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OFFICE OF
PREVENTION, PESTICIOES
AND TOXIC SUBSTANCES

MEMORANDUM

FROM:

TO:

SUBBJECT; Ouinclorac - Submission of an Unscheduled DNA Synthesis Study in Support of Registration

TOX Chem No: 325A
PC No: 128974
DP Barcode Nos: D180499
D180498
D180500

Submission Nos: S421488 S421482

S421482 S421489

William B. Greear, M.P.H. William B File 21/92

Review Section IV, Toxicology Branch I

Health Effects Division (H7509C)

Vickie Walters/Robert Taylor, PM Team # 25

Herbicide-Fungicide Branch Registration Division (H7505C)

THRU: Marion P. Copley, D.V.M., Section Head

Health Effects Division (H7509C)

I. CONCLUSIONS:

The Unscheduled DNA Synthesis Study (UDS) No: 91/10965, dated October 24, 1991 was negative for mutagenic effects.

II. REQUESTED ACTION:

RD has requested that TB-I evaluate a study titled "In Vivo/In Vitro Unscheduled DNA Synthesis in Rat Hepatocytes With Reg. No. 150 732 / BAS 514 H" which was submitted in support of registration by Bob Rohde of the BASF Corporration under a cover letter dated June 25, 1992.

III. DISCUSSION:

An acceptable UDS study #86/0135 dated June, 1986 has already been submitted. It was negative. This new study #91/10965 has been examined and it also appears to be negative. An abbreviated DER is attached. The study is "Acceptable".



moople FAWG Reviewed By: William B. Greear, M.P.H. Review Section IV, Toxicology Branch I (H7509C) Secondary Reviewer: Marion P. Copley, D.V.M. Musseller Review Section IV, Toxicology Branch I (H7509C)

DATA EVALUATION REPORT

Guideline Series 83-3 STUDY TYPE:

Mutagenicity- Unscheduled DNA Synthesis

EPA IDENTIFICATION NUMBERS:

325A Tox Chem No: 128974 PC No: 423707-01 MRID No:

TEST MATERIAL: Reg. No. 150 732 / BAS 514H

SYNONYMS: Quinclorac, FACET

BASF Corporation SPONSOR:

Research Triangle Park, NC 27709-3528

STUDY NUMBER: 91/10965

Cyto Test Research GMBH & Co. TESTING FACILITY:

D-6101 Robdorf, FRG

In Vivo/In Vitro Unscheduled DNA Synthesis in TITLE OF REPORT:

Rat Hepatocytes With Reg. No. 150 732 / BAS

514 H

AUTHOR: Rolf Fautz

REPORT ISSUED: October 24, 1991

Negative for induction of DNA-damage leading to CONCLUSIONS: repair synyhesis at 1000 mg/kg for 4 hrs and at 100 and 1000 mg/kg for 16 hrs.

Classification: Acceptable

SUMMARY: The following summary was abstracted directly from the report #91/10965 page 0010.

"The test article Reg. No. 150 732 / BAS 514 ... H was assessed in the in vivo/ in vitro UDS assay for its potential to induce DNA repair (UDS) in the hepatocytes of rats.

The test article was formulated in 0.5 % CMC. This suspending agent was used as negative control. The volume administered orally was 10 ml/kg body weight (b.w.). After a treatment period of 4 and 16 hours, respectively, the animals were narcotized and sacrificed by liver perfusion. Primary hepatocyte cultures were established and exposed for 4 hours to 3HTdR which is incorporated if UDS occurs (3).

The test article was tested at the following dose levels:

4 hour treatment period: 1000 mg/kg b.w. 16 hour treatment period: 100 and 1000 mg/kg b.w.

For each dose level, including the controls, hepatocytes from three treated animals were assessed for the occurrence of UDS.

No toxic reactions of the animals occured at any of the treatment periods or dose groups. In addition, neither the viability nor the in vitro attachment of the hepatocytes was dramatically affected due to the in vivo pre-treatment with the test article.

No dose level of the test article revealed UDS induction in the hepatocytes of the treated animals as compared to the current negative controls.

An appropriate reference mutagen (2-AAF, 100 mg/kg b.w.) was used as positive control. Treatment with 2-AAF revealed distinct increases in the number of nuclear and net grain counts.

CONCLUSION

In conclusion, it can be stated that during the study described and under the experimental conditions reported, the test article did not induce DNA-damage leading to repair synthesis in the hepatocytes of the treated rats.

Therefore, Reg. No. 150 732 / BAS 514 .. H is considered to be non-effective in this in vivo/in vitro UDS test system."

Comment: TB-I concurs with the authors conclusions.