% fenitrothion): liquid insecticide, slightly toxic (acute oral LD₅₀ for rats: 1400 mg/kg) (Ocskó and Molnár, 1996)]. Applied concentrations (solvent: distilled water): Fusilade S: 0.1; 1.0; 10.0%; Sumithion 50 EC: 0.033; 0.33; 3.33%. The middle concentration corresponded to that usually applied in the chemical plant protection practice. Designation of the groups: control and treated groups I, II and III, respectively, in the order of increasing concentration.

Methods

Eggs: fertile, fresh pheasant eggs (*Phasianus colchicus mongolicus et torquatus*), origin: Pusztaberény, Hungary.

Incubation: in a Ragus type incubator under conventional conditions.

Treatment: on day 12 of incubation 0.1 ml emulsion (final volume) per egg was administered directly into the air space with an Ovijector automatic injector. Before the treatment the egg shell was bored through, then, after the injection, it was sealed with paraffin (Clegg, 1964).

Processing: pathological examination was carried out on day 23 of incubation. Microscopic evaluation was done by stereomicroscopy of skeletal preparations after bone staining (Dawson, 1926).

Statistical analysis: Student's *t*-test (Finney, 1972).

Results

The middle and the highest dose level of Sumithion 50 EC significantly reduced the body weight of embryos. It caused high embryo mortality at the third concentration level (Group III). The incidence of developmental anomalies was increased among the embryos at all doses, but the most remarkable numerical rise was detected at the middle treatment level (Tables 1 and 4).

Table 1

Toxicity of Sumithion 50 EC to pheasant embryos after injection on day 12 of incubation

Test material and concentration (%)	Body mass of the embryo (g) Average \pm S.D.	Number of samples	Embryonic death total/after treatment	Embryos showing developmental anomalies (n)	Live embryos No.	%
Control	16.99 \pm 1.37	20	5/4	1	15	75.00
0.033 I	16.30 \pm 1.50	19	2/1	4	17	89.47
0.333 II	15.68 \pm 1.35 ^b	24	6/5	7	18	75.00
3.333 III	13.85 \pm 1.39 ^b	22	18/17	4	4	18.18

^b = P < 0.01

Fusilade S reduced the average body mass significantly at the medium (II) dose level and caused 100% embryonic mortality in the highest (III) dose. Malformations were observed only at the first (I) level. No skeletal malformations were detected (Tables 2 and 5).

Table 2

Toxicity of Fusilade S to pheasant embryos after injection on day 12 of incubation

Test material and concentration (%)	Body mass of the embryo (g) Average \pm S.D.	Number of samples	Embryonic death, total/after treatment	Embryos showing developmental anomalies (n)	Live embryos No.	%
Control	16.20 \pm 2.02	20	5/1	0	15	75.00
0.1 I	16.32 \pm 2.51	19	1/0	3	18	94.44
1.0 II	14.45 \pm 2.67 ^a	19	5/2	0	14	73.68
10.0 III	–	18	18/13	–	0	0.00

^a = P < 0.05

Combined administration of Sumithion 50 EC and Fusilade S caused a significantly lower average body mass at the two higher concentrations (II, III). The highest embryo mortality was caused by the highest dose (III) and developmental anomalies appeared after treatment with the two higher dose levels (II, III). These typical alterations were demonstrated also in earlier teratological trials using different pesticides (Várnagy, 1995). Bone malformations were not detected but the development stage of bones did not correspond to the normal status on the basis of skeletal staining intensity (Tables 3 and 6).

Table 3

Combined toxicity of Sumithion 50 EC (Su) and Fusilade S (Fu) to pheasant embryos after injection on day 12 of incubation

Test material and concentration (%)	Body mass of the embryo (g) Average \pm S.D.	Number of samples	Embryonic death, total/after treatment	Embryos showing developmental anomalies (n)	Live embryos No.	%
Control	14.50 \pm 1.58	16	4/2	1	12	75.00
Su 0.33 + Fu 0.1 (I)	14.99 \pm 1.62	17	4/2	0	13	76.47
Su 0.33 + Fu 1.0 (II)	12.23 \pm 0.98 ^a	13	5/4	3	8	61.53
Su 0.33 + Fu 10.0 (III)	12.73 \pm 1.05 ^a	17	14/12	2	3	17.64

^a = P < 0.05

Table 4

Incidence of malformations in toxicity test of Sumithion 50 EC in pheasant embryos after injection on day 12 of incubation

Groups	Type of anomalies (number of cases)	Type of skeletal malformations
Control	(0)	–
I	cyllosis (4)	diminished intervertebral space
II	cyllosis (6), cyllosis and lordoscoliosis (1)	diminished intervertebral space
III	cyllosis (4)	adhesion of lumbar vertebrae

Table 5

Incidence of malformations in toxicity test of Fusilade S in pheasant embryos after injection on day 12 of incubation

Groups	Type of anomalies (number of cases)	Type of skeletal malformations
Control	(0)	–
0.1% I	cyllosis and lordoscoliosis (1), cyllosis and nanosomia (1)	–
1.0% II	cyllosis (1)	–
10.0% III	(0)	underdevelopment of the skeleton

Discussion

Fusilade S given alone caused 100% embryo mortality at the highest dose level but the same dose given simultaneously with Sumithion 50 EC (level II) moderated the embryotoxicity to 82.36%. The number of developmental anomalies was higher at the middle level of Sumithion 50 EC given alone, as compared

to the same treatment level given in combination with Fusilade S. Combined treatment with the two pesticides reduced the incidence of embryonic developmental anomalies while this phenomenon was highly expressed in the Fusilade S study.

Table 6

Incidence of malformations in combined toxicity test of Sumithion 50 EC (Su) and Fusilade S (Fu) in pheasant embryos after injection on day 12 of incubation

Groups	Type of anomalies (number of cases)	Type of skeletal malformations
Control	cyllosis (1)	–
Su 0.33% + Fu 0.1%	I (0)	–
Su 0.33% + Fu 1.0%	II lordoscoliosis (1), os frontale planum (1), cyllosis (1)	underdevelopment of skeleton
Su 0.33% + Fu 10.0%	III cyllosis and lordoscoliosis (1), cyllosis (1)	underdevelopment of skeleton

No skeletal malformations were found in embryos that received Sumithion 50 EC and Fusilade S simultaneously, though such malformations had been expected on the basis of the trial of Sumithion 50 EC which induced abnormalities in bone development.

The two pesticides applied in combination in this experiment under standardised circumstances represent a remarkable example of the antagonistic effect of chemicals.

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