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A LABORATORY ANALYSIS OF KEPONE DEPURATION BY SPOT, LEIOSTOMUS XANTHURUS

by Marion Y. Hedgepeth, Robert T. Doyle, and Linda L. Stehlik

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INTRODUCTION

Dectectable residue levels of the pesticide Kepone have been found in resident and migratory finfishes from the James River, Virginia (Bender et al., 1977). As a result, the James River was closed to commercial finfishing in early 1976 (with the exceptions of channel catfish and American shad for a short period of time}. In addition, the United States Food and Drug Administration established an action level of O. 3 ppm of Kepone in finfishes utilized for human consumption.

Residue levels of Kepone in fishes such as spot, Atlantic croaker, bluefish, striped bass and American shad were investigated to determine if these migratory species present a health hazard to the public in areas beyond the James River system. Bender et al. (1977) found that residue levels in finfishes were dependent upon the species of fish and the length of residence in the James River. Also, they maintained that residue levels in finfishes declined as distance from the James was increased.

In 1977, additional Kepone studies were begun at the Virginia Institute of Marine Science (VIMS) to determine the rates of Kepone depuration in contaminated fishes from the James River. In a laboratory analysis of Kepone depuration by Atlantic croaker, Micropogonias undulatus, Doyle et al. (In Press) observed a significant drop in Kepone concentration in the 24th week sample. Furthermore, it was noted that this significant change in mean residue levels coincided with a rise in the ambient water temperature to above 15°C; however, additional studies were needed to confirm this relationship. In our study we chose to observe the effect of temperature on the rate of Kepone depuration by contaminated spot, Leiostomus xanthurus, from the James River.

Bender et al. (1977) reported a mean Kepone level of 0.81 ppm in spot from the James River and the lower Chesapeake Bay. This was attributed to the biomagnification of Kepone through the food chain and/or direct uptake from the water {Schimmel and Wilson, 1977). Bahner et al. (1977) confirmed this belief in a study in which spot were fed live mysids which had grazed on Kepone laden brine shrimp. Consequently, the spot accumulated concentrations of Kepone near that in their diet. Spot which had been exposed to Kepone in water were able to reduce Kepone residues in their tissues to 30-50 percent within 24-28 days in Kepone free water.

MATERIALS AND METHODS

On November 11, 1977, approximately 550 spot were obtained from the lower James River with a 30 foot semi-balloon trawl. They ranged in size from 86 mm in fork length and 8 grams in total weight to 233 mm in fork length and 165 grams in total weight. They were transported to VIMS and distributed randomly to four circular four-foot tanks {approximately 200 gallons each). All tanks were supplied with Kepone-free York River water in a flowthrough system and strong aeration. Fish were fed chopped Keponefree squid daily (8-12 percent body weight).

After one nonth of acclimation at ambient river temperature, three of the tanks were heated with water from a large header

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tank equipped with two 220-volt heaters. Heated and unheated water were combined in mixing boxes, and flow rates were adjusted so that temperatures were maintained at approximately 22°, 17° and 12°, respectively, in the three experimental tanks. The fourth tank remained at ambient temperature except for a period between January and March in which a small heater was added to keep the water above 5°C. All tanks were insulated with cotton padding and aluminium foil. At times, temperatures in the heated tanks fluctuated as a result of sand clogging the pipes and disrupting the established flow rates. In the spring, river water temperature rose until, in June, all tanks were above 22°C. Throughout the experiment, salinity and dissolved oxygen were measured weekly, while temperatures were taken daily. In addition, water samples were analyzed periodically for Kepone.

During the acclimation period, two samples of twenty fish (five per tank) were sacrificed on Day O and Day 31 and analyzed for Kepone. Thereafter, biweekly samples of ten spot per tank were taken for several weeks. Since, it appeared that the spot were depurating slowly, the time interval was increased later to four weeks. Kepone concentrations (whole body, micrograms/gram, µg/g; and parts per million,ppm) were determined by electron capture gas chromatography. Mass spectrometry was utilized when concentrations were high. For the exact methodology of the chemical analysis see Appendix A.

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RESULTS

Contaminated spot depurated considerable amounts of Kepone within a period of two hundred days (Fig. 1, dotted line). A mean Kepone concentration of 1.63 ppm $(N = 20)$ was found for spot sacrificed on the day of collection $(t = 0)$; whereas, a mean Kepone concentration of 0.45 ppm (N = 30) was found for spot sacrificed two hundred days later $(t = 200)$. In a statistical analysis utilizing mean concentration values for the periods $t = 0$ and $t = 31$, spot eliminated approximately 53 percent of the Kepone residues in their tissues; however, 95 percent confidence interyals were broad during this period of acclimation (Fig. 1). Bahner et al. (1977) reported residue declines of 30-50 percent in spot after **24-28** days in Kepone-free water.

Further demonstration of Kepone depuration in the spot was provided by Pearson correlation coefficients (r) of -0.7252 $(p = .001)$ for the variables Kepone concentration with total number of days in tank (t) and -0.6231 (p = .001) for the variables Kepone concentration with total number of days in tank squared (t^2) . A multiple regression analysis of mean Kepone concentrations by t of each tank produced the following regression equation: Kepone concentration+ l.48183-0.145133 t + -5 2 $4.5612 \times 10^{-5} t^2$, $r^2 = 0.6948$, p = .001 and the regression curve (solid line) of figure 1.

