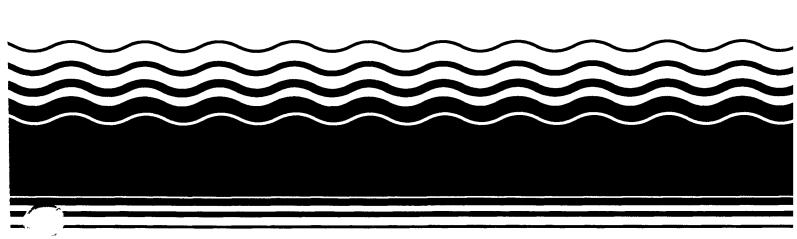
United States Environmental Protection Agency Office of Solid Waste and Emergency Response Publication 9200.6-303(94-1) EPA540/R-94/020 PB94-921199 March 1994

Superfund



# Health Effects Assessment Summary Tables

FY-1994 Annual



9200.6-303(94-1) EPA 540-R-94-020 PB94-921199 March 1994

# HEALTH EFFECTS ASSESSMENT SUMMARY TABLES

FY-1994 Annual

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#### **DISCLAIMER**

This report has been prepared by the U.S. Environmental Protection Agency. The information contained herein has been taken from final documents prepared by the Office of Health and Environmental Assessment for the Office of Solid Waste and Emergency Response and the Office of Water, Washington, DC and the Office of Air Quality Planning and Standards, Research Triangle Park, NC. These documents were reviewed in accordance with Agency policy and approved for publication. Mention of trade names or commercial products does not constitute endorsement or recommendation for use.

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#### INTRODUCTION

This document is the FY94 Annual Update of the Health Effects Assessment Summary Tables (HEAST) prepared by EPA's Office of Health and Environmental Assessment, Environmental Criteria and Assessment Office, Cincinnati, OH for use at both Superfund and RCRA sites. It completely replaces all former editions of the HEAST.

This version of the HEAST will be updated in 1994 by a July 1994 Supplement No. 1 and a November 1994 Supplement No. 2. These supplements will supercede the information in this document, the March 1994 HEAST Annual Update. Therefore, if the supplement(s) are available they should be checked whenever this document is consulted. These supplements, however, will not be produced to stand alone and will not contain the User's Guides or Appendix that are available in the annual update. Thus, the user is strongly encouraged to refer to the March 1994 HEAST Annual Update for this information.

The HEAST is a comprehensive listing consisting almost entirely of PROVISIONAL RISK ASSESSMENT INFORMATION relative to oral and inhalation routes for chemicals of interest to Superfund, the Resource Conservation and Recovery Act (RCRA), and the EPA in general. These entries in the HEAST are limited to chemicals that have undergone review and have the concurrence of individual Agency Program Offices, and each is supported by an Agency reference. This risk assessment information has not, however, had enough review to be recognized as high quality, Agency-wide consensus information.

The Integrated Risk Information System (IRIS) is the Agency's official repository of Agency-wide consensus chronic human health risk information. IRIS evaluations are conducted by the Agency's Work Group Review process, i.e., they have been examined by either the Reference Dose/Reference Concentration (RfD/RfC) Work Group or the Carcinogen Risk Assessment Verification Endeavor (CRAVE) Work Group. These Agency Work Groups conduct a process that leads to internal Agency scientific consensus regarding risk assessment information on a chemical. This information is recorded on IRIS, is considered to be "Work Group Verified", and does not appear on the HEAST. Thus, provisional risk assessment information on the HEAST is subject to possible review and revision by these Agency Work Groups.

There are two exceptions to the above discussion. The HEAST also contains information on chemicals that are a part of the National Ambient Air Quality Standards (NAAQS) or the Drinking Water Criteria Document (DWCD) series. In each of these cases, the chemicals are subject to extensive scientific peer review processes of extremely high quality.

#### **CHEMICAL STATUS DEFINITIONS**

Chemicals reviewed by the Agency Work Groups are classified according to their status as either "verified", "not verifiable", or "under review". The toxicity values (other than NAAQS or DWCD values) listed on the HEAST are considered to be "provisional". The Agency has no official definitions for these terms, but the HEAST user may interpret them as follows:

**Provisional:** A toxicity value or a cancer value is "provisional" if the value has had some form of Agency review, but it does not appear on the IRIS system. These values are generated in several ways. Often they are determined in the course of developing an Agency document on a chemical or on a class of chemicals. Some have been generated through the Work Group process, but have not yet been input to the IRIS system. At the time each value was derived, all available information on the chemical was evaluated, the value was calculated using the most current methodology, and a consensus was reached on the value by Agency scientists.

Brackets are placed around the names of toxicity and carcinogenicity values on the HEAST to distinguish these "provisional" values from information on IRIS. The following names are affected: RfD to [RfD], RfC to [RfC], slope factor to [slope factor], EPA group to [EPA Group] and unit risk to [unit risk].

# These "provisional" values are found on the HEAST. They do not appear on IRIS.

**Verified:** A toxicity value or a cancer value is "Work Group Verified" if all available information on the value has been examined by an Agency Work Group, the value has been calculated using current Work Group methodology, a unanimous consensus has been reached on the value by the Work Group, and the value appears on IRIS.

Some numbers that have achieved unanimous concensus by the Work Group may appear on the HEAST for a short time until they are loaded onto IRIS, at which time they are termed, "verified." During the interim, they are considered to be "provisional" values that are still "under review" by the Work Group.

# These "verified" numbers only appear on IRIS. They do not appear on the HEAST.

**Not verifiable:** A toxicity value is "not verifiable" if an Agency Work Group has considered all available data on a chemical and has unanimously determined that data are inadequate to generate a value that would be suitable for inclusion on IRIS. No toxicity value is calculated; no toxicity value is available for IRIS or the HEAST.

This "not verifiable" status is noted on IRIS, and is sometimes found on the HEAST, with a pointer to the IRIS system.

**Under Review:** A toxicity value is "under review" if an Agency Work Group is in the process of considering all available data on a chemical. All Work Group chemicals will have this status until the toxicity value is placed on the IRIS system. Toxicity values that have been withdrawn from IRIS by a Work Group for further review will have this status.

This "under review" status may be indicated on IRIS or on the HEAST. During this time, "provisional" toxicity values may appear on the HEAST.

In all cases, the status of a chemical may change as new data become available, and the assessment is revisited.

#### **CAUTION**

It is imperative for each user of the HEAST to recognize that the values listed in the toxicity tables and the cancer table are generally considered to be PROVISIONAL RISK ASSESSMENT INFORMATION. The user is referred to IRIS for "Work Group Verified" values. It is also important to remember that the numbers in these tables alone tell very little about the adverse effects of a chemical or the quality of evidence on which risk assessment information is based. Original assessment documents must be consulted by users of the HEAST in order to fully appreciate the strengths and limitations of a specific data base. Original source documents will allow for the most complete characterization of potential toxicity associated with the range of exposure pathways generally evaluated at Superfund and RCRA sites. The Reference Tables point the user to these sources.

# **CONTRIBUTORS**

Chemicals commonly found at RCRA sites as identified by the Office of Solid Waste's (OSW) Technical Assessment Branch are included in the HEAST. The Office of Radiation Programs has provided data on radionuclide carcinogenicity for Tables 4A and 4B of the HEAST. Finally, the Office of Air Quality Planning and Standards (OAQPS) has provided information on chemicals for which Air Quality Criteria Documents and National Ambient Air Quality Standards have been developed.

## **CHEMICALS LISTED**

Most of the chemicals included on the toxicity tables and carcinogenicity table are those for which at least one of the following EPA documents has been written: Health Effects Assessment Document (HEA), Health and Environmental Effects Profile (HEEP), Health and Environmental Effects Document (HEED), Health Assessment Document (HAD), Air Quality Criteria Document (AQCD), Drinking Water Criteria Document (DWCD). A description of each is provided in Appendix A, Section I. In a few cases, the values are supported by other written material, such as Work Group meeting notes or Carcinogen Assessment Group (CAG) Profiles. Radionuclide slope factor values are calculated by the EPA's Office of Radiation Programs.

The names of criteria pollutants that are regulated as National Ambient Air Quality Standards (NAAQS) under the Clean Air Act are listed in the main body of the HEAST, but the actual criteria are included as Section V of Appendix A: Technical Information. The NAAQS were not included in the tables in order to distinguish them from the reference concentration ([RfC]) values. The NAAQS and [RfC]s represent different levels

of review and different methods of calculation and thus, must be interpreted and used differently.

#### HIERARCHY OF SOURCES

It is recognized that at any point in time there may be multiple old and new Agency documents or data bases that present different values on a specific chemical. For chemicals other than those represented by the NAAQS or DWCDs, the following hierarchy of sources is recommended in evaluating chemical toxicity for Superfund sites:

- 1. The Agency's Integrated Risk Information System (IRIS) and cited references. Changes are made in this data base on a monthly basis, but there may be data gaps. Call IRIS USER SUPPORT at (513)569-7254 for further information.
- 2. The Health Effects Assessment Summary Tables (HEAST) and cited references.
- 3. Consultation with the Superfund Health Risk Technical Support Center (TSC) at (513)569-7300.
- 4. <u>Do not consult</u> either the toxicity tables (Appendix A) in the Superfund Public Health Evaluation Manual (SPHEM, U.S. EPA, 1986) or the September 1988 Public Health Risk Evaluation Data Base (PHRED) as these sources are likely to contain numerous values that have since become out-of-date.

#### **QUESTIONS**

## **Chemical Toxicity and Carcinogenicity**

Regional EPA Superfund Staff may direct questions regarding the contents of the chemical toxicity and carcinogenicity tables on the HEAST (e.g., chemicals not covered, chemicals with pending [RfD]s) to EPA's Superfund Health Risk Technical Support

Center (TSC) in Cincinnati, OH at (513)569-7300. Questions from other users must be submitted to the TSC in writing and must contain the following information:

- Superfund site name, site location and twelve-digit site number;
- Name and phone number of the site Remedial Project Manager (RPM) or Regional Risk Assessor/Toxicologist;
- Detailed description of the risk assessment related question.

Please send requests via mail or FAX to:

Superfund Health Risk Technical Support Center US EPA 26 W. ML King Dr. Environmental Criteria and Assessment Office MS 117 Cincinnati, OH 45268 FAX#: (513)569-7159

## **RCRA Chemicals**

Questions about RCRA chemicals may be addressed by calling the Office of Solid Waste at (202)260-4761.

# **Radionuclide Carcinogenicity**

Questions concerning radionuclide carcinogenicity should first be addressed by contacting the appropriate Regional Radiation Program Manager. A listing of these managers and several contacts in the Office of Radiation Programs can be found in Exhibit 2 of the User's Guide - Radionuclide Carcinogenicity.

