LEVEL II

UNITED STATES ARMY
ENVIRONMENTAL HYGIENE
AGENCY
ABERDEEN PROVING GROUND, MD 21010

TOPICAL HAZARD EVALUATION PROGRAM
OF CANDIDATE INSECT REPELLENT AI3-36467
US DEPARTMENT OF AGRICULTURE PROPRIETARY COMPOUND
STUDY NO. 75-51-0899-79
MAY 1976 - JUNE 1978

Approved for public release; distribution unlimited.
A hazard evaluation of candidate insect repellent A13-36467 was performed by means of laboratory studies using rats, rabbits and guinea pigs. The technical grade compound caused mild skin irritation, but no eye irritation, no phototoxic irritation in rabbits, no sensitization reactions in guinea pigs and did not demonstrate an acute ingestion hazard. It is recommended that A13-36467, US Department of Agriculture Proprietary Compound, be approved for further testing as a candidate insect repellent.

A summary of the pertinent findings and recommendations of the inclosed report follows:

A hazard evaluation of candidate insect repellent A13-36467 was performed by means of laboratory studies using rats, rabbits, and guinea pigs. The technical grade compound caused mild skin irritation, but no eye irritation, no phototoxic irritation in rabbits, no sensitization reactions in guinea pigs and did not demonstrate an acute ingestion hazard. It is recommended that A13-36467, US Department of Agriculture Proprietary Compound, be approved for further testing as a candidate insect repellent.

FOR THE COMMANDER:

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1. AUTHORITY.


b. Memorandum of Understanding between the Department of the Army; Office of The Surgeon General; the US Army Health Services Command; the US Army Environmental Hygiene Agency; the Armed Forces Pest Control Board; and the US Department of Agriculture, effective 1970 with Amendment No. I effective August 1974.


3. PURPOSE. The purpose of this program is to provide guidance for further entomological testing of the candidate insect repellent AI3-36467.

4. SUMMARY OF FINDINGS. A hazard evaluation of the candidate insect repellent AI3-36467, US Department of Agriculture Proprietary Compound, was conducted by this Agency using New Zealand White rabbits for skin and eye studies, Hartley guinea pigs for a skin sensitization study and Sprague-Dawley rats for determination of oral toxicity. A tabular presentation of animal toxicity data developed in this Agency follows.*


† The experiments reported herein were performed in animal facilities fully accredited by the American Association for the Accreditation of Laboratory Animal Care.

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Study No. 75-51-0899-79, May 76-Jun 78

TABULAR PRESENTATION OF DATA

<table>
<thead>
<tr>
<th>Test</th>
<th>Results</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td>SKIN IRRITATION STUDIES</td>
<td></td>
<td></td>
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<tr>
<td>Rabbits</td>
<td></td>
<td></td>
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<tr>
<td>Single 24-hour application</td>
<td>Compound A13-36467 produced mild primary irritation of the intact skin and the skin surrounding an abrasion.</td>
<td>USAEHA Category II (ref Appendix)</td>
</tr>
<tr>
<td>to intact and abraded skin</td>
<td></td>
<td></td>
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<tr>
<td>of New Zealand White rabbits.</td>
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<tr>
<td>0.5 ml technical grade</td>
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<tr>
<td>compound applied to each</td>
<td></td>
<td></td>
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<tr>
<td>of six rabbits.</td>
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<td></td>
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<tr>
<td>EYE IRRITATION STUDIES</td>
<td></td>
<td></td>
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<tr>
<td>Rabbits</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Single 24-hour application</td>
<td>Compound A13-36467 did not produce any injury to the cornea and only very slight irritation to the conjunctiva in six out of six rabbits.</td>
<td>USAEHA Category A (ref Appendix)</td>
</tr>
<tr>
<td>of 0.1 ml technical grade</td>
<td></td>
<td></td>
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<tr>
<td>compound to one eye of each</td>
<td></td>
<td></td>
</tr>
<tr>
<td>of six New Zealand White</td>
<td></td>
<td></td>
</tr>
<tr>
<td>rabbits.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>APPROXIMATE LETHAL DOSE (ALD)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Oral</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rats (male) - no diluent</td>
<td>ALD = 4900 mg/kg</td>
<td>Presents little lethal hazard from acute accidental ingestion.</td>
</tr>
</tbody>
</table>
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### PHOTOCHEMICAL SKIN IRRITATION STUDIES

#### Rabbits

A single application (0.05 ml) of a 25 percent (w/v) solution of the compound (AI3-36467) and of a 10 percent oil of Bergamot solution (positive control) in 95 percent ethyl alcohol, were applied to the intact skin of six rabbits. Five minutes after application, the rabbits were exposed to UV light (365 nm) for 30 minutes at a distance of 10-15 cm.

**Control**

Following UV exposures of the rabbits, 0.05 ml of test compound, positive control and diluent were applied to additional skin areas to serve as unirradiated control sites. Application areas were checked for skin irritation at 24, 48 and 72 hours.

