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item D Number	03501 Not Scanned
Author	Schultz, Donald P.
Corporate Author	
Report/Article 11tle	A Review of Literature on the Use of 2,4 -D in Fisheries
Journal/Book Title	
Year	1974
Manth/Day	March
Color	
Number of Images	94
Descripton Notes	Report No. FWS-LR-74-18. PB-235 457

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Schultz, D. P. and P.D. HARMAN

PRICES SUBJECT TO CHANGE

PB-235 457

A REVIEW OF THE LITERATURE ON THE USE OF 2, 4-D IN FISHEPIES

BUREAU OF SPORT FISHERIES AND WILDLIFE

March 1974

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National Technical Information Service

BIBLIOGRAPHIC DATA	1. Report No. FWS-LR=74=18	2,		PB 235 457
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	h Control Laboratory	r		11. Contract/Grant No.
P.O. Box 9 Warm Springs, GA	. 31830			
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2. Sponsoring Organization	Name and Address			13. Type of Report & Period Covered
U.S. Fish and Wi				Final
Division of Popu	lation Regulation Re	search	-	14.
15. Supplementary Notes	22. ·	· · ·		
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U.S. DEPARTMENT OF THE INTERIOR FISH AND WILDLIFE SERVICE

A Review of the Literature on the Use of

2,4-D in Fisheries

by

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March 1974

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The herbicide 2,4-D [2,4-(dichlorophenoxy)acetic acid] was adapted from terrestrial to aquatic use. Recommended treatment rates vary from 2.24 kg/ha to 22.4 kg/ha or even higher for submersed species.

Many formulations of 2,4-D are available, but the one used most commonly is the dimethylamine salt of 2,4-D (DMA-2,4-D). The ester formulations are also used but are 10 to 20 times as toxic to fish and other aquatic organisms as the dimethylamine salt.

A tolarance of 0.1 mg/liter has been issued for DMA-2,4-D that BEEUFS in putable water as a result of applications of UMA-2,4-D to ditch banks in the western United States. Several Federal agencies are presently pursuing the registration of 2,4-D for use on irrigation canal banks and for use in moving water. Since most of the research necessary for registration of 2,4-D has been done, it should be relatively inexpensive to complete the registration procedure.

ABSTRACT

HISTORY OF FISH CULTURAL USE

Research on 2,4-D compounds started during World War II, under war-time secrecy. The chemical has a growth-hormone effect at low concentrations (less than one ug/liter) and a herbicidal effect at concentrations of 0.1 mg/liter and greater.

The first recorded aquatic use of 2.4-D was to control waterhyacinth (Hildebrand, 1946). Saveral years later, Surber (1949) reported that 2,4-D was effective for control of emersed and marginal weeds. Since then, it has been used to control Elodea canadensis (Foret, 1967), Eurasian watermilfoil (Haven, 1963; Smith and Isom, 1967; Whitney, 1970, Wojtalik, et al., 1971) and Hydrilla verticillata (Kleinschmidt, 1969). It also has been tried as a control for Pithophora sp., alligatorweed, duckweed, waterstar grass (Lawrence, 1962), Nymphea odorata (Pierce, 1960), Brasenia spp. (Pierce, 1961); and, in combination with Endothal, Potamageton crispus (Pierce, 1969). Martin et al. (1957) noted tts use in improving duck mersh habitak. Frenk <u>et al.</u> (1464) reported that 2,4-D was useful for controlling water lily, lotus, spatterdock, and penny wort in irrigation canals but would not control Sago pondweed or American pond weed due to its lack of persistence in the soil. The latter authors reported that

2,4-D was somewhat effective for control of the following marginal weeds along irrigation canals: willow, sedge, smartweed, cattail, arrowhead, pickersl weed, bulrush, spikerush, knotgrass, southern watergrass, and needlerush. Bruns (1984, 1987) and Bruns and Clore (1958) investigated possible damage to crops from irrigation water treated with 2,4-D, while DeVaney (1967) reported efficacious dosages for weeds in farm ponds and lakes.

PHYSICAL AND CHEMICAL PROPERTIES

The herbicide 2,4-D [2,4-(dichlorophenoxy)acetic acid] has a molecular formula of $C_{\rm SH_6}Cl_2O_3$, a molecular weight of 221.0, a specific gravity, C, of 1.565 (30° C), a melting point of 135-138° C, a boiling point of 160° C at 0.4 mm Hg, and a vapor pressure of 0.4 mm Hg at 160° C (WSSA, 1970). It is a white, crystalline, odorless substance, and is soluble in many organic solvents but quite insoluble in water. Nelson and Faust (1969) determined the acid dissociation constant of 2,4-D by means of potentiometric titration. They found a pKa value of 2.73 at an ionic strength of 0.05 for 2,4-D.

The acid is prepared from 2,4-dichlorophenol and monochloroacetic acid in the presence of aqueous base. An aqueous solution of the dimethylamine salt (DMA) is prepared by addition of 40% dimethylamine solution to an aqueous slurry of 2,4-D at pH 8. The butoxyethanol ester is synthesized by the reaction of the free acid of 2,4-D with butoxyethanol (WSSA, 1970).

The free acid may be determined by infrared spectroscopy. Information on analytical methods is contained in the Residue Section of this report.

The teratogenic compound tetrachlorodioxin (dioxin) is found in small amounts in commercially prepared 2,4,5-trichlorophenoxyacetic acid, hence, analyses have been conducted on commercial preparations of 2,4-D. Huston (1972) reported three neutral impurities in commercially prepared 2,4-D which could interfere with gas-liquid chromatographic (glc) analyses for dioxins. She concluded that these conteminants were not dioxins nor deleterious. Plinmer and Klingebiel (1971) concluded that under field conditions, dioxins are unlikely products of the lower chlorinated phenols or phenoxyalkanoic acids.

Several formulations of 2,4-D are in common use. These include: water-soluble salts such as sodium, dimethylamine, ethanolamine, and triethanolamine; oil-soluble amine salts, such as the dodecyl-tetradecyl amine salt and the N-oleyl-1,3-propylenediamine salt which are soluble in most organic solvents; and esters, such as the iso-octyl, butoxyethanol, and the propyleneglycolbutylether ester.

The formulation most commonly used in aquatic sites is the dimethylamine salt of 2,4-D (DMA-2,4-D). The molecular formula of DMA-2,4-D is $C_{10}H_{13}Cl_2NO_{3}$; the molecular weight is 266.1; and

the melting point is 85 to 87° C at which temperature it decomposes. DMA-2,4-D is a white, odorless crystal which is highly soluble in water, less soluble in alcohols and acetone, and insoluble in kerosene and diesel oil. It has been reported (Bridges and Sanders, 1963) that DMA-2,4-D will diffuse through polyethylene bags placed in water.

Effects on Organisms

Microorganisms and algae. Lamartiniere et al. (1964) reported that 2,4-D affected cross wall formation and cell division in several bacterial species. They found a rapid cellular death in the stationary phase of growth. Butler (1963) noted no decrease in productivity of natural phytoplankton or unialgal cultures of <u>Duniella euchlora or Platymonas</u> sp. during a 4-hour exposure to a 1-mg/liter concentration of DMA-2,4-D. Poorman (1973) found that 24 hours' exposure to 1- to 10-mg/liter concentrations of 2,4-D hid no effect on the growth of <u>Euglena</u> <u>gracilis</u>. They also reported no effect after 7 days' exposure to 10 or 50 mg/liter but there was a 26% decrease in growth of <u>Euglena</u> when they were exposed to 100 mg/liter for 24 hours. However, when these cells were diluted and transferred to pesticide-free medium, they recovered. When the freshwater alga, <u>Scenedesmus quadricaudata</u>, was exposed to 0.1 or 1.0 mg/liter of 2,4-D there was a decrease in cell density at 4-B days, some indications of carbon fixation stimulation beginning on day 6, and a small reduction in biomass by day 10 (Stadnyk <u>et al.</u>, 1971). Walsh <u>et al</u>. (1970) reported that concentrations of 0.1, 1.0, or 10.0 mg/liter BEE-2,4-D did not alter the rate of photosynthesis of a species of unicellular marine algae.

Plants. Lynn and Barrows (1952) found no increase in hydrocyanic acid content of wild pin cherry leaves sprayed with an ester of 2,4-D (see Willard, 1950). Buck <u>et al.</u> (1961) also reported no increase in hydrocyanic acid in Canadian thistle sprayed with 2,4-D. They found that calves and ewes ate the sprayed, dried plants with no ill effects. Williams (1968) found that spraying spring parsley with 2,4-D detoxified the plants, thereby decreasing the plants capacity to photosensitize chickens.

Insects. Adams (1960) found that coccinellid larvae were susceptible to an amine salt of 2,4-D. She stated that the developmental period was lengthened when older age groups were sprayed and that mortality was twice as great in sprayed larvae as in controls up to pupation, although mortality during pupation was no greater in

sprayed than control larvae. Rexrode <u>et al.</u> (1971) reported that trees treated with herbicides were more subject to attack by oak bestles and that bestle numbers were higher on treated trees. However, trees treated with DMA-2,4-D had less fungal mat than those treated with picloram and 2,4-D or picloram and arsenic.

Maxwell and Harwood (1960) reported an increased rate of reproduction by pea aphids fed on broad beans sprayed with nonlethal doses of 2,4-D. This increased rate may have been due to increased amounts of free alanine, aspartic acid, serine, and glutathion in the growth terminals of the bean plant, the area of greatest aphid development.

Aquatic organisms. Butler (1963) found that 5 mg/liter of DNA-2,4-D irritated juvenile blue crabs after 24 and 48 hours of testing but no decrease in oyster shall growth occurred after treatment at 2 mg/liter for 48 hours. When ponds were treated at 5 and 10 mg/liter of 2,4-D, there was a 2-week delay in bluegill spawning (Cope, <u>et al.</u>, 1970). Higher treatment levels induced some pathology including depletion of liver glycogen, globular deposits in the blood vessels and stasis and engorgement of the brain circulatory system.