#### REFERENCES

Most cited Agency references (e.g., HEAs, HEEPs, HEEDs), are (or will soon be) available through the National Technical Information Service (NTIS), 5285 Port Royal

Road, Springfield, VA 22161 [(703)487-4650]. Carcinogen Assessment Group (CAG) Profiles cited in Table 3 are available through the RCRA docket (202)260-9327.

Drinking water documents are available by calling the Drinking Water Docket at (202)260-3027.

## ORDERING INFORMATION

Limited copies of the HEAST are available for EPA Superfund staff, State Superfund programs and other Federal agencies working on Superfund sites, and EPA contractors working for the EPA Superfund program. Users in these groups can call Syracuse Research Corporation (202)479-0881 to be put on the mailing list.

EPA's Office of Solid Waste (OSW) requests that their users (i.e., OSW staff, contractors, State solid waste programs) call the Health Assessment Section (202)260-4761 to obtain copies of the HEAST. Regional OSW staff are reminded that copies are sent to all EPA Regional libraries.

Users of the HEAST in EPA's Office of Air and Radiation and State air programs should call Kelly Rimer of EPA's Office of Air Quality Planning and Standards at (919)541-2962.

All other users must purchase the document from:

National Technical Information Service (NTIS) 5285 Port Royal Road Springfield, VA 22161 (703)487-4650

For ordering information, call the NTIS Subscriptions Department at (703)487-4630. NTIS normally ships 4th class United States mail. When ordering the

1994 Health Effects Assessment Summary Table annual update from NTIS refer to the

folowing order number:

PB94-963310: Annual HEAST update

STRUCTURE OF THE HEAST

The HEAST Introduction contains explanatory material relative to the quality of

information on the HEAST, its sources, and its availability. This is followed by a listing

of changes since the last HEAST was published and then by User's Guides for both

Chemical Toxicity and Carcinogenicity, and Radionuclide Carcinogenicity. The values

on the HEAST are presented in a series of five tables that contain toxicity information

and three tables of references. The information contained in each table and their

designations are as follows:

HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER

THAN CARCINOGENICITY)

Table 1 lists subchronic and chronic non-cancer toxicity values that were calculated using the current methodology practiced by the RfD/RfC Work

Group.

HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND

CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

Table 2 lists subchronic and chronic non-cancer toxicity values that are found in Agency documents, but were calculated by alternative methods that are not currently practiced by the RfD/RfC Work Group. These values are considered to be adequate provisional values for risk assessment purposes at Superfund and RCRA sites, but are subject to being reviewed by the RfD/RfC Work Group and revised when necessary to reflect current

work group practices.

**HEAST TABLE 3: CARCINOGENICITY** 

Table 3 lists carcinogenicity values that were calculated using the current

methodology of the CRAVE Work Group.

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# HEAST TABLES 4A AND 4B: RADIONUCLIDE CARCINOGENICITY - SLOPE FACTORS

Tables 4A and 4B list ingestion, inhalation and external exposure carcinogenicity slope factors for radionuclides in two equivalent, but different units, picocuries and becquerels, respectively.

# HEAST TABLE 1 REFERENCES: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

The references for Table 1 are numerically coded to associate each toxicity value clearly with its corresponding reference.

# HEAST TABLE 2 REFERENCES: ALTERNATE METHODS – SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

The references for Table 2 are numerically coded to associate each toxicity value clearly with its corresponding reference.

## **HEAST TABLE 3 REFERENCES: CARCINOGENICITY**

The references for Table 3 are numerically coded to associate each toxicity value clearly with its corresponding reference.

Following the tables, a Technical Appendix (Appendix A) is available, containing the following sections:

- Data Sources and Selection Criteria Used in HEAST
- II Dose Conversions on HEAST
- III. Chemical Name and Chemical Abstracts Service Registry Number Cross Reference
- IV. Effect Level Definitions
- V. National Ambient Air Quality Standards (NAAQS)

#### WHAT'S NEW IN THE FY94 HEAST

## **GENERAL CHANGES - CHEMICAL TOXICITY AND CARCINOGENICITY**

The changes in this version of the HEAST reflect changes in IRIS through January 1, 1994. It is also current with RfD/RfC and CRAVE Work Group activities through January 1, 1994.

# CHEMICAL-SPECIFIC CHANGES -- CHEMICAL TOXICITY AND CARCINOGENICITY

A. CHEMICAL-SPECIFIC CHANGES ON HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

## Aldicarb 000116-06-3

The chronic oral RfD has been replaced on IRIS. The chronic oral RfD on IRIS was adopted as the subchronic oral [RfD].

## Bromoform 000075-25-2

A comment was added to indicate that the chronic inhalation RfC is considered not verifiable by the RfD/RfC (02/11/93) Work Group.

# Dichlorobenzene, 1,4- 000106-46-7

An indicator was added to show that the inhalation RfC is now available on IRIS. The chronic inhalation RfC on IRIS was modified to derive the subchronic inhalation [RfC].

## Dichloroethane, 1,2- 000107-06-2

The CAS Registry Number was corrected from 106-06-2 to 107-06-2. No other changes in the table.

#### Glycidaldehyde 000765-34-4

The NOAEL which was inadvertently omitted from the 1993 HEAST has been replaced and converted from 10 ppm to 29 mg/cu m.

#### Hexachlorocyclopentadiene 000077-47-4

No change in the tables. Reference to one of two studies used in the HEED to derive the RfC (Batelle Northwest Laboratories, 1984) was replaced in the reference section.

#### Manganese 007439-96-5

The chronic inhalation RfC has been replaced on IRIS. A subchronic inhalation [RfC] has not been derived.

#### Methoxychlor 000072-43-5

A comment was added to indicate that the chronic inhalation RfC is considered not verifiable by the RfD/RfC (11/07/91) Work Group.

# Trichloro-2'-hydroxydiphenylether, 2,2,4'- 003380-34-5

The Target and Critical effect which were inadvertently omitted from the 1993 Annual HEAST have been replaced.

B. CHEMICAL-SPECIFIC CHANGES ON HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

# Dichlorobenzene, 1,2- 000095-50-1

The subchronic inhalation [RfC] which had inadvertently been omitted from the 1993 Annual HEAST has been replaced.

# Ethoxyethanol acetate, 2- 000111-15-9

The NOEL units were corrected.

# C. CHEMICAL-SPECIFIC CHANGES ON HEAST TABLE 3: CARCINOGENICITY

#### Dimethyl hydrazine, 1,1- 000057-14-7

No EPA Group Classification was provided in the reference document, therefore this compound was removed from Table 3.

# Methyl hydrazine 000060-34-4

No EPA Group Classification was provided in the reference document, therefore this compound was removed from Table 3.

#### Metholachlor 051218-45-2

An indicator was added to show that the EPA Group Classification is now available on IRIS.

D. CHEMICAL-SPECIFIC CHANGES IN THE JULY 1993 AND NOVEMBER 1993 SUPPLEMENTS ON HEAST TABLE 4A AND 4B: RADIONUCLIDE CARCINOGENICITY -- SLOPE FACTORS

# <u>Uranium (92) U-238 +D</u>

The ingestion, inhalation, and external slope factors in Tables 4A and 4B have changed to correct an error in summation for the +D values of U-238.

## Annnouncement of Upcoming Changes to Radionuclide Slope Factors

Slope factors for radionuclides are currently being revised to incorporate EPA's revised methodology for estimating radiogenic cancer risk. The new methodology is being reviewed by experts within and outside the EPA. Users of the HEAST should expect revised radionuclide slope factors to be available sometime during 1994. The new

values and a synopsis of the revised methodology will be published in the HEAST quarterly update when available.

# CHEMICAL SPECIFIC CHANGES MADE IN THE JULY 1993 SUPPLEMENT NO. 1 AND THE NOVEMBER 1993 SUPPLEMENT NO. 2 TO THE MARCH 1993 HEAST ANNUAL UPDATE

The following changes were made in the July 1993 and November 1993 supplemental editions of the March 1993 Heast Annual Update and represent those changes that have occurred between the publication of the March 1993 HEAST Annual Update and this document, the March 1994 Heast Annual Update. Because some HEAST users may have been unaware of the publications of these supplements, the following information will indicate additional changes in toxicity information that should be noted. Note: These changes have been incorporated into the March 1994 Annual Update.

# A. CHEMICAL-SPECIFIC CHANGES IN THE JULY 1993 AND NOVEMBER 1993 SUPPLEMENTS ON HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

#### Acrylonitrile 000107-13-1

Added to Table 1. The chronic oral [RfD] under review by the RfD/RfC Work Group was modified to derive the subchronic oral [RfD].

## Arochlor 1248 012672-29-6

Added to Table 1. A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/20/93) by the RfD/RfC Work Group.

# Atrazine 001912-24-9

An indicator was added to show that the chronic oral RfD is now available on IRIS. The chronic oral RfD on IRIS was adopted as the subchronic oral [RfD].

#### Barium cyanide 000542-62-1

A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/20/93) by the RfD/RfC Work Group.

## Benzenethiol / (Thiophenol) 000108-98-5

Added to Table 1. The chronic oral [RfD] under review by the RfD/RfD Work Group was modified to derive the subchronic oral [RfD].

## Boron 007440-42-8

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Boron was added.

# Bromoethene / (Vinyl bromide) 000593-60-2

Added to Table 1. An indicator was added to show that the chronic inhalation RfC is now available on IRIS. The chronic inhalation RfC on IRIS was adopted as the subchronic inhalation [RfC].

# Bromoform 000075-25-2

A comment was added to indicate that the chronic inhalation RfC is considered not verifiable (02/11/93) by the RfD/RfC Work Group.

# Bromomethane 000074-83-9

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Bromomethane was added.

# Chlorobenzilate 000510-15-6

A comment was added to indicate that the chronic inhalation RfC is considered not verifiable (02/11/93) by the RfD/RfC Work Group.

#### Chromium(III) 016065-83-1

The subchronic oral [RfD] was changed to agree with IRIS. The uncertainty factor was changed to incorporate an additional Modifying Factor of 10 as on IRIS.

# Cresol, p- / (Methylphenol, 4-) 000106-44-5

Study information was changed to be consistent with information under review by the RfD/RfC work group. The NOEL is changed to a NOAEL, critical effects are changed, and the chronic oral [RfD] is adopted as the subchronic oral [RfD].

# Cyanazine 021725-46-2

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Cyanazine was added.

# <u>Dicamba</u> 001918-00-9

Added to Table 1. An indicator was added to show that the chronic oral RfD is available on IRIS. The chronic oral RfD was adopted as the subchronic oral [RfD].