<table>
<thead>
<tr>
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</thead>
<tbody>
<tr>
<td>A 25 percent solution of AI3-36467 in ethanol did not cause a photochemical skin irritation reaction under test conditions.</td>
<td>Compound AI3-36467 did not cause a photochemical irritation reaction under test conditions and is not expected to cause a photochemical irritation reaction in humans.</td>
<td></td>
</tr>
<tr>
<td>Positive control application and irradiation caused greater irritant effects than in unirradiated skin areas.</td>
<td>Ethanol solutions caused a very slight erythema reaction on both non-UV and UV skin sites.</td>
<td>Ethanol solutions of this compound may cause a slight skin reaction in sensitive individuals.</td>
</tr>
</tbody>
</table>
**SENSITIZATION STUDIES**

**Guinea Pigs (male)**

Intradermal injections of 0.1 ml of a 0.1 percent suspension (w/v) of AI3-36467 or of dinitrochlorobenzene (DNCB)* in a mixture containing 1 volume propylene glycol and 29 volumes of saline.

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</thead>
<tbody>
<tr>
<td>Ten test guinea pigs received and challenged with a 0.1 percent solution of AI3-36467.</td>
<td>Challenge dose of test compound (last intradermal injection) did not produce a sensitization reaction.</td>
<td>Compound AI3-36467 did not produce a sensitization reaction under these test conditions and is not expected to cause a sensitization reaction in humans.</td>
</tr>
<tr>
<td>Ten positive control guinea pigs received and challenged with a 0.1 percent suspension of DNCB.</td>
<td>Positive control (DNCB) produced a marked sensitization reaction in 10 out of 10 guinea pigs.</td>
<td></td>
</tr>
</tbody>
</table>

* A known skin sensitizer
5. CONCLUSION. The candidate insect repellent AI3-36467 has a potential for causing some slight skin irritation, but presents no acute hazard from eye, photochemical, or sensitization contact or from acute ingestion. Slight irritation may result from skin exposure to ethanol solutions of AI3-36467.

6. RECOMMENDATION. Under the provisions of the Memorandum of Understanding (paragraph 1b), it is recommended that AI3-36467, USDA Proprietary Compound, be approved for further testing as a candidate insect repellent. The compound should be used with caution around abrasions of the skin. Persons experiencing irritation when working with ethanol solutions of AI3-36467 should wash the site with copious amounts of water.

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APPENDIX

TOPICAL HAZARD EVALUATION PROGRAM
DEFINITIONS OF CATEGORIES OF COMPOUNDS BEING CONSIDERED FOR ACUTE SKIN APPLICATION

CATEGORY I - Compounds producing no primary irritation of the intact skin or no greater than mild primary irritation of the skin surrounding an abrasion. (INTERPRETATION: No restriction for acute application to the human skin.)

CATEGORY II - Compounds producing mild primary irritation of the intact skin and the skin surrounding an abrasion. (INTERPRETATION: Should be used only on human skin found by examination to have no abrasions or may be used as a clothing impregnant.)

CATEGORY III - Compounds producing moderate primary irritation of the intact skin and the skin surrounding an abrasion. (INTERPRETATION: Should not be used directly on the skin without a prophetic patch test having been conducted on humans to determine irritation potential to human skin. May be used without patch testing, with extreme caution, as clothing impregnants. Compound should be resubmitted in the form and at the intended use concentration so that its irritation potential can be reexamined using other test techniques on animals.)

CATEGORY IV - Compounds producing moderate to severe primary irritation of the intact skin and of the skin surrounding an abrasion and, in addition, producing necrosis, vesiculation and/or eschars. (INTERPRETATION: Should be resubmitted for testing in the form and at the intended use concentration. Upon resubmission, its irritation potential will be reexamined using other test techniques on animals, prior to possible prophetic patch testing in humans, at concentrations which have been shown not to produce primary irritation in animals.)

CATEGORY V - Compounds impossible to classify because of staining of the skin or other masking effects owing to physical properties of the compound. (INTERPRETATION: Not suitable for use on humans.)

EYE CATEGORIES:

A. Compounds noninjurious to the eye. INTERPRETATION: Irritation of human eyes is not expected if the compound should accidentally get into the eyes, provided it is washed out as soon as possible.

B. Compounds producing mild injury to the cornea. INTERPRETATION: Should be used with caution around the eyes.

C. Compounds producing mild injury to the cornea, and in addition some injury to the conjunctiva. INTERPRETATION: Should be used with caution around the eyes and mucosa.
D. Compounds producing moderate injury to the cornea. INTERPRETATION: Should be used with extreme caution around the eyes.

E. Compounds producing moderate injury to the cornea, and in addition producing some injury to the conjunctiva. INTERPRETATION: Should be used with extreme caution around the eyes and mucosa.

F. Compounds producing severe injury to the cornea and to the conjunctiva. INTERPRETATION: Should be used with extreme caution. It is recommended that use be restricted to areas other than the face.