Whitehead (1973) reported that growth rates of 0- to 8-Birds. week old brotlers decreased significantly when fed 10, 50, or 100 mg/kg 2.4-D but that food conversion was not affected by any dietary level. Whitehead and Pettigrew (1972) noted a reduction in food consumption and growth levels in chicks fed 250-900 mg/kg dietary 2,4-D. They found that 5,000 mg/kg caused histological changes, but the chicks resumed normal growth when placed on a normal diet. No effect was noticed on plasma calcium or magnesium. Somers et al. (1973) sprayed fertile chick eggs at the recommended and also at 10 and 20 times the recommended application rate. They found no effect on hatching or growth of the chicks. Chickens tolerated 300 mg/kg/day for several weeks without ill effects (Bjorklund and Erne, 1966). The same workers reported that the most noticeable effect was reduced egg production when chickens were given 500 mg/kg in feed or 1,000 mg/liter in water for prolonged periods. In contrast, Lutz-Ostertag and Lutz (1970) reported 40-75% mortalities in eggs of pheasants, gray partridge, and red partridge sprayed with 2,4-D. They also found fetal malformations, reduced fertility, and physiological castration. Sheldon et al. (1964) fed geese 1,000 mg/kg of ¹⁴C-2,4-D and sacrificed geese at intervals from 22 to 230 days. Treated geese gained less than control geese and also had disorganized cellular structure, hepatic cell destruction, and fatty

degeneration in the liver. Two geese treated for 192 days were placed on a control diet and had almost completely recovered after 7 months.

Mammals. When Collins and Williams (1971) fed hamsters 20-100 mg/kg daily they found occasional terata, decreased fetal viability, and decreased litter size. However, these anomalies were not dose-related.

Brody (1973) induced myotonia in rat muscles in 45 minutes by injecting 200 mg/kg body weight of 2,4-D. The animals did not appear ill and the myotonia disappeared within 24 hours. He postulated that 2,4-D caused an increase in K^+ conductance which could lead to myotonia through a compensatory decrease in C1⁻⁻ conductance.

Philleo and Fang (1967) reported that control rats fed 14 Cacetate converted more than 90% of it to 14 CO₂. When rats were fed 10-20 mg/kg 2.4-D the effect on respired 14 CO₂ was insignificant, however, a dose of 400 mg/kg decreased the amount of respired 14 CO₂ and affected the pathway of 14 CO₂ elimination. Heene (1967) found that 2.4-D inhibited glycogen forming enzymes in rat skeletal muscle. Khera and McKinley (1972) noted fetopathy and an increased incidence of skeletal anomalies following daily <u>per os</u> doses of 100-150 mg 2.4-D/kg. However, weight gain and viability of the offspring were within control limits. Schwetz <u>et al</u>. (1971) administered

doses of 2,4-D esters to pregnant rats 6-15 days after conception. The maximum tabulated dose was 87.5 mg/kg/day. High dose levels resulted in decreased fetal body weight, subcutaneous edema, delayed bone ossification, and formation of lumbar and wavy ribs, all of which were probably dose-related ... They found no teratogenic effects at any dose, nor any effect on fertility, gestation, viability, or lactation. Also, neonatal growth and development were not altered by treatment during pregnancy. Rats tolerated 100 mg 2,4-D/kg without 111 effect (Bjorklund and Erne, 1966). The same workers also found that 50 mg/kg/day could be toxic to pigs. Pigs fed 500 mg/kg for up to 12 months suffered growth depression, locomotor disturbances, anemia, albuminuria, and moderate hepatic and renal degeneration. A sow fed 2,4-D through the gestation period showed no fil effects, but the piglets were underdeveloped and 10 to 15 of them died within 24 hours of birth. There was retarded growth and increased mortality in the second generation when pregnant rats and their offspring were given 1,000 mg/kg 2,4-D for 10 months. Hansen et al. (1971) fed rats up to 250 mg/kg 2,4-D in the diet for 2 years. They found no significant effect on growth rate, survival rate, organ weights, or hematologic values. No 2,4-D related effects were found in beagles fed up to 500 mg/kg dietary levels for 2 years. In a

three-generation rat litter reproduction study, no deleterious effect was noted from 100 or 500 mg/kg dietary treatment, and, while 1,500 mg/kg did not affect fertility nor litter size, it reduced the percentage of pups surviving to weaning and the weights of weanlings.

Sheep given 100 mg/kg of the alkanolamine salt of 2,4-D were unaffected after 481 doses (Palmer and Radeleff, 1964). Cattle were unaffected by 112 doses or 50 mg/kg but died after 44 doses of 200 mg/kg. Sheep also were unaffected by 481 doses of 100 mg/kg of an ester of 2,4-D while cattle suffered weight loss when given 10 doses of 250 mg/kg of the same ester. Radeleff (1964) stated that 2,4-D was absorbed from the digestive tract and eliminated by the kidneys in mammals.

Walker et al. (1972) reported that 2,4-D inhibited tumor development in Ehrlich ascites. They further noted an increased survival time for tumorogenic mice treated with 2,4-D.

Humans. Sare (1972) reported a case of headaches and double vision in a worker spraying 2,4-D. Berkley and Magee (1962) reported the case of a farmer who had cutaneous exposure and had possibly inhaled DMA-2,4-D. He stated he had tingling of the hands and feet, aching arms, and stiffness in his hands and

His dextarity was so reduced he could not button his knees. shirt or the his shoes. He also had reduced perception to painful stimuli in the distal extremeties. The authors concluded that neuropathy from 2.4-D exposure is rare and those who show it are probably predisposed to neuropathy or susceptibility to the toxin. Another source (Anonymous, 1956) also mentions possible sensitivity of certain individuals to phenolic compounds which are contained in some formulations. Willard (1950) also mentioned the possible allergy of some people to 2,4-D. Berwick (1970) reported an accidental ingestion of 7,200 mg 2,4-D which resulted in fibrillary twitching and paralysis of the intercoastal muscles. Bonderman et al. (1971) studied the adaptive responses of some esterase enzymes and found no significant difference in the esterase levels between people who had formulated herbicides for up to 20 years versus controls. Poland et al. (1971) studied 73 male employees in a factory where 2,4-D was manufactured and found chloracne in 18% of thos studied. However, the incidence of chloracne was not correlated with job location in the plant. Johnson (1971) noted no genetic effects on 220 men exposed to 30-40 mg 2,4-D/day. Hayes (1971) stated that the oral dose of 2,4-D required to produce illness in man is probably 3-4 g. An intravenous dose of 2.0 g.

produced no illness, but 3.6 g produced coma, fibrillary twitching of some muscles, hyporeflexia, and urinary incontinence. Mammals killed by large doses of 2,4-D are thought to die of ventricular fibrillation. Treatment for 2,4-D poisoning is symptomatic.

Stability and Degradation

Aly and Faust (1964) decomposed 2,4-D with a UV lamp. They found that decomposition was more rapid at pH 9 than pH 7. They did not feel that UV from sunlight would decompose the herbicide. Daly (1971) found that the butoxyethanol ester of 2,4-D (BEE-2,4-D) and also 2,4-D degraded rapidly under conditions of intense light and high temperature. He found that UV light degraded 50% of a solution of BEE-2,4-D to volatile products in 12 hours.

Crosby (1969). Crosby and Tutass (1966), Crosby and L1 (1969) and Crosby and Wond (1973) found that aqueous solutions of 2,4-D are photolyzed through a series of reactive intermediates with replacement of the chlorines by hydroxyl groups and cleavage of the ether bond. Among the photolytic products are 2,4-dichlorophenol, 4-chlorocatechol, 2-hydroxy-4-chlorophenoxyacetic acid, and 1,2,4-benzenetriol. The latter is then converted via a nonphotochemical oxidation to a nontoxic polymeric humic acid.

Crosby and Wong (1973) reported that side-chain degradation was oxygen dependent and was more rapid at pH 8 than at pH 2, but that ring reactions did not require oxygen. They further stated that the displacement of chloride from the photo-excited ring by hydroxide ions was pH dependent and the final degradation products (condensation products) were the result of ionic and oxidative combinations with substrates.

EFFICACY

DMA-2.4-D is recommended for use primarily on broad-leaved plants. The primary aquatic-related uses are for control of weeds on irrigation canal banks, control of waterhyacinth, and control of Eurasian water milfoll. It is also used to control other aquatic plants such as lotus, arrowhead, waterlily, and smartweed, but on a lesser scale. Treatment should be carried out when plants are young and actively growing before the bud or early bloom stage (WSSA, 1970; Corns and Gupta, 1971; DeRigo, 1964; DeVaney, 1967; Thomas and Duffy, 1968). General aquatic application for emersed plants is 2.2 kg/ha in 400-600 liters of water. A "sticking" agent is sometimes employed. Normally, no more than two applications should be made per year except for possible spot treatment. Grover et al. (1972) found that 3-4% of DMA-2,4-D and the butylester drifted off the target area as droplets and an additional 25-30% of the butylester drifted off as a vapor mass within 30 minutes after spraying. Therefore, no treatment should be made when wind velocity exceeds 10 mph or under rainy conditions. Treatment along ditch banks or other moving water should be done in an upstream direction to avoid large concentrations of the chemical. When a body of

water is nearly covered with an undesirable plant, only part of the water should be treated at one time to avoid the possibility of BOD problems from the decomposing vegetation.