# Dichloropropene, 1,3- / (Telone II) 000542-75-6

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for 1,3-Dichloropropene was added.

#### Dinitro-o-cresol. 4.6- 000534-52-1

A comment was added to indicate that the chronic inhalation RfC is considered not verifiable (02/11/93) by the RfD/RfC Work Group.

# Endosulfan 000115-29-7

The chronic oral RfD was withdrawn from IRIS (12/01/92). The chronic oral [RfD] was changed to reflect the value currently under review by the RfD/RfC Work Group (11/04/92). The chronic oral [RfD] was adopted as the subchronic oral [RfD].

# Hexachlorobutadiene 000087-59-1

The chronic oral RfD was withdrawn from IRIS (05/01/93). The chronic oral [RfD] was changed to reflect the value currently under review by the RfD/RfC Work Group (04/01/93).

# Isophorone 000078-59-1

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Isophorone was added.

# Manganese 007439-96-5

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Criteria Document for Manganese was added.

# Methoxyethanol, 2- 000109-86-4

The chronic oral [RfD] and the subchronic oral [RfD] were moved from Table 1 to Table 2 because they were derived from methodology that is not current with the interim methodology used by the RfD/RfC Work Group.

# Methyl ethyl ketone 000078-93-3

An indicator was added to show that the chronic oral RfD is now available on IRIS. The subchronic oral [RfD] and the subchronic inhalation [RfC] were changed to be consistent with IRIS.

## Methyl ethyl ketone peroxide 001338-23-4

Added to Table 1. A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/20/93) by the RfD/RfC Work Group

## Methylene-bis(2-chloroaniline), 4,4'- 000101-14-4

A comment was added to indicate that the chronic inhalation RfC is considered not verifiable (02/10/93) by the RfD/RfC Work Group.

#### Methyl isobutyl ketone 000108-10-1

The chronic oral [RfD] was changed to reflect the value currently under review by the RfD/RfC Work Group (07/22/93). The chronic oral [RfD] was modified to derive the subchronic oral [RfD].

#### Metolachlor 051218-45-2

Added to Table 1 from the 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Metolachlor. The chronic oral [RfD] was adopted as the subchronic oral [RfD].

# Metribuzin 021087-64-9

Added to Table 1. An indicator was added to show that the chronic oral RfD is now available on IRIS. The chronic oral RfD on IRIS was adopted as the subchronic oral [RfD].

## Naphthalene 000091-20-3

The record for Naphthalene, which was inadvertently omitted from Table 1 of the Annual HEAST, has been replaced.

# Nickel cyanide 000557-19-7

Added to Table 1. A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/20/93) by the RfD/RfC Work Group

## Nitroaniline. 2- 000088-74-4

The record for Nitroaniline, 2-, which was inadvertently omitted from Table 1 of the Annual HEAST, has been replaced.

# Osmium tetroxide 020816-12-0

Added to Table 1. A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/22/93) by the RfD/RfC Work Group

# Simazine 000122-34-9

An indicator was added to show that the chronic oral RfD is now available on IRIS. The chronic oral RfD on IRIS was adopted as the subchronic oral [RfD].

# Tetrachloroethane, 1, 1, 1, 2- 000630-20-6

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for 1,1,1,2-Tetrachloroethane was added.

#### Thallic oxide 001314-32-5

A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/20/93) by the RfD/RfC Work Group.

## Thallium Selenite 012039-52-0

The chronic oral RfD was withdrawn from IRIS (08/93). A comment was added to indicate that the chronic oral RfD is considered not verifiable (07/20/93) by the RfD/RfC Work Group

## Trichlorobenzene, 1,2,4- 000120-82-1

The chronic inhalation [RfC] was removed from Table 2. The chronic inhalation [RfC] under review by the RfD/RfC Work Group was added to Table 1. The chronic inhalation [RfC] under review by the RfD/RfC Work Group was modified to derive the subchronic inhalation [RfC].

# Trifluralin 001582-09-8

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Trifluralin was added.

# Zinc (metallic) 007440-66-6

The record for Zinc (metallic), which was inadvertently omitted from Table 1 of the Annual HEAST, has been replaced.

B. CHEMICAL-SPECIFIC CHANGES IN THE JULY 1993 AND NOVEMBER 1993 SUPPLEMENTS ON HEAST TABLE 2. SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

# Methoxyethanol, 2- 000109-86-4

The chronic oral [RfD] and the subchronic oral [RfD] were moved from Table 1 to Table 2 because they were derived from methodology that is not current with the interim methodology used by the RfD/RfC Work Group.

# <u>Trichlorobenzene, 1,2,4 000120-82-1</u>

The chronic inhalation [RfC] was removed from Table 2 and the chronic inhalation [RfC] under review by the RfD/RfC Work Group was added to Table 1.

C. CHEMICAL-SPECIFIC CHANGES IN THE JULY 1993 AND NOVEMBER 1993 SUPPLEMENTS ON HEAST TABLE 3: CARCINOGENICITY

# Chloromethane / (Methyl chloride) 000074-87-3

The record for Chloromethane, which was inadvertently omitted from Table 3 of the Annual HEAST, has been replaced.

#### Coke Oven Emissions 008007-45-2

An indicator was added to show that the EPA Group classification is now on IRIS.

# Cyanazine 021725-46-2

Carcinogenicity information was added to Table 3 of the July 1993 Supplement from the 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Cyanazine. The record was revised (cancers reported and the references changed) on the November 1993 Supplement to reflect CRAVE Work Group review. There was no change in quantitative values.

# Dibromo-3-chloropropane, 1,2- 000096-12-8

The inhalation [slope] and the inhalation [unit risk] were inadvertently reversed on the July 1992 Supplement 1.

# Dichloropropene, 1,3- / (Telone II) 000542-75-6

The record was revised (cancers reported and the references changed) to reflect the information currently under review by the CRAVE Work Group.

## Ethylene thiourea 000096-45-7

The oral [slope] and the oral [unit risk] were changed to reflect the values currently under review by the CRAVE Work Group.

# Methylenebis(benzeneamine), 4,4- 000101-77-9

No [EPA Group] classification was provided in the reference document, therefore the compound was removed from Table 3.

## Metolachlor 051218-45-2

Added to Table 3 from the 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for Metolachlor.

## Mirex 002385-85-5

The oral [slope] and the oral [unit risk] are under review by the CRAVE Work Group and were removed from Table 3.

# TCDD, 2,3,7,8- 001746-01-6

The units for the inhalation [unit risk] were added to Table 3. There are no additional changes to the Table.

# Tetrachloroethane, 1, 1, 1, 2 - 000630-20-6

No change in the tables. Reference to 1993 Revised and Updated Drinking Water Quantification of Toxicologic Effects for 1,1,1,2-Tetrachloroethane was added.

# Trichloropropane, 1,2,3- 000096-18-4

The oral [slope] and the oral [unit risk] being considered by the CRAVE Work Group and were added to Table 3.

D. CHEMICAL-SPECIFIC CHANGES IN THE JULY 1993 AND NOVEMBER 1993 SUPPLEMENTS ON HEAST TABLE 4A AND 4B: RADIONUCLIDE CARCINOGENICITY -- SLOPE FACTORS

No new radionuclide slope factors were added to Tables 4A and 4B, and none of the slope factors listed in the March 1993 HEAST Annual Update were changed.

# **USER'S GUIDE: CHEMICAL TOXICITY**

The HEAST summarizes provisional toxicity and cancer values as well as values developed for the NAAQS and DWCD chemicals. The provisional status of the toxicity and cancer values is indicated by placing brackets around the title of the value. These include provisional reference concentrations ([RfC]) and provisional reference doses ([RfD]) for toxicity from subchronic and chronic inhalation and oral exposure (Tables 1 and 2) and provisional slope factors ([slope factor]), provisional cancer classifications ([EPA Group]) and provisional unit risk values ([unit risk]) for carcinogenicity, based on lifetime inhalation and oral exposure (Table 3). Brackets should be included with the acronym whenever a user quotes the value in an assessment document, and the provisional nature of the value should be noted. A more complete discussion of how Superfund develops and considers the toxicity assessment in hazardous waste sites is presented in Chapter 7 of Risk Assessment Guidance for Superfund Volume 1: Human Health Evaluation Manual, Part A, EPA/540/1-89/002.

The references listed for each chemical in the Reference Tables for Tables 1, 2 and 3 represent the study or studies that are the basis for the [RfC], [RfD], [slope factor], [EPA Group], or [unit risk], as well as the EPA reference that is the source of the Agency analysis or risk assessment information. In some cases, additional EPA documents are also listed as a source of information on the chemical. Work Group verified values found on IRIS are not found on the HEAST, but are indicated in the tables by the word "IRIS" in place of the number.

# TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINO-GENICITY)

The [RfC] or [RfD] is a provisional estimate (with uncertainty spanning perhaps an order of magnitude) of the daily exposure to the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a portion of the lifetime, in the case of a subchronic [RfC] or [RfD], or during a lifetime, in the case of a chronic [RfC] or [RfD]. The [RfC] and [RfD] values are listed in Tables 1 and 2 in columns with the headings "Subchronic" and "Chronic". The critical dose or concentration level is usually a No-Observed-Adverse-Effect Level (NOAEL) or a Lowest-Observed-Adverse-Effect Level (LOAEL) (See Appendix A, Section IV: Effect Level Definitions, for more information). The [RfC] or [RfD] is derived by dividing the NOAEL or LOAEL by an uncertainty factor (UF) times a modifying factor (MF):

$$[RfC]$$
 or  $[RfD] = \frac{NOAEL \text{ or } LOAEL}{UF \times MF}$ 

In Tables 1 and 2, the information listed is the following:

Chemical = Chemical Name/CASRN

Level = Effect Level

Dose = Administered Dose or Concentration

Route = Route of Administration

Species = Tested Species Experiment Length = Length of Exposure

Target = Target Organ(s) Affected at Critical Level

Critical Effect = Effect(s) Observed at Critical Level

Subchronic [RfC] = Subchronic Inhalation [Reference Concentration]

UF = Uncertainty Factor for the Subchronic Inhalation

[Reference Concentration]

Subchronic [RfD] = Subchronic Oral [Reference Dose]

UF = Uncertainty Factor for the Subchronic Oral [Reference

Dosel

Chronic [RfC] = Chronic Inhalation [Reference Concentration]

UF = Uncertainty Factor for the Chronic Inhalation [Reference

Concentration]

Chronic [RfD] = Chronic Oral [Reference Dose]

UF = Uncertainty Factor for the Chronic Oral [Reference Dose]

Reference = Reference Identification Number for All Toxicity Values

on the Same Line.

