Reproductive efficiency in mink (*Mustela vison*) treated with the pesticides lindane, carbofuran and pentachlorophenol

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Mink are carnivores of agroforestry fringe habitats and are exposed to pesticides that biomagnify within the food chain. Some pesticides are thought to disrupt reproductive and endocrine functions. In Expt 1, four groups of mink (*n* = 10) were fed either a control diet, or diets treated with lindane (1 mg kg$^{-1}$ day$^{-1}$), carbofuran (0.05 mg kg$^{-1}$ day$^{-1}$) or pentachlorophenol (1 mg kg$^{-1}$ day$^{-1}$) from before breeding until weaning. Mink were mated twice, at 7–8 day intervals. The treatments had no effect on the proportion of mink accepting the first mating; however, lindane and pentachlorophenol caused a decrease in the percentage of females accepting the second mating. Lindane and pentachlorophenol caused a decrease in whelping rate, although litter size was not affected. Carbofuran had no effect on fertility. Mink that mated only once had a lower whelping rate than mink that mated twice; therefore, it could not be determined whether the decreased whelping rates were due to the lack of a second mating or to increased embryo loss. In Expt 2, two groups of mink (*n* = 15) were fed a control diet or a diet treated with lindane (1 mg kg$^{-1}$ day$^{-1}$) from before mating until weaning. Mink were mated twice on two consecutive days. Lindane did not affect mating response at either mating. Whelping rate, but not implantation rate, was decreased by the lindane treatment. The proportion of embryos lost after implantation (implantation scars not represented by kits at whelping) was increased by the lindane treatment. In conclusion, both lindane and pentachlorophenol decreased fertility in mink, and the lindane effect was primarily a result of embryo mortality after implantation.

Introduction

A number of pesticides may pose a hazard to wild and domestic animals and humans owing to their ability to alter the normal functioning of mammalian endocrine and reproductive systems (Colborn et al., 1993). The reproductive effects of endocrine disrupting chemicals have been identified as a critical research priority in a number of recent government workshops (see Kavlock et al., 1996).

Mink are carnivores of agroforestry fringe habitats and are exposed to endocrine disrupting environmental contaminants that biomagnify within the food chain (Giesy et al., 1994; Heaton et al., 1995). Mink are seasonal breeders with induced ovulation (Hansson, 1947; Sundqvist et al., 1989). In mammals, the early embryonic period may be especially susceptible to disruption by pesticides (Mably et al., 1992; Gray et al., 1994; Guo et al., 1995) and mink may be particularly sensitive owing to their system of delayed implantation; hence mink are often used in studies of reproductive toxicology (Aulerich et al., 1973; Calabrese et al., 1992; Crum et al., 1993; Backlin and Bergman, 1995; Heaton et al., 1995).

Organochlorine pesticides are considered persistent environmental contaminants and are routinely detected in air, dust sediments, ground water and body tissues of animals and humans (Baekloek et al., 1985; Foster, 1995; Sonawane, 1995; Waite et al., 1995; Tate and Heiny, 1996; Thompson and Treble, 1996). Carbamate pesticides are less persistent than organochlorines, but are still present in air, food and ground water and are generally more toxic (Gupta, 1994). Wild animals may be particularly at risk from chronic exposure to subacute concentrations of pesticides as they live in areas of treated farmland and drink water contaminated by agricultural runoff.

In the present study, the effects of three pesticides on reproduction in mink were examined. The organochlorines used were lindane (the gamma isomer of hexachlorocyclohexane) and pentachlorophenol (PCP), and the carbamate used was carbofuran. These pesticides were chosen because they are used extensively in the prairies, and their toxic effects have been previously investigated (Spencer, 1982; Ecobichon, 1991). Lindane is an insecticide present in a variety of agricultural, medical and veterinary products that are used widely, particularly in developing countries, for the control of pests that transmit vector borne diseases (Ecobichon, 1991). Pentachlorophenol has been used extensively as a general biocide and wood preservative (Exon, 1984). Carbofuran is a broad spectrum insecticide used in agriculture and forestry throughout the world (Gupta, 1994). Previous mammalian studies of endocrine and reproductive effects of pesticides are

