

Data Evaluation Record
BRODIFACOU
Wild Mammal Toxicity Test

GUIDELINE NUMBER: 71-3

CITATION: Ringer, R.K. and Richard J. Aulerich. Date no given, but, c. 1979. Determination of oral LD₅₀ of Brodifacoum for mink. Submitted by ICI Americas, Inc., Agricultural Products, Wilmington, Delaware 19897. RR 90-292 B.

REASON FOR SUBMISSION:

FIFRA '88 Reregistration.

RESULTS- Valid _____

Invalid _____

GUIDELINE- Satisfied _____

Partially Satisfied _____

Supplemental X

Not Satisfied X

DISCUSSION:

No DER was found in EEB's files for this study. The two highest concentrations are too widely separated to be of use. Each successively larger concentration should be 1.66 times that of the one before (or be about 0.6 of the one larger than itself). The sixth group is so much larger (4.7 times) than the fifth that the gap between the two groups made the LD₅₀ meaningless. If it had been 3.73 mg/kg (1.66 X 2.24 mg/kg) and the animals had still died, the LD₅₀ would drop to 3.73 mg/kg even though the mortality remained the same. We cannot tell the difference between the two possibilities from this study. The "Protocol for determination of oral LD₅₀ for Mink" specifies .04, .12, .36, 1.08, and 3.24 mg/kg. If it was believed that a higher range of concentrations was needed, then the entire range should have been increased, not just its highest member.

Some animals had bloody stools even at the 0.116 mg/kg level, therefore, the NOEL = 0.04 mg/kg.

If, as stated, the species presented a difficulty because food passes through their digestive tract to quickly, another subject should be chosen. Although, this may be a trait of the Mustelids generally and, therefore, the high LD₅₀ accurately reflects the conditions in the wild.

CONCLUSIONS:

The study is classified as "Supplementary" No LD₅₀ but a NOEL 0.04 mg/kg.

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Pages 2 through 3 are not included in this copy.

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