An example of this information is shown in Figure 1, HEAST Table 1:

Chemical = GLYCIDALDEHYDE/000765-34-4

Level = NOAEL Dose = 10 PPM

Route = INHALATION: INTERMITTENT

Species = RAT

Experiment Length = 12 WEEKS

Target = WHOLE BODY, BLOOD, KIDNEY

Critical Effect = DECREASED WEIGHT GAIN, HEMATOPOIETIC

EFFECTS, EFFECTS

Subchronic [RfC] = 1E-2 mg/cu.m

UF = 300

Subchronic [RfD] = 4E-3 mg/kg/day

UF = 300

Chronic [RfC] = 1E-3 mg/cu.m

UF = 3000 Chronic [RfD] = IRIS UF = IRIS Reference = 005968

Notice that a Chronic RfD for Glycidaldehyde is available on IRIS, so it is not listed here. Also notice that there are footnotes for this chemical that indicate a route-to-route extrapolation was performed and that there is information available on Table 3: Carcinogenicity.

Also given in Figure 1 is an example of the References for Table 1 for the same chemical. The reference is identified by the chemical name (Glycidaldehyde), the CASRN (00765-34-4), and the reference number that links it with the toxicity values (005968).

The uncertainty factor used in calculating the [RfC] or [RfD] reflects scientific judgment regarding the various types of data used to estimate [RfC] or [RfD] values. An

# -22-

#### FIGURE 1

## Example Data and References for Chemical Toxicity

# HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) January 1992

<u>CHEMICA</u> LEVEL	L <u>Dose</u> Route	<u>SPECIES</u> EXPERIMENT LENGT	H TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu_m) (mg/kg/day UF UF		Chr [RfC] (mg/cu m) UF	onic [RfD] (mg/kg/day) UF	REFERENCE	
GLYCIL NOAEL	OALDEHYDE 10 PPM INHALATION INTERMITTENI	RAT 12 WEEKS	765-34-4 WHOLE BODY BLOOD KIDNEY	DECREASED WEIGHT GAIN HEMATOPOIETIC EFFECTS EFFECTS	1E-2 300	4E - 3 300	1E-3 3000	IRIS	005968	

SUBCHRONIC [RfD] COMMENT BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0 5 CHRONIC [RfD] COMMENT. BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0 5 GENERAL COMMENT. ALSO SEE TABLE 3 CARCINOGENICITY

# REFERENCES FOR HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

January 1992

GLYCIDALDEHYDE 000765-34-4

005968 HINE CH. RJ GUZMAN, MK DUNLAP, R LIMA AND GS LOQUVAM 1961 STUDIES ON THE TOXICITY OF GLYCIDALDEHYDE ARCH ENVIRON HEALTH 2: 23-30

US EPA 1989 HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENTFOR GLYCIDALDEHYDE PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON, DC

uncertainty factor of 10 is usually used to account for variation in human sensitivity among populations. An additional 10-fold factor is usually used to account for each of the uncertainties assumed when extrapolating from animal data to humans, when extrapolating from a LOAEL to a NOAEL, and when extrapolating from subchronic to chronic exposure. In order to reflect professional assessment of the uncertainties of the study and the data base not explicitly addressed by the above uncertainty factors (e.g., completeness of the overall data base), an additional uncertainty factor or modifying factor ranging from greater than 0 to less than or equal to 10 is applied. The default value for this modifying factor is 1.

For chemicals for which a chronic [RfC] or [RfD] is presented in Tables 1 and 2, a subchronic [RfC] or [RfD] is usually derived, if not previously derived in the Agency documents that originally addressed the chemical. Subchronic toxicity values are not evaluated by the RfD/RfC Work Group. The subchronic [RfC] or [RfD] is derived in either of two ways: 1) If an uncertainty factor was used to account for extrapolation from subchronic to chronic exposure in the derivation of the chronic [RfC] or [RfD], then, the subchronic [RfC] or [RfD] is derived from the same benchmark concentration or dose without applying the uncertainty factor for subchronic to chronic exposure extrapolation.

2) If the chronic [RfC] or [RfD] was derived without use of an uncertainty factor for extrapolating from subchronic to chronic exposure (e.g., if chronic data were available), then, the chronic [RfC] or [RfD] is adopted as the subchronic [RfC] or [RfD].

Tables 1 and 2 list the uncertainty factor and modifying factor, multiplied together, to form a single factor under the heading "Uncertainty Factor." For example, the uncertainty factor of 3000 listed for the chronic inhalation [RfC] for Glycidaldehyde reflects an uncertainty factor of 1000 (10 for human sensitivity, 10 for extrapolation from

animal to human, and 10 for extrapolation from subchronic to chronic) and a modifying factor of 3 (for an inadequate data base); the uncertainty factor of 500 listed for the subchronic oral [RfD] for cyanide reflects an uncertainty factor of 100 (10 for human sensitivity, and 10 for extrapolation from animal to human) and a modifying factor of 5 (to account for tolerance to cyanide when ingested by food rather than administration by gavage or by drinking water).

[RfC] and [RfD] values are specific for the route of exposure for which they are listed on Tables 1 and 2. In the few instances where an [RfD] or [RfC] has been determined from another exposure route, route-to-route extrapolation is indicated by a footnote.

The current methodology for the derivation of inhalation RfCs is detailed in the document, "Interim Methods for Development of Inhalation Reference Doses" (U.S. EPA, 1990, EPA/600/8-88/066F, NTIS PB90-145723). These methods are different from those used for oral RfDs because of (1) the dynamics of the respiratory system and its diversity across species, and (2) differences in the physicochemical properties of contaminants (such as the size and shape of a particle or whether the contaminant is an aerosol or a gas). Parameters such as deposition, clearance mechanisms and the physicochemical properties of the inhaled agent are considered in the determination of the effective dose delivered to the target organ.

An RfC value calculated using this interim methodology is generally reported as a concentration in air (mg/m³), although it may be converted to a corresponding inhaled dose (mg/kg/day) by dividing by 70 kg (an assumed human body weight), multiplying by 20 m³/day (an assumed human inhalation rate), and adjusting by an appropriate absorption factor. This conversion, however, may often be technically incorrect, and the

appropriateness of doing this must be evaluated on a case-by-case basis. It is recommended that HEAST users that plan to use this technique read a further discussion of the difficulties inherent in this dose conversion that can be found in Appendix A, Section II: Dose Conversions On HEAST.

Inhalation [RfC] values reported in HEAs and early HEEDs that were finalized prior to the implementation of the interim methods were calculated using methods similar in concept to those used for oral [RfD]s. These values are reported both as a concentration in air (in mg/m³ for continuous, 24 hours/day exposure) under the column [RfC], and as a corresponding inhaled dose (in mg/kg/day) in the footnotes called, Chronic (Subchronic) [RfC] Comment. These chemicals are listed in Table 2: Alternate Methods - Subchronic and Chronic Toxicity (Other Than Carcinogenicity).

[RfD] values for oral exposure are reported as mg/kg/day. An oral [RfD] value can be converted to a corresponding concentration in drinking water, assuming human body weight of 70 kg and water consumption of 2 L/day, as follows:

The [RfC] or [RfD] is used as a reference point for gauging the potential effects of other exposures. Usually, exposures that are less than the [RfC] or [RfD] are not likely to be associated with health risks. As the frequency of exposures exceeding the [RfC] or [RfD] increases and as the size of the excess increases, the probability increases that adverse health effects may be observed in a human population. Nonetheless, a clear distinction that would categorize all exposures below the [RfC] or [RfD] as "acceptable" (risk-free) and all exposures in excess of the [RfC] or [RfD] as

"unacceptable" (causing adverse effects) cannot be made. In addition, [RfC] and [RfD] values, and particularly those with limitations in the quality or quantity of supporting data, are subject to change as additional information becomes available.

When [RfC] or [RfD] values are listed in Tables 1 or 2 for chemicals that are carcinogens, a footnote will refer to Table 3 if additional information concerning carcinogenicity is available in that table. [RfC] and [RfD] values that have been derived for carcinogens are based on noncancer endpoints only and should not be assumed to be protective against carcinogenicity.

# TABLE 2: ALTERNATE METHODS — SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

Chemicals are listed in Table 2 when the [RfD] or [RfC] was derived from alternative methods that are not currently practiced by the RfD/RfC Work Group. The table consists primarily of inhalation [RfC] values determined from methodology that does not follow the interim inhalation methods adopted by the Agency, and [RfC] or [RfD] values based on route-to-route extrapolation with inadequate pharmacokinetic and toxicity data. A footnote is added to each chemical to provide a short explanation of the specific methodology used in calculating these provisional toxicity values. Most of these toxicity values were formerly listed in Table 1. In some instances, the chemical may be listed in both Tables 1 and 2 if the chemical has more than one toxicity value. Table 2 follows the same format as Table 1 (refer to Figure 1).

# **TABLE 3: CARCINOGENICITY**

In assessing the carcinogenic potential of a chemical, the Human Health Assessment Group (HHAG) of EPA classifies the chemical into one of the following groups, according to the weight of evidence from epidemiologic and animal studies:

- Group A Human Carcinogen (sufficient evidence of carcinogenicity in humans)
- Group B Probable Human Carcinogen (B1 limited evidence of carcinogenicity in humans; B2 sufficient evidence of carcinogenicity in animals with inadequate or lack of evidence in humans)
- Group C Possible Human Carcinogen (limited evidence of carcinogenicity in animals and inadequate or lack of human data)
- Group D Not Classifiable as to Human Carcinogenicity (inadequate or no evidence)
- Group E Evidence of Noncarcinogenicity for Humans (no evidence of carcinogenicity in adequate studies).

These classifications are shown under [EPA Group] on Table 3.

Quantitative carcinogenic risk assessments are performed for chemicals in Groups A and B, and on a case-by-case basis for chemicals in Group C. Cancer [slope factors] (formerly called cancer potency factors in the Superfund Public Health Evaluation Manual) are estimated through the use of mathematical extrapolation models, most commonly the linearized multistage model, for estimating the largest possible linear slope (within the 95% confidence limit) at low extrapolated doses that is consistent with the data. The [slope factor] or risk is characterized as an upper-bound estimate, i.e., the true risk to humans, while not identifiable, is not likely to exceed the upper-bound estimate and in fact may be lower.