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limited (particularly for carnivore species) (O'Brien et al., 1993). However, studies in rodents have indicated that lindane (Sircar and Lahiri, 1989; McNutt and Harris, 1994) and pentachlorophenol (Welsh et al., 1987) alter embryonic development and oestrous behaviour (Uphouse, 1987). Carbofuran has generally been shown to have no effects on fertility or embryo development in rodents, but chronic treatment with large doses caused testicular degeneration in male beagles and hyperplasia and hydrometria in female beagles (for review, see Gupta, 1994). Other carbamate pesticides have been shown to affect fertility in sheep (Grendon et al., 1994). The doses used in the present study were based on previous studies (Spencer, 1982; Ecobichon, 1991) and our own preliminary study, and were chosen to be low enough to avoid overt toxicity but high enough to show any potential effects on the endocrine and reproductive systems.

In these studies, our specific goal was to look at the effect on fertility of lindane, pentachlorophenol and carbofuran fed to mink from before breeding until their young were weaned. A second experiment was designed to characterize the timing of the embryo loss observed in mink treated with lindane, using a revised mating protocol.

**Materials and Methods**

**Animals**

Female mink kits (9 months old, demibuff colour phase, n = 70; Baldwins mink farm, Starbuck, Manitoba) were housed individually under natural conditions of daylight and temperature at 52°N latitude. They were fed a pelleted complete ration (Pike Lake Fox Feed, Federated Co-op Feed Mill, Saskatoon, SK) supplemented with 5–20% chicken offal (depending on the outside temperature) mixed into a paste with water. Drinking water was provided ad libitum. Care and treatment of animals was carried out according to Canadian Council for Animal Care guidelines.

**Preliminary study**

Doses to be used were based on previous data for rats (Spencer, 1982; Ecobichon, 1991), which were confirmed in preliminary acute dosing studies in mink. On the basis of the results of these tests the following doses of pesticides were administered in the subsequent experiments: lindane and pentachlorophenol, 1 mg kg⁻¹ body weight day⁻¹; carbofuran, 0.05 mg kg⁻¹ body weight day⁻¹.

**Experiment 1**

Forty female mink were weighed and divided into four weight-matched groups (n = 10). The mink were fed either an untreated diet, or a diet treated with lindane (Sanex Agro, Dundos, Ontario), carbofuran (Chem Agro, Etobicoke, Ontario) or pentachlorophenol (Sigma, St Louis, MO). Individuals in each group were housed in adjacent cages with at least one empty cage between each group. The pesticides were evenly sprayed onto the pelleted ration and enough treated feed was prepared in this manner on a weekly basis (thereafter being mixed with the chicken and water each day before feeding). Treated feed was kept in darkness at ≤4°C before use. Treated or untreated feed was given throughout the study commencing 3 weeks before breeding and continuing until weaning (8 weeks post partum). Body weight was recorded every week during the pre-mating period and every 2 weeks during the postpartum period. Feed intake was also monitored at these times by weighing the amount of wet feed given and subtracting any feed wasted.

Mink were mated according to normal ranch practice of an initial mating, followed 7 or 8 days later by a second mating. A similar double-mating strategy has been shown to improve pregnancy rates in yearling females (Lagerkvist, 1992). Each day from 5 March to 20 March, one or two females from each group were selected randomly and mated. Single females were placed in the cage of a single breeding male (n = 10) and left with the male for 20 min. If copulation had been initiated, they were left for a further 30 min. If copulation did not take place, the females were placed with another male and if no copulation occurred on the second attempt, the female was returned to her own cage. The procedure was repeated with any unmated females on two further occasions a number of days later and, if mating still did not occur, the females were classified as non-breeding females. The mated females were mated again to a different male 7 days later. If the second mating was rejected, these females were retried 1 or 2 days later; refusal at this mating meant they were only mated once. If copulation was observed, a vaginal lavage was taken to ensure motile spermatozoa had been deposited. Males were only allowed to mate a single female on any given day.