There are conflicting reports regarding the disappearance of 2,4-D from water. Aly and Faust (1964) reported that 2,4-D persisted for 120 days in lake water that was aerobically incubated in the laboratory, whereas esters of 2,4-D under similar treatment were hydrolyzed biologically to the acid in 9 days. They further reported that B1-85% of the 2,4-D was decomposed biologically in lake muds within 24 hours but only after extensive microbial adaptation techniques. Faust and Suffett (1966) verified these results. However, Daly (1971) reported that BEE-2,4-D degraded to 2,4-D within 24 hours in systems containing living material, pond water, polluted water, or water with watermilfoil. All systems produced ¹⁴CO₂, but polluted water produced the most.

Rogoff (1961) discussed the oxidation by bacteria of aromatic compounds including 2,4-D. Anderson <u>et al.</u> (1968), utilizing 2,4-D as the sole carbon source, isolated a strain of <u>Aerobacter</u> sp. which decomposed 90% of the 2,4-D in 15 days. Bell (1957, 1960) isolated an <u>Achromobacter</u> species which could decompose 2,4-D. He reported that 99% of the chlorine was released as chloride. He further stated that requirements for rapid oxidation included a free ortho position, two or less chlorine substitutions on the ring, a <u>para</u> chlorine substitution, and a free side-chain carboxyl group <u>beta</u> to the ether linkage. Others (Robeck <u>et al.</u>, 1963; Steenson and Walker, 1956 and 1953; Rogoff and Reid, 1954) have reported degradation of 2,4-D by microorganisms in the genera <u>Flavobacterium</u>, <u>Achromobacter</u>, and <u>Coxyne</u>. Aerobic conditions were found to be essential, and degradation was attributed to the formation of adaptive enzymes. Walker and Newman (1956) reported that a culture of <u>Mycoplasma</u> sp. decomposed 2,4-D, while Wedemeyer (1966) found a protozoan and paramecium which could metabolize 2,4-D.

Linscott <u>et al.</u> (1968) stated that the resistance of alfalfa to 2,4-dichlorophenoxybutyric acid (2,4-DB) may result from the synthesis of inactive chlorophenoxy compounds having larger side chains than the parent compound, thus preventing <u>beta-oxidation</u> to the toxic parent chemical.

Faust <u>et al.</u> (1961) reported objectionable odors in water treated with 2,4-D, and concluded the odors resulted from production of phenols from the herbicide. Faust and Aly (1963) reported that 2,4-dichlorophenol (2,4-DCP) was metabolized slowly under either acid or anaerobic conditions. Activated carbon was capable of removing 2,4-D and 2,4-DCP from water, whereas

strongly basic anion-exchange resins sorbed 2,4-DCP and Na-2,4-D but not the esters of 2,4-D (Faust and Suffet, 1966). Inglis and Davis (1968) found that hardnesses ranging from 13-365 mg/liter CaCO3 had no significant effect on the toxicity of organic herbicides to fish.

A brief summary of the efficacy of 2,4-D is listed in Table 1. The data are primarily related to aquatic plants. Exceptions are where the applications may have had an effect on vegetable crops (via irrigation) or effects on wildlife. Most emersed broadleaved aquatic plants are controlled by 2,4-D. Aquatic grasses and submersed species are not controlled or require massive doses of the herbicide for control.

TOXICITY

The toxicity of 2,4-D depends greatly on the formulation. For example, Sanders (1970) reported the following TL₅₀ values at 48 hours for <u>Daphnia magna</u>: propyleneglycobutyl ester, 0.1 mg/liter; DMA-2,4-D, 4 mg/liter; and BEE-2,4-D, 5.6 mg/liter.

Lawrence (1962b) reported that 80% of the largemouth bass exposed to 1 mg/liter DMA-2,4-D died in 72 hours. On the other extreme, Stickel (1964) reported an LD_{50} of 56,776 mg/kg at 94 days for adult <u>Coturnix</u> quail.

Table 2 is a compilation of toxicity data in which specific amounts of the herbicide and specific mortalities were reported by various authors. The remainder of the toxicity data is summarized on the following pages.

Plants. Elder et al. (1970) reported that 2,4-D exhibited low toxicity to all freshwater and marine algal organisms tested at rates approaching the maximum solubility of the herbicide in water. DeRigo (1964) found that 2,4-D injured commercially grown Chinese waterchestnut. A single annual application of 6.7 kg 2,4-D/ha in diesel fuel was found to be effective for

cattail control with no deleterious effects on the biological activity of sewage lagoons from repeated applications (Corns. and Gupta, 1971). Kleinschmidt (1969) found that 2,4-D gave good control of Hydrilla verticillata while others (Daly, 1971; Haven, 1963; Smith and Isom, 1967; Wojtalik et al., 1971) reported control: of Eurasian watermilfoil with DMA-2,4-D or BEE-2,4-D. Hildebrand (1946), Schultz and Harman (1974), and Schultz and Whitney (1974) reported that waterhyacinth could be controlled by 2.4-D. Lapham (1964) controlled alligatorweed with a combination of 2,4-D/and dichlobenil while Foret (1967) used 2,4-D to control Elodea canadensis. Pierce (1960, 1961) found that 2,4-D eliminated Nymphaea odorata and Brasenia sp., reduced Utricularia purpurea, but seemed to accelerate growth of Potamogeton spp. She also reported that 2,4-D in combination with other herbicides was toxic to Potamogeton crispus and several other aquatic plants (1968, 1969).

Several authors (USDA, 1968; Lawrence, 1962a) have compiled references dealing with the toxicity of herbicides to aquatic plants while others (DeVaney, 1967; Lawrence, 1962b; Mullison, 1970; Gangstad, 1972; and Pimentel, 1971) have reviewed the toxicity of 2,4-D to higher aquatic plants, algae, fish, and non-target organisms. The toxicity of 2,4-D to terrestrial plants is well known and too lengthy to document. Keith <u>et al.</u> (1959) reported a decrease in pocket gophers due to the reduction of food plants such as dandelion, agoseris, western yarrow, and penstemon. Krefting and Hansen (1963) improved deer habitat by top-killing mountain maple with 2,4-D. The herbicide also increased conifer sprouts and stimulated regrowth of maple sprouts, both of which are excellent deer browse. Hee <u>et al.</u> (1973) found that the oil-soluble amine formulations were absorbed faster by sunflower leaves than the water-soluble amines, thereby decreasing the length of time for death of the plant.

Microorganisms. Petruk (1965) added 2,4-D to fish nursery ponds and found an increase in microorganisms in 2 to 3 days. The total numbers decreased after 12 days, but reached the initial value again in 23 to 35 days. He stated that the herbicide stimulated hetarotrophic organisms which increased the reservoir productivity.

Invertebrates. Butler (1965) found no effect on the growth of oyster shell after 96 hours' exposure to 2 mg/liter of 2,4-D. He also noted no effect of 2 mg/liter on brown shrimp after 24 hours' exposure. In static bioassays, 10 mg/liter of BEE-2,4-D

had no effect on grass shrimp (Hansen et al., 1973). The same authors reported that grass shrimp avoided 2,4-D but not 5 insecticides. Walsh (1971) found that 5 mg/liter of DMA-2,4-D had no effect on blue crabs after 24 or 48 hours' exposure and that 1 mg/liter had no toxic effect on Eastern oysters at 96 hours' exposure.

Walker (1953) reported that 2,4-D decreased the number of bottom organisms in ponds, presumably through removal of cover or feed. No adverse effects were noted on benthos or other aquatic invertebrates in large-scale field applications of BEE-2,4-D (Beaven et al., 1962; Sears and Meehan, 1971; Smith and Isom, 1967; Thomas and Duffy, 1968; Whitney, 1970). In another study, Rawls (1965) found that only the acetamide formulation was toxic to aquatic invertebrates at field application rates.

Adams (1950) and Adams and Drew (1965) reported that applications of 2,4-D could enhance aphid infestations of oat fields by the toxic effects on the coccinellid larvae, which biologically control the aphid.

Moffett and Morton (1971) and Moffett <u>et al</u>. (1972) found that 2,4-D was nontoxic to honeybees when applied in a water carrier, but oil carriers themselves were toxic. Beilman (1950) sprayed shrubs along roads with 2,4-D and assumed bees could not use the area. Surprisingly, the shrubs were replaced by sweet clover which resulted in good honey production.

There was no effect on the survival of earthworms treated for 2 hours in 100 mg/liter of 2,4-D (Martin and Wiggans, 1959). Neither was the herbicide toxic to wireworms, spring tails, mites, or grasshoppers (Fox, 1964; Putnam, 1949). The reproduction of plant parasitic nematodes was inhibited by 5 mg/liter and impaired by 8 ug/liter of 2,4-D (Webster and Lowe, 1966). Reviews dealing with the effect of 2,4-D on invertebrates incTude those by USDA (1968), Bohmont (1967), Mullison (1970), and Pimentel (1971).

Fish. The earliest report dealing with the toxicity of 2,4-D to fish is that of King and Penfound (1946). They reported that 1 mg/liter was not toxic to bluegills or largemouth bass and 100 mg/liter was only slightly toxic. The herbicide DMA-2,4-D was not toxic to fry of bluegill, green sunfish. Take chubsucker, or smallmouth bass exposed to 25 mg/liter for 8 days (Hilltebran, 1967). Butler (1965) found no effect on longnose killifish treated for 48 hours with 15 mg/liter of DMA-2,4-D. There were no mortalities in bluegill exposed to 5 mg/liter of 2,4-D for 6 weeks (Cope et al., 1970), and 10 mg/liter was not toxic to squawfish (MacPhee and Ruelle, 1969). Sergeant <u>et al</u>. (1971) found that the acid and salt formulations of 2,4-D were non-toxic to

green sunfish at 110 mg/liter. However, BEE-2,4-D was toxic at the same concentration. There was no toxicity to fish noted in large-scale field applications of BEE-2,4-D or DNA-2,4-D (Beaven <u>et al.</u>, 1962; Sears and Meehan, 1971; Whitney, 1970; Wojtalik <u>et al.</u>, 1971). In another field study, Smith and Isom (1967) reported that fish temporarily moved out of an area treated with BEE-2,4-D, but no toxicity was noted.