Quantitative carcinogenic estimates listed in Table 3 include the following:

[slope factor] = risk per unit dose = risk per mg/kg/day

[unit risk] for inhalation exposure = risk per concentration unit in air = risk per  $\mu$ g/m<sup>3</sup>

[unit risk] for oral exposure = risk per concentration unit in water = risk per  $\mu\alpha/L$ 

[Unit risk] estimates for inhalation and oral exposure can be calculated by dividing the appropriate [slope factor] by 70 kg and multiplying by the inhalation rate (20 m<sup>3</sup>/day) or the water consumption rate (2 L/day), respectively, for risk associated with unit concentration in air or water. Hence,

risk per 
$$\mu$$
g/m³ (air) = (risk per mg/kg/day) x 10 x 20 m³/day x 10 (mg/ $\mu$ g) 70 kg

risk per 
$$\mu$$
g/L (water) = (risk per mg/kg/day) x 1 x 2 L/day x 10<sup>-3</sup> (mg/ $\mu$ g) 70 kg

Quantitative estimates of carcinogenic risk are listed under [Unit Risk] or [Slope Factor] in Table 3. Information on the study and data set used for estimation of the [slope factor] is given in the other columns of Table 3.

In Table 3, the information listed is the following:

Chemical = Chemical Name/CASRN Route = Route of Administration

Species = Tested Species **Experiment Length** = Length of Exposure

Target = Target Organ(s) Affected at Critical Level Cancer

= Tumors Observed at Critical Level (Not Specified

if More Than One Type of Tumor)

[EPA Group] = EPA Classification by Weight of Evidence

Oral [Slope Factor] = Risk Per Unit Dose Inhalation [Slope Factor] = Risk Per Unit Dose

Oral [Unit Risk] = Risk Per Concentration Unit in Water Inhalation [Unit Risk] = Risk Per Concentration Unit in Air

Reference = Reference Identification Number for All Toxicity Values on the Same Line.

An example of this information is shown in Figure 2, HEAST Table 3:

Chemical = DIMETHYLHYDRAZINE, 1,2-/000077-78-1

Route = ORAL: DRINKING WATER

Species = MOUSE Experiment Length = LIFETIME

Target = CARDIOVASCULAR SYSTEM

Cancer = TUMORS

[EPA Group] = B2

Oral [Slope Factor] = 3.7E+1 (MG/KG/DAY)-1 Inhalation [Slope Factor] = 3.7E+1 (MG/KG/DAY)-1

Oral [Unit Risk] = 1.1E-3 (UG/L)-1 Inhalation [Unit Risk] = 1.1E-2 (UG/CU M)-1

Reference = 009993

Notice that the inhalation values for 1,2-Dimethylhydrazine was extrapolated from the oral data.

Also given in Figure 2 is an example of the References for Table 3 for the same chemical. The reference is identified by the chemical name (Dimethylhydrazine, 1,2-), the CASRN (000077-78-1), and the reference number that links it with the toxicity values (009993).

Quantitative carcinogenic estimates are specific for the route of exposure for which they are listed on Table 3. Footnotes are used to indicate those instances in which the values for inhalation or oral exposure are based on extrapolation from another route of exposure. The route-to-route conversion required to present inhalation [slope factors] in the units of mg/kg/day is considered by the CRAVE Work Group to be technically incorrect. It is recommended that HEAST users who plan to use this information read a further discussion of the difficulties inherent in this dose conversion which can be found in Appendix A, Section II: Dose Conversions On HEAST.

To estimate risk-specific concentrations in air from the [unit risk] in air as presented in Table 3, the specified level of risk is divided by the [unit

# <del>-</del>30-

#### FIGURE 2

# Example Data and References for Carcinogenicity

# HEAST TABLE 3: CARCINOGENICITY

January 1992

January 1992

CHEMICAL ROUTE	ERIMENT LENG SPECIES	TH TARGET	CANCER	[EPA GROUP]	OŘAL	FACTOR] INHALATION (mg/kg/day)			REFERENCE
DIMETHYLHYDRAZINE,1,2- 000077 78-1 ORAL DRINKING LIFETIME WATER				B2	3 7E+1	3 7E+1	1 1E-3	1 1E-2 (	009993
WATER	MOUSE	CARDIOVASCULAR SYSTEM	TUMORS						

Inhalation [Slope] Comment BASED ON ROUTE TO ROUTE EXTRAPOLATION

# REFERENCES FOR HEAST TABLE 3: CARCINOGENICITY

DIMETHYLHYDRAZINE, 1,2000077-78-1
009993 TOTH B AND K PATEL 1982 CARCINOGENICITY DOSE-RESPONSE STUDY BY CONTINUOUS ADMINISTRATION OF 1,2-DIMETHYLHYDRAZINE
DI-HYDROCHLORIDE IN MICE I LIGHT AND TRANSMISSION ELECTRON MICROSCOPIC STUDY OF COLONOIC NEOPLASMS AM. J OF PATH 84 69-86.

US EPA 1988 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

risk] for air. Hence, the air concentration (in  $\mu$ g/m³) corresponding to an upper-bound increased lifetime cancer risk of 1x10<sup>-5</sup> is calculated as follows:

$$\mu g/m^3$$
 in air =  $\frac{1x10^{-5}}{[unit \ risk]}$  in  $(\mu g/m^3)^{-1}$ 

To estimate risk-specific concentrations in drinking water from the oral [slope factor] values presented in Table 3, the specified level of risk is multiplied by 70 kg and divided by the [slope factor] times 2 L/day. Hence, the water concentration corresponding to an upper-bound increased lifetime cancer risk of 1x10<sup>-5</sup> is calculated as follows:

$$mg/L$$
 in water = 
$$\frac{1x10^{-5} \times 70 \ kg}{[slope factor] \ in \ (mg/kg/day)^{-1} \times 2 \ L/day}$$

# **USER'S GUIDE: RADIONUCLIDE CARCINOGENICITY**

#### INTRODUCTION

EPA classifies all radionuclides as Group A (known human) carcinogens. HEAST Tables 4A and 4B list ingestion, inhalation and external exposure cancer slope factors for radionuclides in units of picocuries and becquerels, respectively. Ingestion and inhalation slope factors are best estimates (i.e., median or 50th percentile values) of the age-averaged, lifetime excess cancer incidence (fatal and nonfatal cancer) risk per unit of activity inhaled or ingested, expressed as risk/pCi or risk/Bq. External exposure slope factors are best estimates of the lifetime excess cancer incidence risk for each year of exposure to external radiation from photon-emitting radionuclides distributed uniformly in a thick layer of soil, and are expressed as risk/yr per pCi (or Bq)/gram of soil. When combined with site-specific media concentration data and appropriate exposure assumptions<sup>2</sup>, slope factors can be used to estimate lifetime cancer risks to members of the general population due to radionuclide exposures.

<sup>1</sup> Slope factors are reported in Tables 4A and 4B in two different units of activity. Table 4A presents slope factors in the customary units of picocuries (1 pCi =  $10^{-12}$  curies (Ci) =  $3.7 \times 10^{-2}$  nuclear transformations per second) for consistency with the system used for radionuclides in the IRIS database; Table 4B presents slope factors in the International System (SI) units of becquerels (1 Bq = 1 nuclear transformation per second; approximately 27 pCi). Users can calculate cancer risks using slope factors expressed in either customary units or SI units with equivalent results, provided that they also use air, water and soil concentration values in the same system of units.

<sup>2</sup> Agency standardized default exposure scenarios and assumptions for use in baseline risk assessment are provided in EPA (1991), *Human Health Evaluation Manual*, *Supplemental Guidance*: "Standard Default Exposure Factors" (Interim Final), Office of Emergency and Remedial Response, OSWER Directive 9285.6-03. [NTIS order number: PB 91-921314.]

## INTENDED USERS AND APPLICATIONS

HEAST users include individuals from the EPA, other Federal agencies, States and contractors who are responsible for the identification, characterization and remediation of sites contaminated with radioactive materials. Radionuclide slope factors are calculated by EPA's Office of Radiation and Indoor Air (ORIA) to assist HEAST users with risk-related evaluations and decision-making at various stages of the remediation process. During site assessment, for example, slope factors are used in EPA's Hazard Ranking System (HRS) to assign toxicity factor values to radionuclides to calculate site scores. During the remedial investigation and feasibility study (RI/FS), slope factors are used to determine baseline site risk, to develop preliminary remediation goals, and to evaluate cleanup alternatives. For further examples on the application of radionuclide slope factors in risk evaluations, users are referred to the following EPA documents:

- Hazard Ranking System (HRS), Federal Register (55 FR 515320), December 1990.
- Risk Assessment Guidance for Superfund; Volume I Human Health Evaluation Manual (RAGS/HHEM), Part A, Baseline Risk Assessment (EPA/540/1-89/002).
- RAGS/HHEM Part B, Development of Risk-Based Preliminary Remediation Goals (OSWER Directive 9285.7-01B). [NTIS order number: PB 92-963333.]
- RAGS/HHEM Part C, Risk Evaluation of Remedial Alternatives (OSWER Directive 9285.7-01C). [NTIS order number: PB 92-963334.]

Copies of RAGS/HHEM Parts A, B and C are available to the public from the National Technical Information Service (NTIS) at (703)487-4650. Copies are available to EPA staff by calling the Superfund Documents Center at (202)260-9760.

# **RADIATION EFFECTS**

Ionizing radiation has been shown to be a carcinogen, a mutagen, and a teratogen. At sufficiently high doses, radiation acts as a complete carcinogen, serving as both an initiator and promoter, and can induce cancers in nearly any tissue or organ in both humans and animals. At lower doses and dose rates, radiation produces delayed responses in the form of increased incidence of cancer long after the exposure period. These delayed responses have been documented extensively in epidemiological studies of Japanese atomic bomb survivors, underground uranium miners, radium dial painters, and patients subject to a variety of radiation treatments. Laboratory animal research and mammalian tissue culture studies have provided additional, collaborative data.

Mutagenic effects of radiation have been demonstrated primarily in animal and tissue culture studies; limited data from studies of A-bomb survivors indicate that humans may be as sensitive or less sensitive than animals to radiogenic mutagenicity. Data are also available from both human and animal studies on the teratogenic effects of radiation. These data show that the fetus is most sensitive to radiation injury during the early stages of organ development (between 8 and 15 weeks for the human fetus). Resultant radiation-induced malformations depend on which cells are most actively differentiating at the time of exposure.

EPA classifies all radionuclides as Group A carcinogens, based on their property of emitting ionizing radiation and on the extensive weight of evidence provided by epidemiological studies of radiogenic cancers in humans. At Superfund radiation sites, EPA generally evaluates potential human health risks based on the radiotoxicity, i.e., adverse health effects caused by ionizing radiation, rather than on the chemical toxicity, of each radionuclide present. These evaluations consider the carcinogenic effects of

radionuclides only. In most cases, cancer risks are limiting, exceeding both mutagenic and teratogenic risks.

