

Releasable

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Atrazine Technical - Addition of Data to Files
Casewell #63
Shaughnessy #080803
Toxicology Branch
Registration Division

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Thru: O.E. Paynter, Ph.D.
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Recommendation

No action required.

Review

1. 28-Day Range Finding Study with Atrazine in Albino Mice - (Industrial Bio-Test, IBT # 8532-08868, 5/28/76, submitted by Ciba-Geigy on 6/2/77, Acc.#230303.)

Ninty Charles River mice weighing 23-29 g. were divided into 9 groups of 10 animals each (5 males, 5 females) and were fed either 0, 10, 30, 100, 300, 1000, 3000, 10000, or 30000 ppm Atrazine Technical in the diet for 28 consecutive days. Males were housed individually, whereas females were housed 5 per cage. Body weight was determined initially and at weekly intervals for the duration of the study. Food consumption data was collected, individually for males and on a group basis for females, during the 4 week study. Observations for abnormal reactions and mortality were made daily. Necropsies were performed on all animals.

Results

MTD = 1000 ppm

One male died at 300 ppm, 2 males died at 3000 ppm, 2 males and 5 females died at 10000 ppm and all animals died when fed 10000 ppm.

Toxic Signs: tremors and generalized weakness among animals fed 30000 ppm after 3 days of feeding - the response was more pronounced in females. The same reactions were seen among mice fed 10000 ppm after 7 days of feeding.

No adverse reactions were seen among animals feed 3000 ppm or less. Body weight gain and food consumption were normal in mice fed up to 3000 ppm.

Necropsy: unremarkable

Classification: Core-minimum data.

2. Acute Oral LD₅₀ of Technical Atrazine in the Rat: (Ciba-Geigy, Project #Siss 4569, 3/10/75, submitted by Ciba-Geigy on 6/2/77), Acc # 230303.

Seventy Tif. RAI rats, ranging in body weight from 160-180 g., were divided into 7 groups of 10 animals each (5 male, 5 female) and administered 600, 1000, 1290, 1670, 3170, 4640 or 6000 mg/kg of the test material by gavage. Animals were housed in groups of 5 and observed for signs of toxicity and/or mortality for a period of 14 days. Necropsies were performed on all animals.

Results

LD₅₀ = 1869 (1405-2487) mg/kg

Toxic Signs: sedation, dyspnea, exophthalmos, curved position and ruffled fur.

Necropsy: unremarkable

Classification: Core-minimum data

- 1) body weight and food consumption were not determined daily.

TOX CATEGORY:III

3. Acute Oral LD₅₀ of Technical Atrazine in the Mouse - (Ciba-Geigy, Project #Siss 4569, 9/7/77, submitted by Ciba-Geigy on 6/2/77), Acc # 230303.

Sixty Tif. MAG mice, ranging in body weight from 20-30 g., were divided into 6 groups of 10 animals each (5 male, 5 female) and administered 1670, 2780, 3590, 4640, 5200 or 6000 mg/kg of the test material by groups. Animals were housed in groups of 5 and observed for signs of toxicity and/or mortality for a period of 14 days. Necropsies were performed on all animals.

Results

LD_{50} = 3992 (3557-4479) mg/kg

Toxic Signs: sedation, dyspnea, curved or ventral position and ruffled fur.

Necropsy: unremarkable

Classification: Core-minimum Data

- 1) body weight and food consumption were not determined daily.

TOX CATEGORY: III

4. Acute Dermal LD_{50} of Technical Atrazine in the Rat - (Ciba-Geigy, Project # Siss 5663, 12/6/76, submitted by Ciba-Geigy on 6/2/77, Acc # 230303).

Twenty Five: RAIF(SPF) rats, ranging in weight from 180-200 g., were divided into 2 groups of 10 animals each (5 male, 5 female) and dermally administered 2150 or 3170 mg/kg of the test material as a 50% aq. suspension to intact skin. The material was allowed to remain in contact with the skin for 24 hrs. under an impervious cuff. Following the 24 hr. exposure the material was washed away and the animals observed for 14 days post exposure. Necropsies were performed on animals at the end of the observation period.

Results

LD_{50} > 3100 mg/kg (highest dose possible) - No deaths occurred.

Toxic Signs: none, no local irritation.

Necropsy: unremarkable

Classification: Core-minimum Data

- 1) since the results are clear at the highest dose level possible was administered, further testing would be pointless. The study adequately reflects dermal hazard.