During the whelping period the mink were checked at least twice per day to determine whether they had whelped (by listening for young in the nest box). If whelping was suspected, the litter size was counted quickly (without disturbing the nest) while the mother was kept away from the nest (since any disturbance immediately after whelping can cause cannibalization of the young). The kits were then counted and weaned every 2 weeks until weaning. At weaning (8 weeks post partum), the adult mink were separated from their litters, and sedated with ketamine (Vetlar, Vet pharm, London, Ontario) and xylazine (Rompun, Bayvet, Etobicoke, Ontario). A blood sample was collected by cardiac puncture, and the animals were killed by an intracardial injection of T61 (Hoechst, Regina, SK). The animals were then necropsied. Samples of ovary, oviduct, uterus, pituitary, adrenal, thymus, pancreas, kidney, liver, lung, and heart were fixed in normal buffered formalin, dehydrated, and embedded in paraffin wax for sectioning, mounted and stained with haematoxylin and eosin before histopathological examination.

**Experiment 2**

Thirty female mink were weighed and divided into two weight-matched groups. They were fed a control or lindane (1 mg kg⁻¹ day⁻¹) treated diet from 6 weeks before mating until weaning, when the adults were killed. Feed was treated in the manner outlined above.

Mink were mated using a revised protocol in an attempt to avoid the confounding effects of treatment-related changes in
the mating response at the second mating seen in Expt 1. The protocol consisted of an initial mating, followed the next day by a second mating. Commencing on 10 March and continuing daily until 25 March, one to nine females from each group were selected randomly and mated. Single females were placed in the cage of a single male (n = 14). If copulation did not take place after 20 min, the female was tried with one other male. This procedure was repeated with any unmated females on up to four consecutive occasions at intervals of 3 days. If copulation did occur, the female was mated again to a different male the next day. If this second mating was rejected, these females were also retried with another male; refusal at this mating meant that they were only mated once. Motile spermatozoa were examined under the microscope from a vaginal lavage to confirm successful copulation. Males were only allowed to mate with a single female on any given day.

Adults and litters were treated as in Expt 1 post partum, except that litters were weaned at 10 weeks of age and weights of uterus, ovary, kidney and liver were also recorded.

**Hormone assays**

Blood samples were allowed to clot overnight at room temperature; serum was separated by centrifugation at 1500 g for 10 min, and stored at −20°C until assayed. Serum concentrations of progesterone and oestradiol were measured as described by Ravindra et al. (1994). The limits of sensitivity were 0.3 nmol l⁻¹ and 3.7 pmol l⁻¹, respectively. The intra-assay and interassay coefficients of variation for reference sera with progesterone concentrations of 1.8 or 5.6 nmol l⁻¹ were 7.2% and 9.0% or 5.3% and 8.1%, respectively, and for oestradiol concentrations of 15.9 or 25.8 pmol l⁻¹ were 3.0% and 8.1% or 5.1% and 12.3%, respectively. Serum concentrations of cortisol were measured as described by Kingsbury and Rawlings (1993). The limit of sensitivity was 12.4 nmol l⁻¹. Serum concentrations of thyroxine were measured as described by Allen et al. (1995). The limit of sensitivity was 5.0 nmol l⁻¹. The intra-assay and interassay coefficients of variation for a reference serum with a cortisol concentration of 40.0 nmol l⁻¹ were 16.0% and 13.4%, and for reference sera with thyroxine concentrations of 27.0 or 51.0 nmol l⁻¹ were 11.1% and 8.1% or 8.2% and 5.6%, respectively.

**Histopathological examination**

Each tissue was examined for any potential treatment-related lesions. When a characteristic of a tissue was observed to be outside of the normal histological limits, the severity of the lesion was evaluated as follows: 0–1 = minimal, 1–2 = mild, 2–3 = moderate, 3–4 = marked. Within any of the above categories, the extent of the lesion within the tissue was scored as follows: for example, 1.25 = mild focal, 1.50 = mild diffuse, 1.75 = mild multifoc. Therefore, each tissue was given a score ranging from 0 to 4.