# **DERIVATION OF RADIONUCLIDE SLOPE FACTORS**

EPA's Office of Radiation and Indoor Air (ORIA) calculates radionuclide slope factor values using health effects data and dose and risk models from a number of national and international scientific advisory commissions and organizations, including the National Academy of Sciences (NAS), the National Council on Radiation Protection and Measurements (NCRP), the United Nations Scientific Committee on the Effects of Atomic Radiation (UNSCEAR), and the International Commission on Radiological Protection (ICRP). A detailed discussion of ORP's approach and assumptions is provided in *Risk Assessment Methodology, Environmental Impact Statement, NESHAPS for Radionuclides, Background Information Document - Volume I* (EPA 520/1-89-005: September 1989).

Radionuclide slope factors are calculated for each radionuclide individually, based on its unique chemical, metabolic and radioactive properties. The calculation uses EPA's computer code RADRISK<sup>3</sup>, life table analyses (described in EPA 520/1-89-005), and cancer risk estimates based largely on the results of the NAS BEIR III report.<sup>4</sup> Ingestion and inhalation slope factors for radionuclides account for:

<sup>3</sup> Dunning, D.E. Jr., Leggett, R.W., and Yalcinatas, M.G. (1980). "A Combined Methodology for Estimating Dose Rates and Health Effects from Exposure to Radioactive Pollutants." ORNL/TM-7105.

<sup>4</sup> National Academy of Sciences (1980). The Effects on Populations of Exposure to Low Levels of Ionizing Radiation, Report of the Committee on the Biological Effects of Ionizing Radiation (BEIR III), National Research Council, Washington, D.C. EPA anticipates that the slope factors for radionuclides in HEAST will be modified in the future to incorporate the results of BEIR V (1990) and/or other relevant sources.

- the amount of radionuclide transported into the bloodstream from either the gastrointestinal (GI) tract following ingestion, or from the lungs following inhalation;
- the ingrowth and decay of radioactive progeny produced within the body subsequent to intake;
- the distribution and retention of each radionuclide (and its associated progeny, if appropriate) in body tissues and organs;
- the radiation dose delivered to body tissues and organs from the radionuclide (and its associated progeny, if appropriate); and
- the sex, age, and organ-specific risk factors over the lifetime of exposure.

The slope factors are the average risk per unit intake or exposure for an individual in a stationary population with vital statistics (mortality rates) of the United States in 1970. (The expected lifetime for an individual in this population is about 70 years.) Consequently, radionuclide ingestion and inhalation slope factors are <u>not</u> expressed as a function of body weight and time, and do <u>not</u> require corrections for gastrointestinal absorption or lung transfer efficiencies.

NOTE: The GI absorption values  $(f_1)$ , ICRP lung classifications (D, W, Y) and radioactive half-lives are provided in HEAST Tables 4A and 4B for reference only and should not be used to correct, modify, or in any way adjust radionuclide slope factors or intake assumptions in risk calculations.

External slope factors provide cancer risk estimates per unit exposure to a uniform radionuclide concentration in soil. These factors, which account for photon energy flux attenuation and buildup in soil, are calculated for each radionuclide using volume and surface dose factors derived using the computer code DFSOIL.<sup>5</sup>

Because of the radiation risk models employed for both internal and external exposures, slope factors for radionuclides are characterized as best estimates (i.e., median or 50th percentile values) of the age-averaged lifetime total excess cancer incidence risk per unit intake or exposure.

## ABOUT THE INFORMATION PROVIDED IN TABLES 4A AND 4B:

Tables 4A and 4B list ingestion, inhalation and external exposure slope factors of principal radionuclides, and provide key parameter values used in the derivation of slope factor values. In both tables, radionuclides are presented alphabetically by element and atomic weight.

Selected radionuclides that have radioactive <u>Decay products</u> are designated within the HEAST tables with the suffix "+D" (e.g., U-238+D, Ra-226+D, Cs-137+D) to indicate

Sjoreen, A.L., Kocher, D.C., Killough, G.G. and Miller C.W. (1984). "MLSOIL and DFSOIL - Computer Codes to Estimate Effective Ground Surface Concentrations for Dose Computations." Oak Ridge National Laboratory. Oak Ridge, TN, ORNL-5974.

that the cancer risk estimates for such radionuclides include the contributions from either <u>all</u>, or only <u>some</u>, of their decay products. The slope factor for a radionuclide marked with a "+D" accounts for risk from all direct progeny with half-lives of less than one year. The slope factor for U-238+D, for example, includes the risk from U-238, Th-234, and Pa-234, but not from U-234 or any of its progeny.

Decay sub-chains included in the HEAST Tables 4A and 4B are identified in Exhibit 1, and members of such sub-chains are noted in the HEAST tables with a dagger ("†"). The rationale for the one year half-life cutoff for members of a sub-chain is that such radionuclides are likely to achieve secular equilibrium with the parent over the time scales commonly encountered in environmental remediation, while radionuclides with longer half-lives may not.

When a radionuclide is in secular equilibrium with <u>all</u> its progeny, one should add the slope factors for all the relevant sub-chains. To assess risk from Ra-226 in equilibrium with all its progeny, for example, one adds the slope factors for Ra-226+D and Pb-210+D.

In most cases, site-specific analytical data should be used to establish the actual degree of equilibrium between each parent radionuclide and its decay products in each media sampled. However, in the absence of empirical data, the "+D" values for radionuclides should be used unless there are compelling reasons not to. For example, the external slope factors for Cs-137 and Cs-137+D are 0.0 and 2x10<sup>-6</sup> (risk per year per pCi/gram), respectively. The value for Cs-137+D is higher because it includes the risk contribution from cesium's short-lived gamma-emitting decay product Ba-137m (half-life, 25.5 minutes) which, under most environmental conditions, will be in secular equilibrium with Cs-137.

Note that there may be circumstances, such as long disposal times or technologically enhanced concentrations of naturally occurring radionuclides, that may necessitate the combination of the risks of a parent radionuclide and its decay products over several contiguous subchains. For example, Ra-226 soil analyses at a site might show that all radium decay products are present in secular equilibrium down to stable Pb-206 (See Exhibit 1). In this case, Ra-226 risk calculations should be based on the ingestion, inhalation and external exposure slope factors for the Ra-226+D subchain, plus the ingestion, inhalation and external exposure factors for the Pb-210+D subchain. For actual sites, users should consult with a health physicist or radiochemist (1) to evaluate the site-specific analytical data to determine the degree of equilibrium between parent radionuclides and decay members of contiguous decay chains, and (2) to assist in the combination of appropriate slope factor values. For health physics and radioanalytical support, HEAST users may contact EPA's Regional Radiation Program Staff, ORIA's National Air and Radiation Laboratory (NAREL) in Montgomery, Alabama, ORIA's Las Vegas Laboratory (ORP-LV) in Las Vegas, Nevada, or the ORIA staff at EPA Headquarters in Washington, DC, listed in Exhibit 2.

A Chemical Abstract System Reference Number (CASRN) is assigned to each radionuclide for identification and reporting accuracy during risk assessments. Users should report a CASRN for <u>all</u> radionuclides (both parent and decay product radionuclides) included in risk calculations. Radioactive half-lives are provided in the HEAST tables for reference.

The designations "D", "W", and "Y" presented under the heading "ICRP Lung Class" in the tables refer to the lung clearance times for inhaled particulate radionuclides, expressed as days (D), weeks (W), or years (Y), as recommended by the International

Commission on Radiological Protection (ICRP). Gaseous radionuclides, e.g., Rn-222, are designated with an asterisk ("\*"). "GI Absorption Factors, f<sub>1</sub>" are the fractional amounts of each radionuclide that may be absorbed from the gastrointestinal (GI) tract into blood following an oral intake. The ICRP lung clearance classifications and GI absorption factors provided in Tables 4A and 4B are the default values that EPA used to calculate radionuclide slope factors for inhalation and ingestion exposures, respectively. These factors are provided for reference only (see the Note Box on the previous page).

## WHERE TO ADDRESS QUESTIONS ABOUT RADIONUCLIDE SLOPE FACTORS

EPA continuously reviews the scientific literature on radiation effects to ensure that the Agency's risk assessment methodologies are consistent with current models and assumptions. As risk assessment methodologies are refined, the slope factors in Tables 4A and 4B will be revised and updated.

HEAST users with questions about radionuclide slope factor values and their use in radiation risk assessments should contact the Remedial Guidance Section of the Radiation Assessment Branch of ORIA at (202)233-9350. Written requests for assistance can be sent by fax to (202)233-9650.

Exhibit 1: Radioactive Decay Chains Included in HEAST Tables 4A and 4B\*

Principal Decay Chain	Subchain*	Members <sup>b</sup>	Half-life <sup>c</sup>
Uranium-238	U-238+D	U-238 Th-234 Pa-234	4.468E+09 Y 2.410E+01 D 1.170E+00 M
	U-234	U-234	2.445E+05 Y
	Th-230	Th-230	7.700E+C4 Y
	Ra-226+D	Ra-226 Rn-222'' Po-218 Pb-214 Bi-214 Po-214	1.600E+03 Y 3.823E+00 D 3.050E+00 M 2.680E+01 M 1.990E+01 M 1.637E-04 S
	Pb-210+D	Pb-210 Bi-210 Po-210	2.226E+01 Y 5.013E+00 D 1.384E+02 D
	Pb-206	Pb-206	[Stable]
Uranium-235	U-235+D	U-235 Th-231	7.038E+08 Y 2.552E+01 H
	Pa-231	Pa-231	3.726E+04 Y
	Ac-227+D	Ac-227 Th-227 [99%] Ra-233 Rn-219 Po-215 Pb-211 Bi-211 Tl-207	2.177E+01 Y 1.872E+01 D 1.143E+01 D 3.960E+00 S 1.778E-03 S 3.610E+01 M 2.130E+00 M 4.770E+00 M
	Pb-207	Pb-207	[Stable]
Thorium-232	Th-232	Th-232	1.405E+10 Y
	Ra-228+D	Ra-228 Ac-228	5.750E+00 Y 6.130E+00 H
	Th-228+D	Th-228 Ra-224 Rn-220 Po-216 Pb-212 Bi-212 Po-212 [64%] T1-208 [36%]	1.913E+00 Y 3.620E+00 D 5.561E+01 S 1.460E-01 S 1.064E+01 H 6.055E+01 M 2.980E-07 S 3.053E+00 M
	Pb-208	Pb-208	[Stable]

Exhibit 1: Radioactive Decay Chains Included in HEAST Tables 4A and 4B

(Continued) \*

Principal Decay Chain	Subchain*	Members <sup>b</sup>	Half-life <sup>c</sup>
Neptunium-237	Np-237+D	Np-237 Pa-233	2.140E+06 Y 2.700E+01 D
	บ-233	U-233	1.592E+05 Y
	Th-229+D	Th-229 Ra-225 Ac-225 Fr-221 At-217 Bi-213 Po-213 [98%] Tl-209 [2%] Pd-209	7.340E+03 Y 1.480E+01 D 1.000E+01 D 4.800E+00 M 3.230E-02 S 4.565E+01 M 4.200E-06 S 2.200E+00 M 3.253E+00 H
	Bi-209	Bi-209	[Stable]
Americium-243	Am-243+D	Am-243 Np-239	7.380E+03 Y 2.355E+00 D
Cesium-137	Cs-137+D	Cs-137 Ba-137m	3.017E+01 Y 2.552E+00 M
Strontium-90	Sr-90+D	Sr-90 Y-90	2.860E+01 Y 6.410E+01 H

<sup>\*</sup> See the discussion on radioactive decay chains in the User's Guide.