Summaries or references to fish toxicity data are given in USDA, 1968; Bohmont, 1967; Lawrence, 1962a; Mullison, 1970; and Pimentel, 1971.

Amphibians. Frogs and turtles showed no toxic effects from 2,4-D treatment of ponds (Pierce, 1961). Sanders (1970) reported that tadpoles of the Western Chorus Frog withstood 50 mg/liter of DMA-2,4-D for 96 hours.

Birds. Fertile hen eggs tolerated injected doses of 50 mg/liter of 2,4-D with no effect on hatching (Dunachie and Fletcher, 1970). In the same study, the only teratogenic effect was feather blanching, which was not fatal. Somers <u>et al.</u> (1973) sprayed fertile chick eggs with the recommended field rate, and 10 and 20 times the recommended rate. They found no toxic effects on incubation performance, hatching, or growth of the chicks. Daily doses of 300 mg/kg of 2,4-D for several weeks were not toxic to chickens (Bjorklund and Erne, 1966). Whitehead and Pettigrew (1972) found that chicks tolerated a dietary level of 5,000 mg/kg for I week without toxic effects. Schultz and Whitney (1974) reported that field application of 2,4-D had no effect on the hatching of boat-tailed grackle eggs or development of fledglings.

Mammals. Hansen et al. (1971) fed rats 1,250 mg/kg and beagles 500 mg/kg of dietary 2,4-D for 2 years with no toxic effects. Seven dosem of the alkanolamine salt of 2,4-D at 500 mg/kg was fatal to sheep and 44 doses of 200 mg/kg was fatal to cattle (Palmer and Radeleff, 1964). In the same study, nine doses of the propylene glycol butyl ether ester of 2,4-D at a level of 250 mg/kg was fatal to sheep. Repeated daily doses of 50 mg/kg of 2,4-D were fatal to pigs (Bjorklund and Erne, 1966). No toxicity to cattle wus noted when they ingested forage from pastures sprayed with normal or 2 to 4 times the recommended rate of the herbicide (Mitchell et al., 1946; Grigsby and Farwell, 1950). Hassall (1965) stated that animals should be excluded from sprayed areas for 2 weeks to eliminate possible toxic effects of the herbicide and also because naturally poisonous weeds may

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be rendered more palatable after spraying. Clegg (1971) reported no effects at 25 mg/kg of 2.4-D and further stated that no problems would be anticipated if dioxin contamination was not a problem.

Human safety. Some humans evidently have individual sensitivity to 2,4-D or the other contents of the formulation (Anonymous, 1956; Berkley and Mages, 1962). One individual accidently ingested about 110 mg/kg body weight without fatal results (Berwick, 1970). Hayes (1971) stated that the oral dose of 2,4-D required to produce illness in man is about 3 to 4 g. An intravenous dose of 2.0 g produced no illness but an intravenous dose of 3.6 g produced coma, hyporeflexia and urinary incontinence. Recovery was marked in 24 hours and complete in 48 hours. An oral dose of 6.5 g led to convulsions, followed by death. No genetic effects were noted in 220 men exposed to 30 to 40 mg of 2,4-D/day over extended periods (Johnson, 1971). RESIDUES

Methodology. The various formulations of 2,4-D are recovered from water by treating the water with acid or base, partitioning with an organic solvent, followed by a column cleanup and derivatization for glc analysis (Devine and Zweig, 1969; Faust and Soffet, 1966; Schultz, 1973; Schultz and Harman, 1974; Schultz and Whitney, 1974; Wilder, 1968). Hesselberg and Johnson (1972) and Rodgers and Stalling (1972) used column extractions for 2,4-D residues in fish. Coakley <u>et al.</u> (1964) and Duffy and Sheldon (1967) gave procedures for extracting BEE-2,4-D from shellfish and fish. Procedures for extracting 2,4-D from mud entail the use of methanol, acetone, or ethyl ether combined with other cleanup procedures (Hesselberg and Johnson, 1972; Schultz, 1973; and Woodham <u>et al.</u>, 1973).

Procedures for extracting 2,4-D residues from fatty foods, animal products, and animal tissues utilize solvent extraction, partitioning with base, soxhelet extraction, and cleanup procedures (U.S. Dept. HEW, 1968; Clark, et al., 1967; Crosby and Bowers, 1966; Yip, 1971). Meagher (1965a, 1966b) gave procedures for extracting 2,4-D, the isopropyl ester of 2,4-D, and a 2,4-D conjugate from citrus peel, while Gutenmann and Lisk (1963) and Yip (1962) gave procedures for determining 2,4-D residues in forage crops.

Rivers et al. (1970) and Shafik et al. (1971) reported methods for determination of 2,4-D in human urine and blood.

Garbrecht (1970) and Scoggins and Fitzgerald (1969) reported esterification procedures for chlorophenoxy acids prior to glc analysis. Purkayastha (1969) detected ionizable herbicides by means of electrophoresis.

Residues. The persistence of 2,4-D in water has been reported by many workers. Averitt (1967) sprayed waters in Louisiana with DMA-2,4-D at the rate of 4.48 kg/ha and found maximum concentrations of 727 ug/liter and 1,020 ug/liter after 1 week in middle and bottom strata of water, respectively. After 3 weeks, concentrations had fallen to 12 ug/liter and 10 ug/liter, respectively. He also sprayed two lagoons at the same rate and found concentrations of 689 ug/liter after 31 days. Grzenda (1963) treated ponds with 0.6 ug/liter. After 62 days residues were less than 1 ug/liter. Kleinschmidt (1969) reported detectable levels of residues in water treated with BEE-2,4-D for up

to 22 days but stated that no residues could be detected after 29 days. Frank and Comes (1967) treated ponds with 1.33 mg/liter BEE-2,4-D and found only 0.024 mg/liter 1 day post= treatment and less than 0.00] mg/liter after 36 days. In ponds treated at 8.96 kg/ha. residues declined from 0.34 mg/liter and 0.69 mg/liter after 1 day in Florida and Georgia ponds, respectively, to less than 0.005 mg/liter by 28 days (Schultz and Harman, 1974). In Missouri ponds treated at the same rate it took 56 days for residues to decline to less than 0.005 ug/liter. The highest residue found in water sprayed at 4.48 kg/ha of an of1-soluble amine salt of 2,4-D was 37 ug/liter, 1 day posttreatment (Schultz and Whitney, 1974). When irrigation ditchbanks were treated at 1.56 to 3.36 kg/ha 2,4-D, the maximum concentration found in the canals was 213 ug/liter and 25 to 61 ug/liter in the irrigation water (Bartley and Hattrup, 1970; Frank et al., 1970). Manigold and Schulze (1969) reported a residue of 0.35 ug/liter in 15 out of 20 stations, while Brown and Nishioka (1967) found no herbicide residues from streams analyzed for 2,4-D as part of the National Pesticide Monitoring Program.

Schwartz (1957), utilizing $^{14}C-2,4-D$, reported that 60% of the ^{14}C remained in sewage effluent for 3 to 6 months. However, he determined only ^{14}C and not actual 2,4-D. Oysters and clams

contained between 3.5 and 3.7 mg/kg of 2.4-D after exposure for 3 days to an application of 33.6 kg/ha (Coakley <u>et al.</u>, 1964). Smith and Isom (1967) reported a maximum concentration of 1.12 mg/kg in mussels taken from water treated at 44.8 to 112 kg/ha of BEE-2,4-D. Oysters contained residues of 1.45 mg/kg after treatment of 44.8 kg/ha of BEE-2,4-D (Thomas and Duffy, 1968). Wojtalik <u>et al.</u> (1971) reported that plankton contained 3.6 ug/kg 4 weeks after treatment at 44.8 kg/ha DMA-2,4-D. Whitney (1970) reported that the highest residue in benthic organisms (grass shrimp, damselfly nymphs, and scud) was 0.23 mg/kg 24 hours post-treatment and that residues in blue crabs never exceeded 0.10 mg/kg.

Sears (1971) found the maximum concentration was 0.5 mg/kg in fish from streams adjacent to cut-over land sprayed with 2,4-D, while Cope <u>et al.</u> (1970) could detect no 2,4-D in fish 4 days after treatment with up to 10.0 mg/liter of 2,4-D in ponds. Walsh (1971) reported no residues of 2,4-D in fish fed 10 mg/kg in their foud for 2 weeks and 8.4 mg/liter in fish fed 100 mg/liter for 2 weeks. No residues were detected in the latter fish after being fed a herbicide-free diet for 2 weeks. Rodgers and Stalling (1972) found that whole body or tissue residues in rainbow trout, bluegill, and channel catfish were proportional

to the exposure concentration. Only the liver contained detectable quantities of BEE-2,4-D, whereas the other organs contained only the acid with the highest concentration in bile. Schultz (1973) exposed fish to 14C-DMA-2,4-D and found radioactive residues ubiquitous for 84 days. However, the actual 2.4-D content was negligible. Among three groups of ponds treated at rates of 2.24, 4.48, or 8.96 kg/ha of 2,4-D, Schultz and Harman (1974) found no residues in fish 14 days post-treatment in Florida ponds, a residue of 0.075 mg/kg in bluegill from Georgia ponds, and no residues 28 days post-treatment in fish from Missouri ponds. Of fish taken from Florida canals sprayed at 4.48 kg/ha of 2.4-D, only 3 contained residues as high as 0.1 mg/kg, 16 had less than 0.01 mg/kg, and 41 had no detectable residues (Schultz and Whitney, 1974). Whitney (1970) reported that the highest residues in fish were 0.23 mg/kg in largemouth bass as a result of treating 200 acres of Eurasian watermilfoil with BEE-2,4-D, while Smith and Isom (1967) reported a maximum residue of 0.20 mg/kg in fish from an area treated for milfoil control.