**Statistical analysis**

Whelping rate is expressed as the number of mated mink that subsequently whelped. Litter size is expressed as the mean number of kits at birth, where data from only the females that produced litters are considered. The percentage embryo loss after implantation was calculated for each litter as the number of kits born subtracted from the number of implantation sites divided by the number of implantation sites. The proportion of mink in each treatment group mating, whelping, or exhibiting implantation sites was compared with the proportion in the control group by chi-squared analysis (True Epistat, Epistat services, Richardson, TX). The effects of treatment on litter size, implantation rate, embryo loss and hormone concentrations were examined by Student’s t test (single comparisons of each treatment group with the control group; True Epistat). Proportions are expressed as percentages and all other data are presented as mean ± SEM. The severity scores for the histopathological lesions were analysed by analysis of variance followed by Dunnett’s ad hoc test (Sokal and Rohlf, 1969).

**Results**

**Experiment 1**

No overt signs of toxicity were noted. During the post-partum period, mink treated with lindane appeared to be more hyper-reactive, especially when faced with the stress of handling, but this was not tested in a quantitative fashion. The pesticides did not cause any significant decrease in body weight during the premating or postpartum periods. Body weights for controls, and mink treated with lindane, carbofuran, and pentachlorophenol at the start of the study were 1139 ± 51, 1101 ± 34, 1119 ± 50, 1136 ± 45 g, respectively, and at weaning were 1019 ± 48, 1119 ± 34, 1023 ± 18, 1069 ± 21 g, respectively. Feed intake was similarly unaffected by the pesticide treatments, although it did vary greatly with environmental temperature changes and demands of lactation (overall mean of all time points were: control, 256 ± 31; lindane, 244 ± 33; carbofuran, 240 ± 37; pentachlorophenol, 249 ± 31 g wet feed day⁻¹).

**Fertility.** Fertility data are summarized (Fig. 1). The treatments had no significant effect on the proportion of mink that accepted the first mating or the date on which the mink first mated. However, the proportion of the mink mated at the first mating that subsequently accepted the second mating was significantly decreased in the groups treated with lindane (P < 0.001) and pentachlorophenol (P < 0.01). This resulted in the average last mating date being significantly earlier (P < 0.05) in the group treated with lindane (control, March 14.3 ± 0.8; lindane, March 10.1 ± 1.4; carbofuran, March 13.1 ± 1.3; pentachlorophenol, March 11.7 ± 0.4). The proportion of mated mink that whelped subsequently was reduced by the lindane (P < 0.01) and pentachlorophenol (P < 0.01) treatments. However, the proportion of mated mink that had implantation sites visible when killed at weaning and the mean number of implantation sites per mink (control, 6.75 ± 1.22; lindane, 6.60 ± 1.41; carbofuran, 5.44 ± 0.90; pentachlorophenol, 4.55 ± 1.31) did not differ significantly between groups. Mink that mated only once had a significantly lower whelping rate than mink mated twice (overall, 53% versus 86%, P < 0.001). There was no significant effect of pesticide
treatment on the proportion of mink with implantation sites that whelped subsequently (control, 88%; lindane, 75%; carbofuran, 100%; pentachlorophenol, 71%); the mean percentage of embryos lost after implantation in each litter (control, 40.5 ± 12.6%; lindane, 61.4 ± 14.3%; carbofuran, 43.0 ± 9.6%; pentachlorophenol, 46.7 ± 16.3%); or the litter size of mink that whelped (control, 4.45 ± 1.00; lindane, 3.80 ± 1.50; carbofuran, 3.38 ± 0.60; pentachlorophenol 3.40 ± 0.68) (P > 0.1). Mink in the groups treated with lindane and pentachlorophenol whelped later (P < 0.05) than control mink (date of whelping ± days: control, April 30.0 ± 1.1; lindane, May 2.7 ± 0.6; carbofuran, May 1.5 ± 0.9; pentachlorophenol, May 4.2 ± 1.8). The duration of pregnancy (when calculated as the number of days from the last mating until whelping) was extended (P < 0.05) in the group treated with lindane (control, 46.6 ± 1.0; lindane, 51.2 ± 1.5; carbofuran, 48.8 ± 1.5; pentachlorophenol, 52.8 ± 2.5). There were no significant differences in the proportion of male or female kits weaned or the growth rate of kits up to weaning at 8 weeks (kit weight at 2 weeks: control, 66.2 ± 9.1 g; lindane, 44.3 ± 10.4 g; carbofuran, 57.2 ± 4.6 g; pentachlorophenol, 64.5 ± 3.7 g; Kit weight at weaning: control, 504.0 ± 46.3 g; lindane, 482.0 ± 76.2 g; carbofuran, 588.9 ± 23.2 g; pentachlorophenol, 564.8 ± 33.5 g).

