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105001 Shaughnessey No.

## EEB REVIEW

| DATE: IN August 3, 1988 OUT December 15, 1988                       |
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| FILE OR REG. NO. 241-GRU  |
| PETITION OR EXP. NO.  |
| DATE OF SUBMISSION June 10, 1988                                    |
| DATE RECEIVED BY EFED August 2, 1988                                |
| RD REQUESTED COMPLETION DATA <u>August 29, 1988</u>                 |
| EEB ESTIMATED COMPLETION DATE August 29, 1988                       |
| RD ACTION CODE/TYPE OF REVIEW 165                                   |
| TYPE PRODUCTS(S): I, D, H, F, N, R, S <u>Insecticide/Nematicide</u> |
| DATA ACCESSION NO(S). 406607-07                                     |
| PRODUCT MANAGER NO. J. Tice / M. Mautz                              |
| PRODUCT NAME(S) Counter XL  |
| COMPANY NAME American Cyanamid Company                              |
| SUBMISSION PURPOSE Proposed new formulation (20G) for use           |
| on corn, sorghum and sugar beets. Response to previous EEB          |
| review  |
| SHAUGHNESSEY NO. CHEMICAL AND FORMULATION % A.I.                    |
|   |

### DATA EVALUATION RECORD

1. TEST MATERIAL:

Terbufos

(S-[[(1,1-dimethylthyl)thio]methyl]0,0-diethyl phosphorodithioate)

2. STUDY MATERIAL - Counter 15G

Terbufos 15 W/W % Inert ingredients 85 100%

3. STUDY TYPE- Avian Dietary Single-dose Oral LD50. Species tested-

Bobwhite quail <u>Colinus virginianus</u>

4. STUDY IDENTIFICATION:

Fletcher, D.W. 1987. 21-day acute oral toxicity study with Counter 15G in Bobwhite quail. Bio-Life Associates, Ltd. Submitted by American Cyanamid Company, Princeton, NJ MRID 406607-07.

5. REVIEW BY:

James J. Goodyear Signature: Biologist Ecological Effects Branch Environmental Fate and Effects Division (TS-796C)

6. APPROVED BY:

Raymond W. Matheny Head, Section 1 Ecological Effects Branch Environmental Fate and Effects Division (TS-796C)

Signature:

SEP 3 0 1988 Date:

7. CONCLUSIONS:

The study meets the requirements of the Avian Single-Dose Oral LD50 guidelines. Because the Terbufos formulated product has an  $LD_{50}$  of 305.0 mg/kg it is considered to be moderately toxic to bobwhite quail. It does not, however, meets the requirements for the registration of Counter 20P since it was done with Counter 15G.

- 8. RECOMMENDATIONS- N/A.
- 9. BACKGROUND:

The study was submitted for the registration of Counter 20P.

- 10. DISCUSSION OF INDIVIDUAL TEST- N/A.
- 11. MATERIALS AND METHODS:
  - A. Test animals:

Bobwhite quail (21 weeks old) from Oak Ridge Game Farm, Route 2, Gravette, Arkansas 72736.

B. Dose:

Counter 15G was administered in five dosages (147, 215, 316, 464 and 681 mg/kg).

C. Design:

There were five test groups of five male and five females (one for each for each test level) plus a control group of five males and five females housed in a 53.3 cm x 45.7 cm x 38.1 cm pens in a heated room in which lighting "was provided by fluorescent lights which were left on eight hours per day". The birds were observed and acclimated for 15 days and fasted for 22 hours before the dosing with gelatin capsules.

D. Statistics:

Litchfield, J.T., Jr. and F. Wilcoxon. 1949. A simplified method of evaluating dose-effect experiments. J. Pharmacology and Experimental Therapeutics. Vol. 96.

12. REPORTED RESULTS:

 $LD_{50} = 305.0 \text{ mg/kg}$  95% C.I. = 258.5 - 359.9 mg/kg The NOEL was not given.

13. STUDY AUTHORS' CONCLUSIONS/QA MEASURES:

"The acute oral median  ${\rm LD}_{50}$  of Counter 15G in Bobwhite quail was determined to be 305.0 mg/kg of body weight with 95% confidence limits of 258.5 to 359.9 mg/kg of body weight", and that,

"In accordance with Bio-Life Associates, Ltd. Laboratories' intent that all toxicity tests conducted by our facility follow good laboratory practices, Bio-Life Associates, Ltd's study director for the above test herein confirms that the study was conducted in compliance with the US EPA Good Laboratory Practice Regulations; Pesticide Programs (40 CFR 160)."

# 14. REVIEWER'S DISCUSSION AND INTERPRETATION OF THE STUDY:

#### A. Test Procedures:

The procedures were not in complete accordance with the guidelines for testing avian single-dose oral LD50.

Errors include; the chemical names of the pesticide were not given, it was not specifically stated if the test was done with the end-use product or the technical grade chemical, the per cent of active ingredient in the test substance was not given and it was not stated if the  ${\rm LD}_{50}$  is in milligrams of ai or end-use product. However, Marilyn Mautz of the Registration Division contacted the Cyanimid Company and found that all dosages and the  ${\rm LD}_{50}$  are in milligrams of Terbufos end-use product.

Only eight hours of light (rather than ten) were provided. The report only states that lighting was provided for eight hours per day, not that an 8 hour light / 16 hour dark photoperiod was provided. This leaves open the possibility that the room had natural light and, therefore, the ducks had a natural photoperiod. This might be of major importance in an avian dietary study but it is considered to be of minor importance in this acute toxicity study.

### B. Statistical Analysis:

The  $LD_{50}$  was calculated from the registrant's data using a computer program from "Stephan, et al. 1978. Computer program for calculating  $LD_{50}$ ; probit method". The  $LD_{50}$  was found to be 294.6 (215 - 464) mg (EUP)/kg.

### C. Discussion/Results:

Counter 15G formulated product can be characterized as being moderately toxic orally to bobwhite quail.

D. Adequacy of the Study:
Classification- Core.

Rational - The study is scientifically sound and meets the requirements of the guidelines. However, there are no current outstanding requirements for the 15% granular formulation. This study may be used to meet requirements of a Second Round Review, should one be initiated.

Repair- N/A.

- 15. COMPLETION OF ONE-LINER FOR STUDY: Yes, see the attached sheets.
- 16. CBI APPENDIX- N/A.