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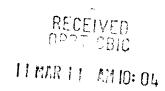
8EHQ-07-16913 89110000189 3/11/11

COMMENTS: COMMUN S (DECLASS)

DOES NOT CONTAIN CBI

MR# 334148





Andrea V. Malinowski

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March 10, 2011

VIA FEDERAL EXPRESS

8EHQ-0311-16913H DCN: 89110000189

Attn: TSCA Declassification Coordinator U.S. Environmental Protection Agency Office of Pollution Prevention and Toxics Document Control Office (7407M) Washington, D.C. 20460



Re:

Declassification Activity - TSCA §8(e) Submission

Originally Assigned 8EHQ Number: 8EHQ-07-16913 (letter dated 07.26.07)

Originally Assigned Bar Code: 88070000331

CAS number: 647-42-7

Supplemental Submission - Revised Public Copy of Submission

Dear TSCA Declassification Coordinator:

This submission is made in connection with the EPA 2010 CBI Declassification Challenge initiative.

Please find enclosed a revised public copy of the above-identified submission. Any information still claimed as confidential business information (CBI) in the attached report has been redacted and replaced by brackets. The originally assigned 8EHQ number has been added by the submitter to the first page of the enclosed revised public copy of the submission.

Very truly yours,

Andrea V. Malinowski

Killialla

Enclosure





Revised Public Copy - Submitted 03.10.11 Originally Assigned 8EHQ Number: 8EHQ-07-16913 Originally Assigned Bar Code: 88070000331

DuPont Haskell Laboratory for Health and Environmental Sciences Elkton Road, P.O. Box 50 Newark, DE 19714-0050

July 26, 2007

Via Federal Express

Confidential Business Information

Document Processing Center (Mail Code 7407M)
Room 6428
Attention: 8(e) Coordinator
Office of Pollution Prevention and Toxics
U.S. Environmental Protection Agency, ICC Building
1201 Constitution Ave., NW
Washington, DC 20460

Dear 8(e) Coordinator:

1-Octanol, 3,3,4,4,5,5,6,6,7,7,8,8,8-tridecafluoro-CAS# 647-42-7 Generic Name: Fluorinated Aliphatic Alcohol

This letter is to inform you of the results of a recently conducted acute oral toxicity study (up-and-down method) in rats with the above referenced substance.

The test substance was administered neat by oral gavage to 1 fasted female rat at a dose of 175 mg/kg, to 3 fasted female rats at a dose of 550 mg/kg, to 4 fasted female rats at a dose of 1750 mg/kg, and to 2 fasted female rats at a dose of 5000 mg/kg. Death occurred in 2 rats dosed at 1750 mg/kg (1 found dead, 1 sacrificed *in extremis*) and in both rats dosed at 5000 mg/kg (both found dead). The oral LD₅₀ is 1750 mg/kg.

One surviving rat dosed at 1750 mg/kg exhibited lethargy on the day of dosing and high posture up to 6 days after dosing. The other surviving rat dosed at 1750 mg/kg exhibited ataxia, lethargy, and high posture on the day of dosing. Another rat dosed at 1750 mg/kg exhibited ataxia on the day of dosing, hyperreactivity 2 days after dosing, high posture up to 2 days after dosing, and excessive vocalization 2 days after dosing. This rat was moribund and was sacrificed *in extremis* 3 days after dosing. The remaining rat dosed at 1750 mg/kg exhibited high posture the day of and day after dosing, lethargy 1 and 2 days after dosing, and leaning as well as ataxia 2 days after dosing. This rat was found dead 2 days after dosing. One rat dosed at 5000 mg/kg exhibited ataxia, lethargy, and high posture on the day of dosing and was found dead the day after dosing. The remaining rat

dosed at 5000 mg/kg was found dead on the day of dosing. No effects were observed in animals treated with 550 mg/kg or lower doses.

Substantiation of our claim of confidentiality is enclosed.

Sincerely,

A. Michael Kaplan, Ph.D.

Director - Regulatory Affairs and Occupational Health

AMK/CFC: clp (302) 366-5260

From: (302) 773-0071 Doris Duffy E. I. du Pont de Nemours & Co. 1007 Market Street D-7096-1 Wilmington, DE 19898

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