

observed in the first discharge were 0.2-0.5 and 3-5 ppm in the two channels respectively. The latter figures are an order of magnitude higher than the permitted maximum. However, the high concentrations were attenuated rapidly as discharge increased and contributed relatively little to the total displacement of diuron into the drainage system. Alternative estimates of contamination, obtained by calculating average concentrations integrated over the first 4 hours of discharge and assuming complete mixing, were about 0.1 and 0.5 ppm. In the latter trial, a longer discharge period would be required to further dilute average concentrations to meet the 0.1 ppm tolerance. In practice, however, dilution with untreated water in Mirrool Creek and Barren Box Swamp before re-use is expected to reduce concentrations to negligible levels.

In these two trials, by discharging the contaminated water to waste, about 9% and 61% respectively of the residual diuron was removed to the drainage system and diuron concentrations in the water at the resumption of irrigation were reduced to about 0.04 and 0.25 ppm respectively. The effect of these residue levels on crops, particularly flood-irrigated germinating rice, may need further investigation.

ASPECTS OF THE USE OF ACROLEIN IN WATER

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Acrolein (acrylaldehyde) is widely used by government authorities to control submerged aquatic plants in lakes and irrigation channels. This paper reports on some recent observations on the fate of acrolein in water and summarizes the progress being made towards rationalizing the use of this herbicide with regard to economy and safety.

The reaction of acrolein with water was investigated initially in laboratory experiments in which the processes of volatilization and adsorption were excluded. A metabolite which is relatively non-volatile and non-phytotoxic was detected using a colorimetric reaction which is specific for carbonyl groups. At normal concentrations of acrolein the intermediate product

was transient and no hazards are anticipated in practice. Experimental observations and published reports suggest that polycondensation or reversible hydrolysis to hydroxypropionaldehyde occurs, followed by microbiological oxidation to a carboxylic acid.

The decay of acrolein in buffered solutions approximated to first order kinetics and was pH-dependent, with rate constants of 0.0139 hr^{-1} at pH 6.6 to 0.0182 hr^{-1} at pH 8.6, corresponding to half lives of 50 and 38 hours respectively. In comparison, rates of dissipation observed in flowing water in irrigation channels were an order of magnitude greater, suggesting major losses by volatilization and/or adsorption onto weeds or sediment.

In irrigation channels, the herbicide is usually mixed with the water at a convenient point upstream and allowed to flow over the submerged weed beds. It is proposed that in any given length of channel effective weed control should be achieved by maintaining the product of concentration and contact time above the minimum lethal dose (E_L) and that an addition above this minimum is required to compensate for dissipation. Observations by the United States Bureau of Reclamation (USBR) suggest that acrolein dissipation from flowing water is a first order process. It follows that the theoretical dosage (E_0) required at the injection point may be calculated using equation (1) :

$$E_0 = E_L \exp(Kt) \quad \text{————— (1)}$$

where K is the first order rate constant for dissipation and t is the time for the herbicide to travel through the treated length.

Values of K obtained for six irrigation channels in Australia agreed quite closely with data from two trials reported by the USBR; the mean was 0.16 hr^{-1} , corresponding to a half-life of 4.4 hours.

Preliminary information on E_L was obtained by calculating the residual dosage at the downstream limit of several successful treatments. The data suggested that the minimum dosage of acrolein required for weed kill was largely independent of the separate values of concentration and time providing the product was maintained at approximately 2 ppm.hr.

It is recognized that several factors (including weed species and density, temperature and turbulence) have not been considered and that experience and intuition are essential in assessing appropriate dosages for particular situations. In

practice, routine weed control in several supply channels in the M.I.A., where ribbonweeds and pondweeds dominate, required acrolein additions three to eight times greater than calculated from equation (1). In contrast, in the irrigation districts of southern New South Wales control of *Elodea canadensis* has been achieved with dosages several orders of magnitude smaller than predicted. Explanations for these discrepancies may be related to differences in weed species in the two areas. Their relative sensitivities to acrolein and effects on the hydrological characteristics of the channels are now being investigated.

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