Breast muscle and liver of Florida gallinules contained 2,4-D residues of 0.30 and 0.68 mg/kg, respectively, 1 day, and no detectable residues 4 days after canals at Loxahatchee National Wildlife Refuge were sprayed with 2,4-D for waterhyacinth control (Schultz and Whitney, 1974).

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Residues were detected in mud for up to 6 weeks from ponds treated at 0.1 mg/liter to 10 mg/liter (Cope et al., 1970). Frank and Comes (1957) reported residues of 4.96 mg/kg, 0.06 mg/kg, and less than 0.005 mg/kg in mud from ponds 1, 36, and 85 days post-treatment with a 1.33-mg/liter concentration of 2,4-D. Mud from Florida and Georgia ponds treated at 2.24, 4.43, or 8.96 kg/ha of 2.4-D contained no residues exceeding 0.05 mg/kg. Mud from Hissouri ponds treated similarly contained a maximum residue of 0.17 mg/kg at 3 days and no detectable residue 28 days after treatment (Schultz and Harman, 1974). The maximum residue in mud from Loxahatchee Refuge was 0.005 mg/kg 15 days post-treatment with 4.48 kg/ha of 2,4-D (Schultz and Whitney, 1974). Nojtalik et al. (1971) reported residues of 0.30 mg/kg in mud 2 months after treatment at 22.4 or 44.8 kg/ha of DMA-2,4-D and residues less than 0.10 mg/kg 6 months post-treatment.

Cope et al. (1970) reported that vegetation from ponds treated with 0.1 to 10.0 mg/liter of 2,4-D contained residues for up to 6 weeks. Smith and Isom (1957) found that Eurasian watermilfoil contained 8.26 mg/kg 24 hours after treatment at either 44.8 or 112 kg/ha of BEE-2,4-D. When rats were fed 100 mg 14 C-2,4-D, all tissues and organs examined contained radioactive material 6 to 8 hours after the dose. This radioactive residue persisted for 17 hours. The largest amount of residue was 1,690 ug/0.1 g dry tissue found in the stomach of rats 24 hours after a dosage of 80 mg (Khanna and Fang, 1966). Zielinski and Fishbein (1967) reported that esters of 2.4-D disappeared more rapidly than the free acid in mice fed various 2,4-D formulations. Only about 10% of the 2,4-D was recoverable 24 hours after treatment. No 2,4-DCP was detected in animals injected with 2,4-D or its butyl or isooctyl ester.

Less than 0.05 mg/kg 2,4-D or 2,4-DCP was found in the milk or cream of cows fed 30 to 1,000 mg/kg dietary 2,4-D for 2 to 3 weeks and then placed on untreated feed for 1 week (Leng, 1972). Negligible residues (less than 0.1 mg/kg) were found in the muscle and fat of slaughtered animals fed 2,4-D for 4 weeks at 300 mg/kg. Cows fed 300, 1,000, or 3,000 mg/kg of 2,4-D contained residues of 1.1, 4.9, and 5.3 mg/kg, respectively, in their kidneys. Residues in sheep were similar to those found in cows. There was no withdrawal period in the latter two studies (Leng, 1972). When a steer was fed 113.5 mg of 2,4-D, 100.65 ug was recovered in the urine (Lisk et al., 1963). Gutermann et al. (1963) fed a Jersey cow 5 mg/kg of 2,4-D in grain for 5 days. No residues were detected in daily samples of milk or feces. They also fed a heifer 5 mg/kg and took samples from a fistula over a 23-hour period. Recovery of 2,4-D dropped from 3.5 initially to 0.5 mg/kg at the end of the experimental period.

Metabolites and degradation products. Extensive studies have been conducted on the degradation and metabolism of 2,4-D by bacteria. Bell (1957, 1960) reported that an Achromobacter species isolated from soil would degrade 2.4-D via exidation. He reported that about 94% of the chlorine from 2.4-D was released as chloride. Anderson and Okrend (1968) found an Aerobacter species which degraded 2,4-D by 90% in 15 days. A species of Arthrobacter cleaved the ether linkage of 2,4-D acetate resulting, through oxidation, in 2,4-DCP (Tiedje and Alexander, 1969). The enzyme responsible for this cleavage converted ¹⁴C-acetate-2.4-D to alanine and a volatile product. The authors proposed that glyoxylate was the initial product formed in cleavage of the side chain and that alanine was produced by condensation of two molecules of either glyoxylate or glycine. Tiedje et al. (1969) reported that 2,4-D was metabolized by an enzyme preparation to succinic acid. An enzyme from an extract of an Arthrobacter species dehalogenated 2,4-D and produced 2,4-DCP which was, in turn, further metabolized (Loos et al., Other workers found that extracts from an Arthrobacter 1967).

species would degrade 2.4-DCP, and other secondary degradation products of 2,4-D (Bollag et al., 1968a, 1968b, Duxbury et al., 1970).

Faulkner and Woodcock (1954) stated that Aspergillus niger would metabolize 2,4-D to 2.4-dichloro-5-hydroxyphenoxy-acetic acid while Steenson and Walker (1958) reported transformation of 2,4-D into 2,4-DCP by a strain of Flavobacterium peregrinum. Rogoff and Reid (1954) reported a Coxyna bacterium which metabolized 2,4-D in buffered solution with a quantitative yield of chloride ion. Two Pseudomonas strains from soil utilized 2.4-D as a sole carbon source (Evans et al., 1971). One of the pseudomonads converted 2,4-D to 2,4-DCP, 2-chlorophenol,3,5dichlorophenol, and alpha-chloromuconate. The second Pseudomonad metabolized 2.4-D to 2.4-dichloro-6-hydroxyphenoxy-acetate, 2.4-DCP, 3.5-dichlorocatechol and alpha-gamma-dichloromuconate. The authors found that dechlorination had to occur at the 4 (para) position before ring fission could occur. Gaunt and Evans (1971) and Gamar and Gaunt (1971) found that a 2,4-D analog was metabolized in similar fashion to that above. Apparently, the formation of adaptive enzymes is necessary for the rapid degradation of 2,4-D (Robeck et al., 1963; Schwartz, 1967; Steenson and Walker, 1955; and Walker and Newman, 1956).

Daly (1971) found that BEE-2,4-D was degraded to 2,4-D in systems containing living material, pond water, polluted water, and water with milfoil, while Rodgers and Stalling (1972) reported that the hydrolysis of BEE-2,4-D to 2,4-D was accelerated by the presence of fish. The sodium salt of 4-(2,4-dichlorophenoxybutyric acid) [4-(2,4-DB)], on analog of 2,4-D, was converted to 2,4-D by fish (Gutenmann and Lisk, 1965), also Lisk et al. (1963), reported that steers fed 4-(2,4-DB) eliminated it in the urine as 2,4-D. When fish were exposed to DMA-2,4-D, one of the major metabolites was a glucuronide (Schultz, 1973).

The majority of the 2,4-D fed to rats was excreted unchanged in the urine (Shafik et al., 1971). Mitchell et al. (1946) reported finding 2,4-D (probably as a salt) in the blood serum of cows fed forage treated with 2,4-D. Eisner et al. (1971) reported finding 2,5-dichlorophenol in the defensive froth of grasshoppers and postulated that it was a degradation product of the 2,4-D applied to vegetation. Sheep fed ¹⁴C-2,4-D yielded approximately 96% of the intact molecule in the urine in 12 hours while less than 1.4% was excreted in the feces (Clark et al., 1964). These authors found little radioactivity in edible tissue and concluded that 2,4-D was excreted essentially unchanged by sheep.

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In an early report of 2,4-D breakdown, Winston and Ritty (1961) stated that under field conditions 2,4-D was not degraded to 2,4-DCP but, rather, to CO2, HCT, and H20. Thomas et al. (1964) reported that phenoxyacetic acids with a chlorine atom at position 4 were not readily hydroxylated. A heat= labile, insoluble, conjugated form of 2,4-D was found in citrus peel by Meagher (1966b). He speculated that the herbicidal molecule was probably conjugated with pectin. Linscott et al. (1968) and Hagin et al. (1970) reported that 4-(2,4-DB) and 2,4-D were converted to inactive compounds by resistant grasses. These inactive compounds had longer side chains than the parent compound thus preventing beta-oxidation and also preventing subsequent translocation to the site of action. Freed and Montgomery (1969) reported that the principal routes of metabolism of phenoxyacetic acids in plants were via conjugation and hydroxylation. Menzie (1969) and Loos (1969), compiled extensive reviews of the degradation of 2,4-D in soil, plants, bacteria, and animals.

REGISTRATION STATUS

In the April 27, 1972 <u>Federal Register</u> (Vol. 37, No. 82) the Environmental Protection Agency and Food and Drug Administration (KEW) announced the issuance of proposed tolerances. for dimethylamine salt, 2,4-D (DMA-2,4-D) used in control of weeds on irrigation ditch banks. A 0.1-mg/liter (negligible residue) tolerance was issued for DMA-2,4-D that may be present in potable water only as a result of application of DMA-2,4-D to ditch banks in western United States in programs of the following groups: Bureau of Reclamation; cooperating water user organizations; the Bureau of Sport Fisheries and Wildlife, USDI, and the Corps of Engineers, USDD (21 CFR Part 121). Livestock are to be excluded from treated ditch banks, and there is no reasonable expectancy of 2,4-D residues in milk or meat as a result of ditch bank treatment.