Endocrinology. The serum concentration of progesterone (nmol l⁻¹: control, 0.45 ± 0.06; lindane, 0.54 ± 0.16; carbofuran, 0.73 ± 0.13; pentachlorophenol, 0.45 ± 0.13) and cortisol at weaning (nmol l⁻¹: control, 27.1 ± 5.4; lindane, 56.8 ± 18.5; carbofuran, 57.0 ± 13.1; pentachlorophenol, 23.0 ± 10.2) were significantly increased (P < 0.05) in the mink treated with carbofuran. There was no effect of pesticide treatment on oestradiol concentration (pmol l⁻¹: control, 13.4 ± 1.7; lindane, 15.0 ± 2.4; carbofuran, 15.0 ± 3.6; pentachlorophenol, 17.6 ± 1.8) or thyroxine concentration at weaning (nmol l⁻¹: control, 32.6 ± 1.5; lindane, 33.4 ± 1.3; carbofuran, 33.9 ± 1.8; pentachlorophenol, 31.7 ± 1.1) (P > 0.05).

Histopathology. Few treatment specific lesions were seen and only two uterine lesions appeared to be related to treatment. All other noted lesions did not differ significantly in severity between treated and control mink. The presence of cystic uterine glands was observed in a number of animals in all groups; however, the severity of this lesion was greatest in the mink treated with pentachlorophenol (severity scores for control and pentachlorophenol were 0.19 ± 0.11 and 1.33 ± 0.47, respectively; P < 0.05). There was no incidence of mucosal inflammation in any mink treated with pesticide; however, a number of mink in the control group exhibited mild mucosal inflammation such that the severity of this lesion was significantly more pronounced in the control group (P < 0.05) than in each of the groups treated with pesticide (control, 0.58 ± 0.30; lindane, carbofuran, pentachlorophenol, 0.00 ± 0.00).

Experiment 2

No overt signs of toxicity were noted. lindane did not cause any significant change in body weight during the prepartum period. During the postpartum period, body weight for the control mink was decreased relative to the mink treated with lindane (body weight at weaning: control, 887 ± 31 g; lindane, 1003 ± 36 g), probably owing to the increased metabolic cost of rearing more and larger litters.