<sup>\*</sup> Radioactive decay chains included in HEAST Tables 4A and 4B. Radionuclides marked with the suffix "+D" include risks from decay chain members, assuming secular equilibrium (i.e., equal activity concentrations) in the environment.

b The chain of decay products of a parent radionuclide extends to (but does not include) members of the next subchain (e.g., U-238+D includes U-238, Th-234 and Pa-234, but not U-234). Note that there may be circumstances when it may be necessary to combine the risks for a parent radionuclide over several contiguous subchains, depending on the conditions of equilibrium. Branches in the decay chain are indicated in square brackets with branching percentages in parentheses.

Tradon-222 decay subchain, Rn-222+D, is also included in the HEAST tables, comprised of ingestion, inhalation and external exposure slope factors for Rn-222 plus the corresponding slope factors for each of its decay products (Po-218, Pb-214, Bi-214 and Po-214). For the ingestion and external exposure slope factors for Rn-222+D, decay products are assumed to be in secular equilibrium. For the inhalation slope factor, decay products are assumed to be in 50% equilibrium.

c Radioactive half-life in years (Y), days (D), hours (H), minutes
(M) or seconds (S).

# Exhibit 2. EPA Radiation Program Staff

(617) 565-4502 Tom D'Avanzo Radiation Program Manager, Region 1 U.S. Environmental Protection Agency John F. Kennedy Federal Building Room 2311 Boston, MA 02203 (212) 264-4110 Paul A. Giardina Radiation Branch Chief, Region 2 U.S. Environmental Protection Agency Room 1005 (AWM-RAD) 26 Federal Plaza New York, NY 10278 (215) 597-8326 Lewis K. Felleisen Radiation Program Manager, Region 3 Special Program Section (3AM12) U.S. Environmental Protection Agency 841 Chestnut Street Philadelphia, PA 19107 (404) 347-3907 Chuck Wakamo Radiation Program Manager, Region 4 U.S. Environmental Protection Agency 345 Courtland Street, NE Atlanta, GA 30365 (312) 886-6175 Jack Barnette Radiation Program Manager, Region 5 U.S. Environmental Protection Agency 230 S. Dearborn Street Chicago, IL 60604 Donna M. Ascenzi (214) 655-7224 Radiation Program Manger, Region 6 U.S. Environmental Protection Agency Air Enforcement Branch (6T-E) Air, Pesticides and Toxics Division 1445 Ross Avenue Dallas, TX 75202-2733 Robert Dye (913) 551-7605 Radiation Program Manager, Region 7

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Office of Radiation and Indoor Air U.S. Environmental Protection Agency 1504 Avenue A Montgomery, AL 36115-2601 Jed Harrison, Acting Director (702) 798-2476 Office of Radiation and Indoor Air Las Vegas Facility (ORP/LVF)
U.S. Environmental Protection Agency P.O. Box 98517 Las Vegas, NV 89193-8517 (202) 233-9350 Nicholas Lailas, Chief Office of Radiation and Indoor Air Radiation Assessment Branch Radiation Studies Division (6603-J) U.S. Environmental Protection Agency

401 M Street, SW Washington, DC 20460

# Exhibit 2. EPA Radiation Program Staff (Concluded)

Anthony B. Wolbarst, Chief (202) 233-9392 Office of Radiation and Indoor Air Remedial Guidance Section Radiation Assessment Branch Radiation Studies Division (6603-J) U.S. Environmental Protection Agency 401 M Street, SW Washington, DC 20460

March 1994

	<u>ECIES</u> Ent length	TARGET C	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
ACENAPHTHENE NOAEL 175 MG/KG/DAY ORAL: GAVAGE	OOOO83- MOUSE 90 DAYS	-32-9 LIVER	HEPATOTOXICITY		6E-1 300	IRIS	010165
ACENAPTHYLENE GENERAL COMMENT:	OOO208 DATA INADEQUAT	<del>-</del>	/E RISK ASSESSMENT				005202
ACEPHATE LOAEL 2 PPM ORAL: DIET  GENERAL COMMENT:	O3O56O-19 RAT 13 WEEKS ALSO SEE HEAST	-1 BRAIN TABLE 3: CARCING	DECREASED CHOLINESTERASE ACTIVITY DGENICITY.		4E-3 30	IRIS	005833
ACETONE  NOEL 100 MG/KG/DAY  ORAL: GAVAGE	000067-64- RAT 90 DAYS	1 LIVER KIDNEY KIDNEY	INCREASED WEIGHT INCREASED WEIGHT NEPHROTOXICITY		1E+0 100	IRIS	005204
ACETONE CYANOHYDRIN  NOEL 10.8 MG CN/KG/DAY  ORAL: DIET	I 000 RAT 104 WEEKS	O75-86-5  WHOLE BODY THYROID CENTRAL NERVOUS SYSTEM	ALTERED WEIGHT EFFECTS EFFECTS		7E-2 500	7E-2 500	005776

SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

Subchronic Chronic CHEMICAL SPECIES [RfC] [RfC] [RfD] REFERENCE [RfD] ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF ACETONITRILE 000075-05-8 NOAEL 100 PPM MOUSE INHALATION: 92 DAYS **ERYTHROCYTES** 6E-2 IRIS 005210 DECREASED CELL COUNT INTERMITTENT BLOOD DECREASED HEMATOCRIT 300 LIVER **HEPATIC LESIONS** SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) SUBCHRONIC (RfD) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.5. CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.5. UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE. **ACETOPHENONE** 000098-86-2 NOAEL 10,000 PPM RAT 005212 IRIS ORAL: DIET 17 WEEKS NONE OBSERVED 1E+0 300 010874 CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (06/25/92) BY THE RfD/RfC WORK GROUP. **ACROLEIN** 000107-02-8 NOAEL 15.6 MG/KG/DAY RAT 2E-2 010390 ORAL: WATER 90 DAYS 1000 GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. 010856 IRIS SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. **ACRYLAMIDE** 000079-06-1 NOEL 0.2 MG/KG/DAY RAT 005835 ORAL: DRINKING 90 DAYS NERVE DAMAGE 2E-3 IRIS 100 WATER GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. 010876 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (09/20/90) BY THE RFD/RFC WORK GROUP.

March 1994

CHEMICAL LEVEL	<u>Dose</u> Route e)	<u>SPECIES</u> (PERIMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC] [R· (mg/cu m) (mg/k	fD] [RfC]	onic [RfD] (mg/kg/day) UF	REFERENCE
ACRYLIC	ACID	000079	-10-7					
NOAEL	. 83 MG/KG/DAY ORAL: DRIN WATER		WHOLE BODY ORGANS, MAJOR	DECREASED WEIGHT ALTERED WEIGHT		BE-1 100	IRIS	005836
LOAEL	5 PPM INHALATION INTERMITTE		NASAL MUCOSA	LESIONS	3E-3 100	IRIS		010346
ACRYLO	NITRILE	000107	7-13-1					
NOAEL	. 1 MG/(KG-DAY ORAL: GAVA		TESTES TESTES	DECREASED SPERM COUNTS SEMINIFEROUS TUBULE DEGENERATION		IE-2 100	1E-3 1000	010939
			E CHRONIC [RfD] UNI AST TABLE 3: CARCI	DER REVIEW BY THE RFD/RFC WON NOGENICITY	RK GROUP WAS MODIFIE	ED TO DERIVE THE S	UBCHRONIC [R	fD].
ADIPONI	TRILE	000111-	69-3					
	GENERAL COM			IVE RISK ASSESSMENT				005157
ALACHLO NOEL	OR 1 MG/KG/DAY	015972- pog	60-8					

1E-2

100

IRIS

005837

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

SITES, MULTIPLE HEMOSIDEROSIS

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

BLOOD

1 YEAR

ORAL: CAPSULE

ANEMIA

Subchronic Chronic CHEMICAL SPECIES DOSE [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

**ALDICARB** 000116-06-3

> NOAEL 0.01 MG/KG-DAY HUMAN

ORAL ACUTE CENTRAL NERVOUS SWEATING 1E-3 010960 IRIS 10

SYSTEM

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

GENERAL COMMENT: CLINICAL SIGNS OF ACETYL CHOLINESTERASE INHIBITION INCLUDING SWEATING, PINPOINT PUPILS, LEG WEAKNESS, NAUSEA, DIARRHEA

AND OTHER EFFECTS WERE OBSERVED IN THE PRINCIPAL AND SUPPORTING STUDIES.

ALDRIN 000309-00-2

> LOAEL 0.025 MG/KG/DAY RAT

ORAL: DIET 2 YEARS 3E-5 IRIS 005159 LIVER LESIONS 1000

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

ALLIDOCHLOR 000093-71-0

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.