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At present there is a request for a label for use of DHA-2,4-D for control of weeds on irrigation canal banks. This petition is sponsored by the Bureau of R-clamation and cooperating water users and the Bureau of Sport Fisheries and Wildlife, USDI, Agricultural Research Service, USDA; and the Corps of Engineers, USDD.

The Corps of Engineers, USDD is also presently seeking a label for the use of DNA-2,4-D for control of waterhyacinth in moving waters. Petitions have also been supported by, or submitted by, Amchem Products, Inc. and the Dow Chemical Company for other formulations of 2,4-D.

A copy of one proposed label has been appended to this manuscript.

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Species	Concentration	Exposure	Formulation	Comments	Citations
Red Mexican beans	2.24 kg/ha 6.72 kg/ha	<u></u>	Acid	0.91 kg during seedling stage injured top growth and roots but didn't reduce yield. 2.73 kg re- duced yield 40%. 0.91 kg at bloom stage did not reduce yield. 6.72 kg reduced yield 29%.	Bruns, 1954
Sugar beets	0.91 & 2.72 kg in water for irrigation		Acid	Both concentrations lethal during seedling stage and reduced stands in 1 in. diameter stage. 2.72 kg at 1 in. diameter stage reduced sucrose.	Bruns, 1957
Concord grape '	2.24, 4.48, 8.96, 17.9 kg/ha		Acid .	Lethal to 13, 33, 65, & 54% of plants respective- ly, lesser conc. No effect. Root systems of in- jured plants partially destroyed.	Bruns & Clore, 1958
Cattail .	6.7 kg/ha		BEE	Single annual application in 112.3 liters diesel fuel/ha was effective for control. No effect on biological activity at nearby sewage lagoons.	Corns & Gupta, 1971
Eurasian watermilfoil	4.48 kg/ha	·	BEE		Daly, 1971
Lotus Spätterdock Water chestnut Arrowarum	2.24-4.48 kg/ha 6.72-8.96 kg/ha 6.72-7.84 kg/ha		DMA		DeVaney, 1967
Needlerush Hibiscus Groundselbush	33.6 kg/ha 7.84 kg/ha		. -		
Hightide bush Swamp loosestrife Willow	1.81 kg/378.5 liter water		•		
Elodea Canadensis	4.48 and 6.72 kg/ha	2 months	DMA combined with / molasses.	4.48 kg/ha gave 50% control in 2 months. 6.72 kg/ha gave 74% control in 2 months	Foret, 1967

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Table 1.-- Efficacy data for 2,4-D on plants. Arranged alphabetically by authors.

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Species	Concentration	Exposure	Formulation	Comments	Citations
Waterlily, lotus, spatterdock, pennywo willow, sedge, smart weed, cattail, arrow head, pickerel weed, bullrush, spikerush, knot grass, needler southern water grass Sago pondweed, American pondweed	t- ī. ∾- · ī. •	•	Acid	Not good control for the last two. Not good for use in irrigation canals due to lack of persist- ence in the soil. The first four were well con- trolled by 2,4-D. On all plants between the first 4 and the last 2 control was marginal.	Frank <u>et al</u> ., 1963
Eurasian watermilfo	11 Not given	.•	2-ethylhexyl and Granular BEE	Some reduction in amphipods and molluses probably due to secondary effects such as smothering and habitat changes.	Haven, 1963
Waterhyacinth	1:1430 (acid:water W/ 1:1140 (acid:water W/		Acid	Got 90-100% kill	Hildebrand, 1946
Dandelion Agoseris Western yarrow Rydberg Penstemon	Not given			Reduction was by 91%, 99%, 85%, and 88% re- spectively.	Keith <u>et al</u> ., 1959
Hydrilla	97 kg/ha		Granular contain- ing 20% BEE of 2,4-D	Good control	Kleinschmidt, 1969
	0.91 kg/13.25 liter wat aerial spray	er	BEE	Used to top kill maple and stimulate undergrowth used for deer browse.	Krefting and Hansen, 1963
Alligatorweed	8.96 kg/ha		DMA	Excellent control after 16 weeks when used with dichlobenil.	Lapham, 1964
Pithophora, Alli- gatorweed, water- hyacinth, duckweed, waterstar grass	5 mg/l in water, 100 in water, 5 mg/l spra plants		DHA	% control for 5 mg/l in H ₂ O was 40, 40, 90,40, 90 re- spectively. % control for 100 mg/l in H ₂ O was 95, 90, 100, 100, 100 respectively. % control for 5 mg/l sprayed on plants was 35, 80, 100, 45, 95, re- spectively.	Lawrence, 1962b

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Spectes	Concentration	Exposure	Formulation	Comments	Citations
iyacinth Wijigatorweed	2.24-4.48 kg/ha 8.96 kg/ha	- -	Amine salt - DMA	" Used for control in 1945 & 1948-50 Used in 1950-52 with spotty results.	Martin <u>et al</u> ., 1957
lymphaea odorata Potamogeton Utricularia purpurea	1.5 mg/1. 3.1 mg/1	11 months	Granular	Nymphaea odorata eliminated by all 3 concen- trations. Potamogeton accelerated by 1.5 mg/1. 3.1 and 6.2 mg/1 reduced but did not eliminate U. purpurea.	Pierce, 1960
Nymphaea odorata Brasenia sp. Utricularia purpurea :	10% a.e. at 0.9, 1.3 & 1.8 mg/1; 20% a.e. ; 1.8, 2.6, & 3.6 mg/1.	it	Aqua granular	N. odorata and Brasenia sp. eliminated at all concentrations. U. purpurea reduced but not eliminated by 1.8, 2.6, & 3.6 mg/l.	Pierce, 1961
Surface plants	1 mg/1; 2 mg/1.	8 weeks		Applied in combination with fenac. Resulted in 50% reduction in surface weeds. Sprayed in combination with endothal at 2 mg/l. 50% clearing of surface noted in 8 weeks. No effect in Chara, Potamogeton, common sedge, fish or inverts.	
Potamogeton crispus	2 mg/1.	3 weeks	• •	Applied with endothal. Hastened winter bud formu- lation and/or death of the plant in 3 weeks.	Pierce, 1969
Waterhyacinth	4.48 kg/ha		Amine(oil)- Emuls- amine 3 DMA-2,4-D	General application was with the Emulsamine 3 and spot treatment was with DMA-2,4-D	Schultz and Whitney, 1974
Eurasian watermilfoil	44.8-112 kg/ha		2,4-D BEE	Ne adverse effects noted on mussels, clams, aquatic fauna or water quality. Fish moved out of treated a	Smith & Isom, 1 area _ 1967
Eurasian watermilfoil	10.16 metric tons 20% a.e./80.94 ha.		BEE	Improved Habitat. Destroyed the milfoil. Residues in various flora and fauna from the treated area are given.	Whitney, 1970
Spring parsley	Not given		DMA, PGBEE	2,4-D and 2,4,5-T controlled and detoxified spring parsley.	Williams, 1968
Eurasian watermilfoil	22.4-44.8 kg/ha		DMA-2,4-D	No distinguishable response noted in zoo or phytoplankton, benthic macroinvertebrates or fish.	Wojtalik <u>et al.</u> , 1971

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Organisms	<u>Concent</u> Treatment	ration Toxicity	Exposure	Water Temp	Loading	Testing regime	Comments •	Citations
Concord grapes	2.24 kg/ha 4.48 " 8.96 " 17.9 "	13% killied 33% " 65% " 54% "	•	- <u></u>		•	Lesser concentrations showed no effect. Roots partially destroyed.	Bruns & Clore 1958
Dyster Tarva	0.025 mg/1 0.05 " 0.10 " 0.025 " 0.025 " 0.10 " 0.25 " 0.25 " 1.00 " 10.00 " 50.00 "	<pre>% mortality -5 -3 -20 -2 -16 -2 -7 -8 -12 -11 -3 -48 -55</pre>	•	. ·	•		First 4 mortality figures from tests with a 2,4-D ester. The rest of mor- talities from tests with an unnamed 2,4-D salt.	Davis & Hidu, 1969
Saphnia magna		·IC50- >100 ug/g	1	•			Immobilization concentration	Crosby & Tucker, 1966
Daphnia magna		TL50-0.10 mg/1 " -4.0 " -5.6 "	48 hr.	2]°C	•		PGBE DMA BEE	Sanders, 1970b
Seed shrimp		TL50-0.32 " " -8.0 " " -1.8 "	48 hr.	2]°C		-	PGBE DMA BEE	·
Scud		TL50-2.6 " " ~>100.0 " " -2.2 "	af ja na	64 2 61 58			PGBE DMA BEE	
Sowbug		TL50~2.2 " " ->100.0 " * -3.9	48 hr.	15.5° C			PGBE DMA BEE	
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Table 2.-- Toxicity of 2,4-D.