Fertility. Fertility data are summarized (Fig. 2). The treatment had no significant effect on the proportion of mink that accepted the initial mating (control, 100%; lindane, 87%) or the
date on which the first or last mating was accepted. Of the mink that mated at the first mating, the percentage of mink that accepted the second mating was also not affected by treatment (control, 87%; lindane, 92%). When all mink (mated and non-mated) were considered, treatment with lindane caused a significant decrease in the whelping rate (control, 73%; lindane, 47%; \( P < 0.05 \)). When only the mated mink were considered, treatment with lindane also tended to cause a decrease in whelping rate (control, 73%; lindane, 54%; \( P < 0.1 \)). However, the proportion of mated mink with implantation sites (control, 73%; lindane, 77%) and the number of implantation sites per mink (control, 5.07 ± 1.08; lindane, 3.69 ± 0.98) were not affected by treatment with lindane \(( P > 0.1 \)). The proportion of mink with implantation sites that whelped subsequently was reduced by treatment with lindane (control, 100%; lindane, 70%; \( P < 0.05 \)). The proportion of embryos lost after implantation in each litter (implantation scars not represented by kits at whelping) was significantly greater in the group treated with lindane compared with the control group (control, 30.5 ± 8.3%; lindane, 64.9 ± 10.0%; \( P < 0.01 \)). The increased incidence of embryo loss was mostly accounted for by the loss of complete litters in three of the mink treated with lindane. If the results from these three mink were removed from the data set, the rate of embryo loss was similar in both groups (control, 30.5 ± 8.3%; lindane, 40.6 ± 9.4%; \( P > 0.1 \)). The mean litter size of mink that whelped was not significantly reduced by lindane (control, 4.73 ± 0.98; lindane, 3.29 ± 0.64; \( P > 0.1 \)). Mink in the group treated with lindane tended to whelp later than control mink (control, May 4.6 ± 1.5; lindane, May 8.7 ± 2.7; \( P < 0.1 \)). However, duration of gestation was not affected by the lindane treatment (control, 52.2 ± 1.8; lindane, 54.9 ± 3.2 days; \( P > 0.1 \)). There were no significant differences in the proportion of male or female kits weaned or the body weight of kits at 2 weeks (control, 72 ± 4.8 g; lindane, 82 ± 4.2 g) and 10 weeks (control, 760 ± 91 g; lindane, 704 ± 41 g) of age.

**Endocrinology.** Serum concentration of cortisol (nmol l\(^{-1}\); control, 61.5 ± 9.8; lindane, 23.5 ± 3.8) was significantly lower \(( P < 0.01 \)) in the mink treated with lindane. Concentrations of oestradiol (pmol l\(^{-1}\); control, 11.8 ± 1.8; lindane, 7.71 ± 1.8) progesterone (nmol l\(^{-1}\); control, 0.45 ± 0.06; lindane, 0.57 ± 0.19) and thyroxine (nmol l\(^{-1}\); control, 27.5 ± 1.3; lindane, 31.6 ± 2.1) did not differ between groups \(( P > 0.05 \).)

**Histopathology.** Only one uterine lesion appeared to be related to treatment. All other noted lesions did not differ statistically in severity between treatment and control groups. The most significant finding was that haemosidrosis of the uterus appeared to be inhibited in the treated animals (severity score for control = 0.82 ± 0.27; lindane = 0.02 ± 0.02; \( P < 0.01 \)). The severity of haemosidrosis was most marked in the control group mink which carried large litters to term.

**Discussion**

Experiment 1 demonstrated that exposure to both lindane and pentachlorophenol had a detrimental effect on fertility; however, it was not clear whether the lower whelping rates were the result of a direct effect on embryo survival or an indirect effect of the lack of a second mating on embryo loss. Although neither the lower implantation rate nor the lower survival of embryos after implantation in the mink treated with lindane and pentachlorophenol were statistically significant compared with control mink, the combined effect of these decreases contributed to the reduced whelping rate. The decrease in the number of mink that accepted a second mating 1 week after the first mating appeared to be the result of an unwillingness to copulate on the part of the females when placed with a male, since when the males were exposed subsequently to an untreated female (not involved in the current experiment) copulation took place. The second mating is used routinely in normal ranch practice since it has been demonstrated to improve whelping rates (Lagerkvist, 1992). Therefore, a missed second mating would be expected to reduce whelping rates and it was clear from the results of the current study that mink that mated only once had a significantly lower whelping rate than mink that mated twice. Experiment 2 showed that when the mating response was equivalent in lindane and control mink, there was still a decreased whelping rate. Again, implantation rate was unaffected by treatment and, therefore, the majority of embryo loss was deemed to have occurred after implantation. Similar results have been observed in mink exposed to polychlorinated biphenyls (Backlin and Bergman, 1995).