TOX CATEGORY: III

5. Primary Eye Irritation of Technical Atrazine -
(Ciba-Geigy, Project # siss 5663, 11/24/76, submitted by
Ciba-Geigy on 6/2/77, Acc#230303)

100 mg. was inserted into the conjunctival sac of the left eye of each of six Himalayan rabbits. The treated eye of 3 of the 6 rabbits was flushed with 10 ml of water 30 sec. after treatment. Ocular irritation was recorded 1, 2, 3, 4, and 7 days past treatment.

Results

Unwashed Eyes: 0.0/110 at 1, 2, 3, 4 and 7 days
Washed Eyes: 0.0/110 at 1, 2, 3, 4 and 7 days.

Classification: Core-Minimum data

- 1) although readings were made on 3 unwashed eyes, these readings together with the 3 readings on the washed eyes indicate that the technical material is not an eye irritant.

TOX CATEGORY: IV

6. Primary Skin Irritation of Technical Atrazine -
(Ciba-Geigy, Project #siss 5663, 11/24/76, submitted by
Ciba-Geigy on 6/2/77, Acc#230303)

500 mg. of test material was applied to one abraded and one intact skin site on 6 Himalayan rabbits and held in contact with the skin under an impervious cuff for 24 hrs. Dermal irritation was scored at 24 and 72 hrs. by the method of Draize.

Results

P.I. = 0.2/8.0

Classification: Core minimum data

- 1) readings were not made on 2 intact and 2 abraded skin sites.

TOX CATEGORY: IV

7. Mutagenicity Screening of Pesticides in the Microbial System -
(Shirash, Moriya, Kato, Furuhashi, Kada; Mutation Research
10 (1976) 19-30)

A survey on the mutation induction capacity was made in the microbial system on 166 pesticides including 57 fungicides, 63 herbicides and 46 insecticides. The screening methods consisted of the rec-assay procedure, a sensitivity test utilizing H17 Rec + and M45 Rec - strains of *Bacillus subtilis*, as well as the reversion assays on plates utilizing auxotrophic strains of *Escherichia coli* (WP2) and *Salmonella typhimurium* (Ames series).

Rec Assay

Test strains (*B. subtilis* H17 Rec+ and M45 Rec-) were grown overnight in broth B-2. Two cultures were streaked on the B-2 agar plates and the "starting points" were covered with a 10 mm diameter paper disc containing 0.02 ml. of each sample. All plates were incubated for 24 hrs. at 37°C and the length of the inhibition zones measured.

Reversion Assay

0.1 ml. sample of each bacterial culture was spread on the surface of MB or 9BB agar and a 10 mm diameter paper disc containing 0.02 ml. of each sample was placed in the center of each plate. Each plate was incubated for 2 days at 27°C and the formation of revertant colonies around each disc observed. For samples showing positive effects the "soft agar" procedure was used. 0.1 ml of the bacterial culture and 0.1 ml of drug solution were added to 2nd soft agar at 45°C and poured into a petric dish containing 50 ml of 9BB agar. Colonies formed by incubation at 37° for 2 days were counted.

Media

Liquid broth B-2 (10g. meat extract, 10g. polypeptone and 5g. NaCl in 1000 ml. water pH 7.0 solidified with 1.5% Difco agar was used for the rec-assay. For reversion-assays with *E. coli* WP2 strains, a modified Vogel-Banner salts mixture was supplemented with 1% Difco broth and solidified with agar (MB agar). For the reversion assay with *Salmonella* strains, M9 minimal medium was supplemented with 1% Difco broth and 1mg/ml biotin (9BB agar).

A list of the pesticides assayed follows:

Do not type

TABLE 1

(methyl)acetanilide
 5,8,8-hexahydro-endo-1,4-oxa
 6-methyl-thio-1,3,5-triazine
 amate
 ale
 oxlamino-1,3,5-triazine
 (mate)
 dithiolate
 5,6-dinitro-p-toluidine
 diisopropyl phosphorothio-
 hamate
 chloride
 methylcrotonate
 (ate) ethylmethylthioacetaminate
 ate
 methyl acetanilide
 1-cyclohexene-1,2-dicarboximide
 2-cyclohexene-1,2-dicarboximide
 ate
 amide
 dicarbonate
 inate
 vinyl diethyl phosphate
 oxide
 hexformamidine
 1-ethylmethylurea
 dimethyl phosphorothiolothionate
 1,3-thiazole
 (1) vinyl dimethyl phosphate
 acetate
 ediazosulfonate
 elperidine
 (methyl) ethane
 6-methyl-thio-1,3,5-triazine
 oximidyl phosphorothionate
 mate
 (dimethylsulfamoyl)-aniline