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Organisms	Concentr Treatment	ation Toxicity	Exposure	Water Temp.	Loading	Testing regime	Comments .	Citations
Class shrimp		TL50-2.7 mg/l " - >100.0 " " - 1.4 "	48 hr.	21°C	•		PGBE . DMA BEE	•
Crayfish		7150->100.0 " >100.0 >100.0	48 hr.	15,5° C "	• .		PGBE DMA BEE	
Gammarus lacustris (Scud)		LC50-1400 ug/1 -2100 -6800 LC50-760 -1800 -4600 -4600 -440 -1600 -2400 -2400	24 hr. " 48 hr. " 96 hr.	21°C """"""""""""""""""""""""""""""""""""	.	Ti at	2,4-D-BEE " -PGBE " -IOE 2,4-D-BEE " -PGBE " -IOE 2,4-D-BEE " -IOE 2,4-D-BEE " -IOE " -IOE here was no effect from 100mg/1 DMA fter 96-hr.	Sanders, 1969
Fiddler crab (Uca pugnax)	10,000 mg/kg 5,000 2,500 " 1,000 ") 100%(50%/72 hrs) 100%(50%/96 hrs) 100% 100%(10-20%/2 wk	108 days 10 days s) 17 days	•	-	Removed and washed after 12-hrs.	2,4-D Acid. Injections of the Na-Salt showed toxicity to be greater than 0.4 mg/g.	Sudak & Claff 1960
Stone fly naiad		LC50~8.50 mg/1	24 hr. 48 hr. 96 hr. 24 hr. 48 hr. 96 hr.		•		2,4-D 8EE " 2,4-D	Sanders & Cope 1968
Coccinellid larvae	8 oz ae/acre	e 31 of 77 died		.	•	•	Used mixed amine salts. Mortality mor than 2 times as great in treated than controls up to pupation. During pupat mortality was no greater. Deformity y greater in larvae sprayed at later sta development.	in. tion vas

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Organisms T	Concentration reatment Toxicity	Exposure	Water Temp.	Loading	Testing regime	Comments	Citations
Honey Bee (Apis mellifera)	LD20-71.45 ug. LD50-104.50 " LD90-186.60 "	24 hr.	 			Na-Salt of 2,4-D toxicant was fed to the bees	Jones & Connell, 1954
Harlequin fish and	TLm-7000 ma/1	24 hr.				Shell 2,4-D QR pellets	Alabaster, 1969
Rainbow trout Harlequin fish Rainbow trout	" - 4800 " " - 1160 " " - 250 "	48 hr. 24 hr. 24 hr.			,	2,4-D NaSalt	
Harlequin fish	" - 210 " " - 1.0 " " - 1.0 "	48 hr. 24 hr. 48 hr.		•		2,4-D triethanol amine 2,4-D BEE	• •
Trout	TLm-250 mg/1 " -210 mg/1	24 hr. 48 hr.	. '			2,4-D-triethanol amine salt	Alabaster, 1959
Fathead minnow eggs	TLm-5.6 mg/1 " - 1.5 mg/1	95 hr. 48 hr.			Static	Egg mortality was 100%.	Mount & Stephan, 1967
Killifish Bream (Lepomis gibbosu Bullhead (Ictalurus nebulosus)	LD50-2000 mg/k s) LD50-1000 mg/k LD50-2000 mg/k	ğ	20-25° ("	: 1 fish/	gal	2,4-D acid	Harrisson & Rees 1946
Striped bass (Morone saxatilis)	LC0 -0.1 larva 2.0 LC50-0.15 " 3.0 LC100-0.25 " 4.0) 10 10	•			2,4-D Butyl ester safe for use with fry. Nothing safe with larva.	Hughes, 1973
Green sunfish	5x10 ⁻⁴ m	60 min.				BEE	Sergeant, 1971
Gluegfll	TL50->100.0 mg " - 1.1	1 48 hr.	' 24° C			DMA BEE	Sanders, 1970b

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'Organîsms	Concentration Treatment Toxicity	Exposure	Water temp.	Loading	Testing * regime	Comments	Citation
Bluegill .	Tim 650 mg/1 550 " 600 - " 500 " 475 " 450 " 230 " 230 " 470 " 425 " 327 " 327 " 265 " 265 " 200 " 200 "	24 and 48 hŕ.		-		Weedar-64 " Crop Rider 6D-2 Crop Rider Amine 4D-2 Ortho 2,4-D 2,4-D Amine 4 Amine Weed Rhap A-4 Producers given. TLm ba on active ingredients DMA-2,4-D	Hughes & Davis, 1966 ased
Bluegill	TLm 542 mg/1 458 " 500 " 416 " 390 " .353 " 273 " 273 " 220 " 220 " 166 " 160	24. and 48 hr.	25° C.	10 fish/25	1 lake water	Commercially supplied. DMA salt; TLm based on a equivalent.	2,4-D Hughes & Davis, acid 1963 -
Bluegf11	TLm-542 mg/1 650 " 500 " 600 " 394 " 475 " 390 " 470 " 273 " 327 " 220 " 265 " 166 " 200 " 458 " 550 " 416 " 500 " 373 " 450 " 353 " 425 " 273 " 327 " 220 " 265 " 166 " 200 " 353 " 425 " 273 " 327 " 220 " 265 " 166 " 200	24 hr.		•		Weedar 64 Crop Rider 6D-2 Ortho 2,4-D 2,4-D Amine-4 2,4-O Amine Weed Rhap A-4 Weedar 64 Crop Rider 6-D-2 Ortho 2,4-D 2,4-D Amine 4 2,4-D Amine 4 2,4-D Amine Weed Rhap A-4 The toxicity figures given are the acid equivalent and the act ients of DMA 2,4-D. Producers	tive ingred-
Bluegill	TLM-188 mg/1	24 hr. 48 hr.	•			DMA-2,4-D	Hughes & Davis, 1964
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Organisms	Concent Treatment	ration Toxicity	Exposure 🕤	Water temp.	Loading	Testing regime	Comments .	Citation
Bluegill		TLm-350 mg/kg " 390 "	24 hr.	25° C	10 fish/25 1	Water from Quachita River	DMA TLm's based on 4 reps.	Davis & Hard- castle, 1959
Largemouth bass		" 375 " " 350 " " 375 " " 350 "	48 br. 24 br. 24 hr. 48 hr.	13 13 10		(24) & Cayou DeSiard (24 & 48). Water aerated.	÷	
Largemouth bass	1_mg/1 5_mg/1 10_mg/1	% mortality 80 80 50 70 40 60	72 hrs. 96 hrs. 72 hrs. 96 hrs. 72 hrs. 96 hrs.	-			DMA-2,4-D Fathead minnows showed no mortality after 96 hr. at 1, 5, and 10 mg/1	Lawrence, 1962b.
Frog		7L50-100 mg/1	96 hrs.			Static assay	Weedar 64 (DMA)	Sanders, 1970 a
Bobwhite quailyoun adult	g 5,000 mg/kg	LD50-28,000 mg/ " > 38,000	'kg <100 days		•		DMA-2,4-D. Treatment level is based on amount in diet. 1D50 based on amount eaten	DeWitt <u>et al</u> ., 1962
Adult pheasant	5,000 mg/kg >2, ⁵⁰⁰ "	" 8,250 " " >35,000 " " >16,500 "					by the time 50% mortality oc- curred. Treatment concentrat is the amount of DMA-2,4-D in the diet in mg/kg.	ion
Bodwhīte quail Young Ring neck pheasant Young Adult Mallard, young		LD50-8,250 mg/l LD50-19,780 mg/ >6,500 22,100	'kg <100 days ' <100 days	•	, .		LD50's are based on average amount eaten by the time 50% mortality oc- curred. Treatment concentration is amount of DMA-2,4-D in the diet in m	

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Organisms	Concent Treatment	ration Toxicity	Exposure	Water temp.	Loading	Testing regime	Comments *	Citations
Young Bobwhite		% mortality					· · ·	
quail	2,500 mg/kg	12 29 42	10 days 30 days 100 days		24/pen		Treatment conc. based on amount of DMA-2,4-D in feed. % mortality based on number dead after 10, 30	Stickel, 1964
Young Coturnix	2,500 "	12	10 days 30		25/pen		or 100 days. Exposure time for LC50 is time to 50% mortality.	
Adult Coturnix	5,000 "	7 27 33	10 " 30 " 100 "		15/pen		· · · · · · · · · · · · · · · · · · ·	
H 68	2,500 mg/kg L	100 .D50-56,776 mg/kg	98 days 1 94 days		.16/pen			
Young pheasant	5,000 mg/kg	8 84	10 " 30 "					
ET PE	. LD 2,500 mg/kg	150-15,998 mg/kg 28 35	19 daýs 30 " 100 "		25/pen		· · · ·	
Mallards "	LD	150->1000 mg/kg 150->2025 " 150-ca2000 "				Male & female 3-5 mo. old Male-7 mo.	Technical acid "Na salt. 4 lbs a.e./gal. amine	Tucker & Crabtree, 1970
Pheasants Coturnix Pigeons Mule deer	12	' -472(340-654)n " -668(530-842) " -668(530-842) " -400-800	ng/kg "			Male-3 mo-4 mo. Male-2 months Male & female Female 8-11 mo.	Technical acid	·
Rats, mice, guinea		" - 300-1000 mg/	/kg			Fed in H ₂ 0, olive	Dow formulations	Rowe & Hymas, 1954
pigs, rabbits		• •		•		oil, corn oil, capsule or plain	· .	
Rat, Guinea pig	L)50-375-666 mg/kg ' -1000	9	•				Johnson, 1971
Man	6.5 g						Orally administered. Convulsions precede death. Death thought to be from ventricular fibrillation.	Hayes, 1971

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APPENDIX A - COMMON AND TECHNICAL NAMES OF ORGANISMS

PLANTS

Common name

WATER PLANTAINS Arrowhead

AMARANTHS Alligatorweed

ARUMS Arrowarum

SEDGES Softstem bulrush Sedge Slender spikerush

GRÁSSES Southern watergrass

WATER MILFOILS Eurasian watermilfoil

FROGBITS Florida elodea

DUCKWEEDS Duckweed

BLADDERWORTS Bladderwort

LOOSESTRIFES Swamp loosestrife

MÀLLOWS Hibiscus Technical name

ALISMACEAE Sagittaria latifolia

AMARANTHACEAE Alternanthera philoxeroides

ARACEAE <u>Peltandra virginica</u>

CYPERACEAE Scirpus validus Cyperus spp. Eleocharis acicularis

GRAMINEAE Hydrochloa caroliniensis

HALORAGIDACEAE Myriophyllum spicatum

HYDROCHARITACEAE Hydrilla verticillata

LEMNACEAE Lemna minor

LENTIBU LARIACEAE Utricularia vulgaris

LYTHRACEAE Decodon verticillatus

MALVACEAE Hibiscus moscheutos Hibiscus militaris Hibiscus Tasiocarpos

Common name

PONDWEEDS Curlyleaf pondweed Sago pondweed American pondweed

WATERLILIES Fragrant waterlily Watershield Lotus Spatterdock

BUCKWHEATS Smartweed

PICKEREL WEEDS Waterstar grass Waterhyacinth Pickerelweed

WATER CHESTNUTS Water chestnut

CATTAILS Cattail

CARROTS Water pennywort

Technical name

NAJADACEAE Potamogeton crispus Potamogeton pectinatus Potamogeton nodosus

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NYMPHAEACEAE Nymphaea odorata Brasenia schreberi Nelumbo lutea Nuphar advena