This study is the first to show that lindane causes increased embryo loss in mink; however, the mechanism by which lindane has this effect is unclear. Previous studies in rabbits (Palmer et al., 1976a) and mice (Sircar and Lahiri, 1989) using short-term treatments with high doses of lindane at various stages of pregnancy showed a marked negative effect on fertility. However, when progesterone and oestradiol were given with lindane during early pregnancy, both implantation and fetal development were normal (progesterone alone had no effect; Sircar and Lahiri, 1989). Lindane decreased the rate of steroidogenesis in mice by inhibition of cholesterol side chain cleavage (Sircar and Lahiri, 1990) and has been shown to...
decrease progesterone concentrations (Srivastava and Raizada, 1993). Lindane has also been shown to be anti-oestrogenic, in that it blocks the response of oestrogen-dependent tissues to oestradiol (Chadwick et al., 1988; Cooper et al., 1989; Laws et al., 1994). However, in rats, lindane does not appear to exert its effect by binding to the oestrogen receptor (Uphouse and Williams, 1989). Lindane has also been shown to have a direct degenerative affect on mouse (Alm et al., 1994) and bovine (Alm et al., 1996) embryos in culture. Hassoun and Stohs (1995) indicated that certain fetotoxic effects of lindane may be due to superoxide production induced by oxidative stress. The decreased incidence of haemosiderosis in mink treated with lindane in the present study may have been a symptom, rather than a cause, of the decreased whelping rate in the second experiment. Alternatively, it may have been a direct effect of lindane, suggesting an inhibition of the monocyte–macrophage system or an alteration in endothelial integrity, which may play a role in the increased incidence of embryo loss. It is unlikely that the increased incidence of embryo loss observed merely represents a symptom of general pesticide toxicity, since no significant lesions related to the treatments were observed in the histological examination and no detrimental effects on feed intake or body weight were recorded.

In the present study, whelping rate in mink was also decreased by pentachlorophenol. In rats, pentachlorophenol has been shown to increase the incidence of fetal reabsorption and to cause an increase in fetal abnormalities when given during organogenesis and at doses at least tenfold higher than those given in the present study (Schwetz et al., 1974; Exon, 1984; Welsh et al., 1987). Exon (1984) suggested that the major mode of action of pentachlorophenol was its ability to uncouple oxidative phosphorylation. The increased severity of cystic uterine glands may have contributed to the increased incidence of embryo loss in the mink treated with pentachlorophenol. Cystic uterine glands may be associated with uterine infection and this could indicate an immunosuppressive activity of pentachlorophenol on the uterus. In the present study, carbofuran had no effect on reproductive function in mink, in agreement with work in rats (Barnett et al., 1980).

The doses of lindane and pentachlorophenol used in the present study were lower than those used in some previous studies in which no effects were observed on embryo loss in rats (Palmer et al., 1978b; Welsh et al., 1987). Mink may be particularly susceptible to lindane and pentachlorophenol owing to their extended period of embryonic diapause. During the period of embryonic diapause, circulating oestriadiol concentrations remain high and further ovulations can be induced (Lagerkvist et al., 1992). In mink, an increase in prolactin terminates embryonic diapause by initiating progesterone production (Murphy et al., 1981). The increased incidence of embryo loss in mink treated with lindane may be the result of inadequate concentrations of steroid hormone or altered responsiveness to ovarian steroid hormones. The progesterone concentration may be critical since there appears to be a decline in uterine progesterone receptors immediately after implantation in carnivores (Mead and Eroschenko, 1995). McRae (1994) concluded that inadequate progesterone secretion or excessive oestriadiol secretion may be important causes of preimplantation embryo loss in ranned mink. Unfortunately, we were unable to collect blood samples at the time of implantation since the stress of disturbance involved in this procedure was likely to have caused even further embryo loss (Tauson, 1988).

No effect of lindane on serum oestradiol or progesterone concentrations was apparent at the time of weaning in this study; however, steroid concentrations were very low at weaning. The later whelping date in mink treated with lindane and pentachlorophenol in this study mirrors the effect seen in rats where lindane is known to decrease uterine contractions by an inhibitory effect on the gap junctions essential for successful parturition (Kamrin et al., 1994). This effect appears to be mediated by an increase in arachidonic acid (Criswell and Loch-Caruso, 1994).