Common name	Use	Chemical name
Dichlorocyclopentadiene (DDVP)	I	2,2-dichloro-5-vinyl-1,3-dioxolane
Dichlorodimethylsilane	I	3,3,3-trichloro-1,1,1-trimethyl-2,2,2-trifluoroethane
Dichlorodimethylsilane	I	2,6-dichloro-1,3,5-triazine
Dieldrin	I	1,2,3,4,10,10-hexachloro-6,7-epoxy-1,2,3,4,5,6,7,8,9-octachloro-1,1,1,1-tetrahydro-2,3,4-benzodioxin
Dimecholate	I	dimethyl (N-methylcarbamoylmethyl)phosphorothiolothionate
Dinitro	I	1,1-bis(p-chlorophenyl)ethanol
Dioxathion	I	S,S-1,1-dioxo-2,2-dithio-1,3-dithiolane-2-thiolate
Diphenamid	II	N,N-dimethyl-2,2-diphenylacetamide
Diphenylhydrazine	II	6,7-dihydro-5H-benz[5,6]cyclohepta[1,2-b:4,5-b']pyrazinediimide
Dithianon	I	2,3-dicyano-1,1-dithioanthraquinone
Duron	II	3-(3,1-dichlorophenyl)-1,1-dimethylurea
DMTP	I	S-[5-methoxy-2-oxo-2,3-dihydro-1,3,4-thiadiazolyl-(3)-methyl] phosphorothiolothionate
DFAS	I	poly[methylbis(thioacetamoyl)urea]
Durban	I	O,O-diethyl O-2,5,6-trichloro-2-pyridyl phosphorothionate
EAP	I	hexafluoroantimonyl trifluorophosphate
Edalox	I	quinoxaline-2,3-dithiolothionate
ESBP	I	S-benzyl ethyl phenylphosphorothiolothionate
ESTP	I	S-benzyl ethyl 1,1-dichloro-2-phosphorothiolothionate
ETP	I	tetraethyl S,S'-methylurea bis(phosphorothiolothionate)
ETM	I	N,N-ethylmethylthioacetaminate
Enthion	I	dimethyl 4-methylthio-2-ethyl phosphorothiolothionate
Entazon	I	3-benzylideneamino-1-phenylthiazole-2-thione
Erban	I	non dimethylthioacetaminate
Etape	I	N-(trichloromethyl)phthalimide
ETI	II	N-(2-hydroxyethyl)phthalazine
Heptachlor	I	1,1,1,2,2,4,4,4-heptachloro-3,4,5,6-tetrahydro-1,2,3,4-dioxin
Hinosan	I	O-ethyl dimethylphosphorothiolothionate
Hopside	I	o-chlorophenyl methylcarbamate
Hydrol	I	4-methylamino-2,5-xylol methylcarbamate
IRP	I	S-benzyl diisopropyl phosphorothiolothionate
Imidol	I	dimethyl S-(phosphorothiolothionate)
Ioxanocarbonate	II	1-cyano-2,6-dioxo-1,3,5-triazine
IPC	II	isopropyl isobutylphosphorothiolothionate
Karathane	I	potassium reaction mixture of 2,4-dinitro-6-ethylphenyl crotonate and 2,6-dinitro-1-ethylphenyl crotonate
Kasopoxin	I	2,2,2-trichloro-1,1,1-tris(p-chlorophenyl)ethanol
Kelthane	I	3-cyclohexyl-5,6-trimethylurea
Lenal	II	3-(3,1-dichlorophenyl)-1-methoxy-1-methylurea
Lanuron	II	calcium methane arsonate
MAC	I	3,5-xylol methylcarbamate
Malchal	I	non methane arsonate
MAF	I	S-1,2-bis(ethoxycarbonyl)ethyl dimethyl phosphorothiolothionate
Malathion	I	sodium methane arsonate
MAN	I	ethyl 1-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
MCPB (ethyl)	II	sodium 1-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
MCPB sodium	II	1-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
MCPN	II	2-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
MCP	II	1-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
MCPFA	II	1-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
MCP	II	potassium 2-(1-chloro-2,4-dichloro-5-hydroxyphenyl)acetate
Mercuriam	I	S-(N-ethoxycarbonyl-N-methylglucosaminyl) dimethyl phosphorothiolothionate
Methyl	I	2,1-xylol methylcarbamate
Methion	I	[ethylmethylthioacetaminate] copolymer
	I	[methylmethylthioacetaminate] copolymer

Result

The pesticides showing positive effect in broth rec-assay and reversion-assay were:

Captofol
Captan
2-Hydroazinoethane 1
(HEH)

Dexon
NBT
Vamidothion

Dichlovas,
Folpet
5-Nitro-1-naphthonitrile

Atrazine did not exhibit a mutation induction capacity.

Classification: Core Minimum Data.

William Greear