The levels of Kepone concentrations in spot varied by period (t) and by tank (Appendix B). The appearance of a rise in concentrations after day 31, when heat was applied to tanks

1, 2 and 3, was attributed to no net loss of Kepone while spot were reacclimating to the rise in temperatures (Appendix C} and to possible random samples of highly contaminated fish. Thus, the actual acclimation period for the spot might be considered as the first sixty or seventy days. Once the tanks had achieved their respective temperatures (between days 59 and 73) mean Kepone concentrations in the spot samples began to change. Spot in the warmer tanks demonstrated lower mean Kepone concentrations. In fact, spot in Tank 1 (22°C} generally exhibited lower concentrations (Fig. 1). Unfortunately, Tank 1 was discontinued after a short period of excessively high temperatures which caused a high mortality among the spot.

No significant.relationships were found between the level of Kepone residues (ppm} in spot and the length, weight, or sex of the fish. Values of micrograms of Kepone per gram fish (Appendix C} produced comparable results in statistical analyses. Furthermore, no substantial growth was observed in the spot during the study period. Thus, dilution of Kepone residue in the tissues due to growth was not a factor in the rate of depuration.

Although spot and Atlantic croaker are closely related species, Kepone concentrations in spot were generally higher than those Doyle et al. (1978} found in Atlantic croaker. Both species were collected from the James River at approximately the same time of year (October-November} although in different years (1976-1977). Initially, spot depurated Kepone at a faster rate than Atlantic

croaker (Fig. 2). In fact, there was no significant decrease in Kepone levels of Atlantic croaker until after a period of fiftysix days. On the other hand, Atlantic croaker that were sacrificed after a period of one hundred and fourteen days, depurated at a slightly faster rate than spot from Tank 4 (ambient temperature).

CONCLUSIONS

Spot, like other fishes, depurated Kepone at a slower rate than some invertebrate species (see: Bahner et al. 1977). A mean loss in Kepone residues of 72 percent occurred between the initial spot sample $(t = 0)$ and the eight spot sample $(t = 200)$. A plot of the variables, mean Kepone concentration by period (t) (Figure 1) demonstrated the fact that a negative relationship existed between Kepone concentration in spot and the amount of time a spot was allowed to depurate in Kepone free water. Nonetheless, only 30 percent of the spot $(N = 309)$ utilized in the test were below the established action level for human consumption (0.3 ppm). Therefore, it appears that it would be impractical to remove spot from a contaminated area and to maintain them in a holding facility for the purpose of depuration and later commercial sale. Whether wild spot from the James River and the lower Chesapeake Bay can or cannot eliminate Kepone from their bodies while in the overwintering grounds of Virginia and North Carolina is still another question. To answer this question and other management questions, we would have to establish the Kepone levels in fish from offshore and returning populations which would be very difficult and costly.

Temperature was an important factor in the rate of Kepone depuration in spot. Spot held in warmer water exhibited lower mean Kepone concentrations: however, we were unable to observe the effect of the lower temperature extremities for any length of time in the cooler tanks because of the rise in temperature during the later spring months. In response to the warmer temperatures, Kepone concentrations in spot indicated that the rate of elimination of Kepone from body tissues is probably a function of the rate of an individual's metabolism. Thus, an increase in the matabolic rate as a result of an increase in body temperature may cause an acceleration in the depuration rate; however, it may not be apparent until after a period of acclimation.

It is regretable that the cost of Kepone body burden analysis is so high that sample sizes must remain small. In the future, we should take a closer look at the processes of uptake and accumulation of Kepone in eggs, larvae, juvenile and adult life stages. Also, we must have a better understanding of how Kepone concentrations in fish are related to uptake, accumulation and the lipid composition of fish.

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Figure 1. Depuration of Kepone from Spot, Leiostomus xanthurus.

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Appendix A

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Chemical Analysis for the Pesticide Kepone

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Appendix A

Chemical Analysis for the Pesticide Kepone

Whole fish were ground in a meat grinder into hamburger consistency. A mixture of anhydrous sodium sulfate and Quso^R G-30 (precipitated silica, Philadelphia Quartz Co.) was added for desiccation. The proportions of sample to the desiccants were: 30 g fish - 54 g Na₂ SO₄ - 6 g Quso. Then samples were frozen at -5°C for 24 hours to rupture the cells. After thawing the desiccated samples were ground with a blender to a powdery consistency and transferred to pre-extracted paper thimbles for Soxhlet extraction. Extraction was carried out using 1:1 ethyl ether-petroleum ether for 16 hrs. Extracts were then concentrated by evaporation and cleaned by activated fluorisil column chromatography (EPA, 1975). The Kepone containing elutriate was analyzed by electron capture gas chromatography utilizing packed columns with one or more of the following liquid phases: 4% SE-30 + 6% OV 210; 1.5% OV-17 + 1.95% QF-1 + 3% CV-1. On occasion, when concentrations were sufficiently high, Kepone presence was confirmed by mass spectrometry.

Appendix B

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Descriptive Statistics of Kepone Concentrations Broken Down by Tank and Sampling Period

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Appendix C

Water Quality Analysis During the Experimental Period

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Appendix C2. Dissolved oxygen concentrations for all four tanks.

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Appendix C3. Average salinity for all four tanks.