In the present study, lindane and pentachlorophenol also caused a reduction in the number of mink accepting a second mating 7 or 8 days after the initial mating. In untreated mink, after the initial mating, a new cohort of ovarian follicles develops over the next few days (Douglas et al., 1994). The follicles secrete oestriadiol, increasing sexual receptivity, and the female accepts a second mating 7–8 days later and ovulates a second cohort of follicles (Hansson, 1947). As a significant proportion of the mink treated with lindane and pentachlorophenol did not accept the second mating, we can speculate that follicle growth was disrupted after the initial mating had induced ovulation, leading to a decline in oestriadiol production. Higher doses of lindane have been shown to inhibit cell proliferation and DNA synthesis in an in vitro system involving cultured bovine granulosa cells (Tiemann et al., 1996), and relatively low doses of lindane can reduce ovulation rates in rabbits (Lindenaau et al., 1994). The initial mating was not affected by the treatment with lindane or pentachlorophenol, possibly because the follicles present at this mating had developed before the start of treatment. Lordosis and proceptive behaviour were also reduced in rats administered high doses of lindane during pro-oestrus (Uphouse, 1987) or dioestrus (Uphouse and Williams, 1989), probably as a result of direct neurobehavioural effects since such doses cause overt convulsions. The behaviours seen in mink treated with lindane that failed to mate at the second attempted breeding (rearing, vocalization, and aggression towards the male) were similar to those seen in rats treated with lindane and those seen when an anoestrous female mink is placed with an active male. There do not appear to be any previous studies showing an effect of pentachlorophenol on receptivity and oestrous behaviour in mammals. Despite the fact that the decreased second mating response in the group treated with lindane did not totally account for the decreased whelping rate, it is still of interest because a decrease in libido in a wild population could have serious effects at the population level. Mink are likely to be exposed to much lower doses of lindane and pentachlorophenol in their natural habitat; however wild populations may be exposed to a mixture of pesticides with possible additive and synergistic effects. Tissue concentrations of pesticides in mink fed the same doses of lindane and pentachlorophenol during a longer term study (A. Beard, unpublished) and in calves given similar treatment (Hughes et al., 1985) were 1–5 mg kg⁻¹. These concentrations are generally 10–1000 times higher than those measured in wild animals (Franson et al., 1974; Krynski et al., 1982; Falandysz and Kannan, 1992).
Serum cortisol concentration was increased in mink treated with carbofuran in Expt 1 and reduced in mink treated with lindane in Expt 2; however, single time point measurements of cortisol are relatively unreliable compared with frequent sampling as cortisol secretion is episodic (Goodman, 1994) and may vary in response to stress. Studies have detected increased plasma concentrations of corticosterone in rats exposed in utero and neonatally to low doses of carbofuran, but not in rats given high doses (Crammer et al., 1978). Lindane treatment in rats has been shown to affect adrenal function adversely, leading to a reduction in plasma glucocorticoid concentration (Lahiri and Sircar, 1991); however, in rabbits, lindane treatment causes an increase in cortisol secretion (Anand et al., 1990).

In conclusion, this study showed that exposure of mink to the organochlorine pesticide lindane caused an increase in the number of embryos lost after implantation. The mechanism of this effect is unclear but may involve a disruption in oestrogen-dependent changes normally associated with pregnancy as is seen in mice treated with lindane (Sircar and Lahiri, 1989). Lindane and pentachlorophenol also decreased the number of females accepting a second mating when this was attempted 1 week after the initial mating. Pentachlorophenol also caused a decrease in whelping rate, but it is unclear whether this is a direct effect on embryo loss or a result of the reduction in mating response. Carbofuran did not appear to have any effect on reproduction. Pregnancy in mink appears to be particularly sensitive to the ingestion of low concentrations of pesticides and this may be due to particular characteristics of gestation in mink, such as the extended period of embryonic diapause. Environmental contamination with organochlorine pesticides may pose a risk to wild populations of mink owing to disruptive effects of these substances on fertility.

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