POLYGONACEAE Polygonum amphibium

PONTADERIACEAE Heteranthera dubia Elchornia crassipes Pontaderia cordata

TRAPACEAE Trapa natans

TYPHACEAE Typha latifolia

UMBELLIFERAE Hydrocotyle umbellata

ANIMALS

Common name

MOLLUSKS Eastern oyster

CRUSTACEANS

DAPHNIA Daphnia

OSTRACODS Seed shrimp

ASELLIDAE Sowbug

ASTRACIDAE Crayfish

PALAEMONIDAE Grass shrimp

GAMMARIDAE Scud

DECAPODS Blue crab Fiddler crab Brown shrimp

INSECTS

Pteronardidae Stonefly naiads

ACRIDADAE Grasshoppers

APIDAE Honey bees Technical name

MOLLUSCA Crassostrea virginica

CRUSTACEA

DAPHNIDAE Daphnia magna

Asellis spp

Orconectes spp

Palaemonetes kadiakensis

Gammarus lacustris

DECAPODA Callenectes sapidus Uca pugnax Penaeus aztecus

INSECTA

Pteronarcys californica

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Romalea microptera

Apis mellifera

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FISHES

Common name

TROUTS Rainbow trout

MINNOWS AND CARPS Squaw fish Fathead minnows

SUCKERS Lake chubsuckers

FRESHWATER CATFISHES Channel catfish Brown bullhead

KILLIFISHES Longnose killifish

TEMPERATE BASSES Striped bass

SUNFISHES Green-sunfish Pumpkinseed Bluegill Smallmouth bass Largemouth bass

AMPHIBIANS

FROGS Western Chorus Frog

BIRDS

PARTRIDGES Red partridge Gray partridge Technical name

SALMONIDAE Salmo gairdneri

CYPRINIDAE Ptychocheilus spp Pimephales promelas

CATOSTOMIDAE Erimyzon sucetta

ICTALURIDAE Ictalurus punctatus Ictalurus nebulosus

CYPRINODONTIDAE Fundulus similis

PERCICHTHYIDAE Morone saxatilis

CENTRARCHIDAE Lepomis cyanellus Lepomis gibbosus Lepomis macrochirus Micropterus dolomieui Micropterus salmoides

HYLIDAE Pseudacris triseriata

PERDIDIDAE <u>Perdix rufa</u> Perdix perdix

Common name

PHEASANTS Pheasant

RAILS Florida gallinule

GRACKLES Boat-tailed grackle Technical name

PHASIANIDAE Phasianus colchicus

RALLIDAE Gallinula chloropus

ICTERIDAE Cassidix mexicanus

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APPEND IX

PROPOSED LABEL COPY (front panel)

2,4-D ANINE WEED KILLER

FOR

CONTROL OF WEEDS ON IRRIGATION CANAL BANKS

CAUTION

KEEP OUT OF THE REACH OF CHILDREN

Active Ingredient:

*Dimethylamine salt of 2,4-dichlorophenoxyacetic acid . . .

Inert Ingredients:

*2,4-dichlorophenoxyacetic acid equivalent 41.2% by weight

or 4 pounds per gallon

from Reproduced available cop

Manufactured for use of the Bureau of Reclamation and cooperating water users organizations and the Bureau of Sport Fisheries and Wildlife, U.S. Department of the Interior; Agricultural Research Service, U.S. Department of Agriculture; and the Corps of Engineers, U.S. Department of Defense.

PROPOSED LABEL COPY (Interpanel)

GENERAL DIRECTIONS

Recommendations made are for noncrop use to control annual and perennial broadleaf herbaceous weeds and woody plants on banks of irrigation canals and irrigation drainage ditches.

<u>Control of Broadleaf Annual and Perennial Weeds</u>. Apply 1 to 2 quarts of 2,4-D Amine Weed Killer per acre in a sufficient volume of water to provide good spray coverage of vegetation;20 to 100 gallons per acre id usually sufficient. Treat when weeds are young and actively growing before the bud or early bloom stage. For harder-to-kill weeds, a repeat spray may be needed after 3 to 4 weeks for maximum results. Apply no more than 2 treatments per season.

<u>Control of Woody Brush and Patches of Perennial Broadleaf Weeds</u> gallon of 2,4-D Amine Weed Killer in 150 gallons of water. Wet follage thoroughly using about 1 gallon of solution per square rod.

Spraying Instructions. Apply with low-pressure (optimum 10 psi and not over 40 psi) high-volume power spray equipment mounted on trucktractor, or boat. Spray operation is to be made traveling upstream to avoid concentrating chemical in the water. Spray when the air is fairly calm (5 mph or less). Do not use on small canals (less than 10 cubic feet per second), when water will be used for drinking purposes. Boom spraying onto the water suface must be held to a minimum. Much spraying shoreline weeds, avoid spraying over the water in the stream accept to cover shoreline vegetation. Allow no more than 2 feet of spray onto water with an average of less than 1 foot of spray over the water surface to prevent introduction of greater than negligible amounts of chemical in water. Avoid cross-stream spraying to the opposite bank. Do not spray the banks on both sides of the stream at the same time. Spray each bank separately.

Convert pounds 2,4-D acid equivalent recommendations into terms of 2,4-D AMINE WEED KILLER by the following table:

2,4-D	1 15	3/4 1b	1/2 15 3/8 15 1/4 15	1/6 16
2,4-D AMINE WEED KILLER	2 pt	1-1/2 pt	l pt 3/4 pt 1/2 pt	3/8
• • • • • • • • • • • • • • • • • • •				

PROPOSED LABEL COPY. (right panel)

Typical ditchbank weeds controlled by 2,4-D Amine Weed Killer are as follows:

Annual and Blennial Weeds

beggarsticks bull thistle cocklebur kochia lambsquarter lettuce (wild) mallow morningglory marsh elder mustards parsnip peppergrass pigweed prickley lettuce primrose ragweed Russian thistle sunflower vetch

Perennial Weeds

bindweed catnip Canada thistle dandelion dogbane dock goldenrod hoary cress nettles plantains water hemlock

Du not allow dairy animals to graze on treated areas for at least 7 days after spraying. Water within treated banks should not be fished. Harmful if swallowed. Avoid contact with skin, eyes, or clothing. Avoid spray drift to susceptible plants, such as: cotton, beans, tomatoes, and ornamentals, as this produce may cause injury. Coarse sprays are less likely to drift. Thoroughly clean spray equipment with suitable chemical cleaner before using for other purposes (or do not use spray equipment for other purposes). Do not store near fertilizers, seeds, insecticides, or fungicides.

DO NOT USE THIS HERBICIDE AT RATES OR FOR METHODS OF APPLICATION OTHER THAN THOSE RECOMMENDED ON THIS LABEL TO PREVENT SERIOUS CONTAMINATION OF IRRIGATION WATER. DO NOT ADD INGREDIENTS OTHER THAN WATER TO THE HERBICIDE FORMULATION.

DIRECTIONS FOR USE OF 2,4-0 AMINE WEED KILLER FOR CONTROL OF WEEDS ON IRRIGATION CANAL BANKS

The amine formulation of 2,4-D has been used for over 20 years to control broad-leaved weeds on irrigation canal banks of the western United States. The following directions summarize treatment procedures used by the Bureau of Reclamation and cooperation water users organizations.

Amount of Rates of Application

Broad-leaved Annual and Perrennial Weeds. Apply 1 to 2 quarts per acre in sufficient volume of water to provide adequate coverage of vegetation; 20 to 100 gallons per acre is usually sufficient. Treat when weeds are young and actively growing before the bud or early bloom stage. Appeat treatments may be needed in 3 os 4 weeks for maximum control. For control of woody brush and patches of perennial broadleaf weeds, mix 1 gallon of 2,4-D with 150 gallons of water. Apply with hand gun or similar treat of spot application equipment. Wet foliage thoroughly using about 1 gallon of solution per square foot.

Frequency and Timing of Application

Most irrigated areas of the western United States require only one application of 2,4-D amine in spring to early summer to control broad-lawed ditchbank weeds. Treatment timing will vary locally but should be under when the weeds are young and actively growing before the bud or early bloom stage. A second application may be required after 3 or 4 weeks for maximum results or when higher than normal rainfall encourages a second crop of ditchbank weeds. The second seasonal treatment can often be managed by spot treatment of the weedy vegetation, particularly perennial weeds or woody plants. Intervals between treatments will vary according to local conditions ranging from as little as 3 to 4 weeks or as much as 6 weeks to 2 months.

In areas where 2,4-D sensitive crops such as cotton are grown, make the first general ditchbank treatment so that it will precede emergence of the crops concerned.

Restrictions on Use and Application Procedures - Special Application

Apply with coarse low-pressure (optimum 10 psi and not over 40 psi) power spray equipment mounted on truck, tractor, or boat. Spraying operations are to be made traveling upstream to avoid concentrating chemical spray that gets into the stream. To prevent introduction of greater than negligible levels of chemical into the stream, boom spray and manually operated spray gun patterns should be made so as not to spray over the water surface more than 24 inches from the edge of the water.