

US EPA ARCHIVE DOCUMENT



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
WASHINGTON, D.C. 20460

SEP 22 1992

OFFICE OF
PESTICIDES AND TOXIC
SUBSTANCES

MEMORANDUM

SUBJECT: MAT 7484 (Phostebupirim) - Rabbit Developmental
Toxicity Study - 6(a)(2) Submission ("Adverse Effects")

DP Barcode: D182216 **Chemical No.:** 129086
Submission No.: S424808 **Identification No.:** 003125-URR
Case: 030678 **MRID No.:** 424578-01

FROM: Alan C. Levy, Ph.D., Toxicologist *Alan C. Levy*
Review Section IV, Toxicology Branch II *9/16/92*
Health Effects Division (H7509C)

TO: Robert Forrest/Marilyn Mautz PM 14
Registration Division (H7505C)

THRU: Elizabeth A. Doyle, Ph.D., Section Head *E. A. Doyle*
Review Section IV, Toxicology Branch II *9/16/92*
Health Effects Division (H7509C)

and

Marcia van Gemert, Ph.D., Branch Chief
Toxicology Branch II *M van Gemert 9/16/92*
Health Effects Division (H7509C)

Registrant: Miles Agricultural Division, Kansas City, MO

REQUEST: Comment on the Registrant's submission of 6(a)(2)
(adverse effects) data regarding a rabbit developmental
toxicity study with MAT 7484 (Phostebupirim).



COMMENTS:

An addendum to a rabbit developmental toxicity study with MAT 7484 (MRID No. 420054-55), was submitted because of "adverse effects". This is a NEW CHEMICAL and a review of the original submission of the studies has not been completed as of the date of this Memorandum. A cover letter from John S. Thornton (Miles) to Document Processing Desk [6(a)(2)] of EPA with a copy to Robert A. Forrest (August 26, 1992), stated, in part, "... we would like to bring these results to your attention in conformance with the reporting requirements of FIFRA 6(a)(2)." "Unfortunately, this addendum is in German and will not be translated in time to meet the 6(a)(2) reporting deadlines."

Toxicology wishes to indicate the following:

1. The submitted report does not constitute 6(a)(2) data because the chemical is not currently registered in the United States. Therefore, the product can not present a hazard in any way.
2. The Registrant is to submit to the Agency only those documents which are in English or for which a translation is provided.
3. Because of the above two statements, Toxicology will not review the submitted addendum at this time.

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