ALLYL ALCOHOL 000107-18-6

NOEL 50 PPM RAT

ORAL: DRINKING 15 WEEKS LIVER **EFFECTS** 5E-2 IRIS 005839 100 WATER KIDNEY **EFFECTS** 

ALLYL CHLORIDE 000107-05-1

NOAEL 17 MG/CU M RABBIT

> INHALATION: 5 MONTHS **NERVOUS SYSTEM** NEUROTOXICITY 1E-2 IRIS 010369 300 INTERMITTENT

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

**ALUMINUM** 007429-90-5

> GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005162

March 1994

005838

<u>CHEMICAL</u> LEVEL		<u>SPECIES</u> IMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
ALUMINU	IM PHOSPHIDE	020	859-73-8				
NOAEL	0.43 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	WHOLE BODY UNSPECIFIED	ALTERED WEIGHT ALTERED CLINICAL PARAMETERS	4E-4 100	IRIS	010255
AMETRY	N	000834-12	2-8				
NOEL	10 MG/KG/DAY ORAL: GAVAGE	RAT 13 WEEKS	LIVER	EFFECTS	<del>9</del> E-2 100	IRIS	005841
AMINO-2	-NAPHTHOL, 1 GENERAL COMMEN		34-92-6 ATE FOR QUANTITA	TIVE RISK ASSESSMENT.			005842
AMINO-2	-NAPHTOL HYI GENERAL COMMEN		-	8-27-2 ITIVE RISK ASSESSMENT.			005843
AMINOPH	IENOL, M-	00059	1-27-5				
	1300 PPM ORAL: DIET	RAT 13 WEEKS	WHOLE BODY THYROID	ALTERED WEIGHT ALTERED WEIGHT	7E-1 100	7E-2 1000	005844
AMINOPH	•	OOOO95		TIVE RISK ASSESSMENT.			005845
AMINOPH	- •	OOO123		TIVE RISK ASSESSMENT.			005846
AMINOPY	'RIDINE, 4-	000504	24-5				
NOAEL	•	RAT 90 DAYS	LIVER BRAIN	INCREASED WEIGHT INCREASED WEIGHT	2E-4 1000	2E-5 10000	005847

Subchronic Chronic DOSE ROUTE CHEMICAL **SPECIES** [RfC] [RfD] REFERENCE [RfC] [RfD] LEVEL EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF **AMMONIA** 007664-41-7 NOAEL 34 MG/L HUMAN ORAL: DRINKING SENSORY TASTE THRESHOLD 34 MG/L 34 MG/L 005166 WATER SUBCHRONIC [RfD] COMMENT: GIVEN AS CONCENTRATION IN DRINKING WATER. SPECIFICALLY RELATED TO ORGANOLEPTIC THRESHOLD. SAFE CONCENTRATION MAY BE HIGHER, BUT DATA ARE INADEQUATE TO ASSESS. CHRONIC [RfD] COMMENT: GIVEN AS CONCENTRATION IN DRINKING WATER. SPECIFICALLY RELATED TO ORGANOLEPTIC THRESHOLD. SAFE CONCENTRATION MAY BE HIGHER, BUT DATA ARE INADEQUATE TO ASSESS. NOAEL 6.4 MG/CU M HUMAN INHALATION: 010392 NASAL CAVITY RHINITIS 1E-1 IRIS INTERMITTENT 30 LUNGS **PNEUMONIA** LUNGS LESIONS **ANILINE** 000062-53-3 NOAEL 19 MG/CU M MOUSE INHALATION: 20-26 WEEKS 010370 SPLEEN **PATHOLOGY** 1E-2 IRIS INTERMITTENT 300 RAT 20-26 WEEKS SPLEEN **PATHOLOGY** GUINEA PIG 20-26 WEEKS SPLEEN **PATHOLOGY** GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. **ANTHRACENE** 000120-12-7 NOEL 1000 MG/KG/DAY MOUSE ORAL: GAVAGE IRIS 010166 90 DAYS NONE OBSERVED 3E+0 300 **ANTIMONY PENTOXIDE** 001314-60-9 LOAEL 0.46 MG/KG/DAY RAT ORAL: DRINKING 5E-4 005174 LIFETIME WHOLE BODY INCREASED MORTALITY 5E-4 WATER ALTERED CHEMISTRIES 1000 1000 BLOOD

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

Subchronic Chronic **SPECIES** [RfC] [RfC] [RfD] REFERENCE CHEMICAL DOSE [RfD] EXPERIMENT LENGTH LEVEL ROUTE TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF UF UF ANTIMONY POTASSIUM TARTRATE 000304-61-0 LOAEL 0.91 MG/KG/DAY LIFETIME WHOLE BODY 9E-4 9E-4 005234 ORAL: DRINKING INCREASED MORTALITY WATER BLOOD ALTERED CHEMISTRIES 1000 1000 SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. **ANTIMONY TETROXIDE** 001332-81-6 RAT LOAEL 0.44 MG/KG/DAY 005238 ORAL: DRINKING LIFETIME WHOLE BODY INCREASED MORTALITY 4E-4 4E-4 WATER BLOOD ALTERED CHEMISTRIES 1000 1000 SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. ANTIMONY TRIOXIDE 001309-64-4 LOAEL 0.42 MG/KG/DAY RAT 005242 LIFETIME WHOLE BODY **INCREASED MORTALITY** 4E-4 4E-4 ORAL: DRINKING 1000 1000 WATER BLOOD ALTERED CHEMISTRIES SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ANTIMONY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. ANTIMONY, METALLIC 007440-36-0 LOAEL 0.35 MG SB/KG/DAY RAT 005170 IRIS ORAL: DRINKING LIFETIME WHOLE BODY INCREASED MORTALITY 4E-4 1000 ALTERED CHEMISTRIES WATER BLOOD

CHEMICAL LEVEL		<u>ecies</u> Ent length	TARGET	CRITICAL EFFECT	[RfC] (mg/cu m) ( Uf	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/der UF UF	REFERENCE
ARAMITE	:	000140-57-	R					
	100 PPM ORAL: DIET	RAT 104 WEEKS	LIVER	INCREASED WEIGHT			5E-2 100	005850
NOAEL	500 PPM ORAL: DIET	DOG 52 WEEKS	LIVER	DEGENERATION		1E-1 100		005849
	GENERAL COMMENT:	ALSO SEE HEAS	TABLE 3: CARCI	NOGENICITY.				
AROCHLO	= -	012672 MMENT: THE CHR	·	IS CONSIDERED NOT VERIFIABLE	E (07/20/93) BY	THE RfD/RfC	WORK GROUP.	010940
	, INORGANIC 0.009 mg/l oral	007440 Human	O-38-2 SKIN SKIN	KERATOSIS HYPERPIGMENTATION		3E-4 3	IRIS	010434
	SUBCHRONIC [RfD] GENERAL COMMENT:			WAS ADOPTED AS THE SUBCHRONOGENICITY.	NIC ORAL [RfD].			
ATRAZIN	E	001912-24	9					
NOEL	3.5 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		3.5E-2 100	IRIS	010855
	SUBCHRONIC [RfD] GENERAL COMMENT:			WAS ADOPTED AS THE SUBCHRONOGENICITY.	NIC ORAL [RfD].			
BARIUM		007440-39-	3					
	0.21 MG/KG/DAY ORAL: WATER	HUMAN 10 WEEKS	CARDIOVASCULAR SYSTEM	INCREASED BLOOD PRESSURE		7E-2 3	IRIS	010348
				2: ALTERNATE METHODS SU WAS ADOPTED AS THE SUBCHRO	NIC ORAL [RfD].		TY (OTHER THAN CARCINOC	

CHRONIC [RFC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

March 1994

Subchronic Chronic

UF

UF

CHEMICAL SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

BARIUM CYANIDE 000542-62-1

> CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/20/93) BY THE RfD/RfC WORK GROUP. 010941

UF

**BENEFIN** 001861-40-1

> NOAEL 25 MG/KG/DAY DOG

1 YEAR 3E-1 IRIS 005852 ORAL: DIET **ERYTHROCYTE** DECREASED COUNT 100

SUBCHRONIC (RfD) COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

BENZAL CHLORIDE 000098-87-3

> 005853 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.

BENZALDEHYDE 000100-52-7

> NOEL 200 MG/KG/DAY RAT

1E+0 IRIS 005854 ORAL: GAVAGE 13 WEEKS KIDNEY **EFFECTS** 100

LESIONS FORESTOMACH

SUBCHRONIC (RfD) CPMMENT: THE CHRONIC ORAL RFD WAS MODIFIED TO DERIVE THE SUBCHRONIC ORAL [RfD].

BENZALDEHYDE CYANOHYDRIN 000532-28-5

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005781

BENZENE 000071-43-2

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

000108-98-5 BENZENETHIOL / (THIOPHENOL)

LOAEL 0.1 MG/(KG-DAY)

1E-5 010942 1E-4 ORAL: GAVAGE 90 DAYS LIVER CENTRILOBULAR EOSINOPHILIC

1000 10.000 CHANGES

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] UNDER REVIEW BY THE RFD/RFC WORKGROUP WAS MODIFIED TO DERIVE THE SUBCHRONIC [RfD].

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Subchronic Chronic CHEMICAL SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF

BENZIDINE 000092-87-5

> LOAEL 2.7 MG/KG/DAY MOUSE

ORAL: DRINKING 33 MONTHS 3E-3 IRIS 005830 BRAIN **CELLULAR CHANGES** WATER LIVER 1000 **CELLULAR CHANGES** 

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

010877 IRIS

CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (03/28/91) BY THE RFD/RFC WORK GROUP.

BENZOIC ACID 000065-85-0

> NOAEL 312 MG/DAY HUMAN

> > 005260 ORAL: DIET IRIS NONE OBSERVED 4E+0

SUBCHRONIC [RfD] COMMENT: THE HUMAN DAILY PER CAPITA INTAKE WAS USED AS THE CRITICAL DOSE LEVEL. THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfD] COMMENT: THE HUMAN DAILY PER CAPITA INTAKE WAS USED AS THE CRITICAL DOSE LEVEL.

**BENZO[B]FLUORANTHENE** 000205-99-2

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

BENZYL ALCOHOL 000100-51-6

LOAEL 286 MG/KG/DAY

RAT ORAL: GAVAGE

005855 103 WEEKS **FORESTOMACH** EPITHELIAL HYPERPLASIA 3E-1 1000

NOAEL 143 MG/KG/DAY

RAT

1E+0 005856 ORAL: GAVAGE 13 WEEKS WHOLE BODY DECREASED WEIGHT

100

<u>CHEMICAL</u> <u>DO</u> LEVEL RO		<u>cies</u> Nt length	TARGET (	CRITICAL EFFECT	Subchro [RfC] [RfD (mg/cu m) (mg/kg/ UF UF	] [RfC] /day) (mg/cu_m) (mg	[RfD] R	EFERENCE
BERYLLIUM		007440-41-	7					
NOAEL 0.54	MG/KG/DAY RAL: DRINKING ATER	RAT LIFETIME		NONE OBSERVED	5E 10	:-3 00	IRIS	005262
			HRONIC ORAL RfD TABLE 3: CARCIN	WAS ADOPTED AS THE SUBCHRONIC OGENICITY.	ORAL [RfD].			
BIPHENYL, 1	,1′	000092-52-	4					
NOAEL 50 P	NG/KG/DAY RAL: DIET	RAT 700 DAYS	KIDNEY	DAMAGE		:-2 000	IRIS	005857
su	BCHRONIC [RfD] (	COMMENT: THE C	HRONIC ORAL RfD	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].	IRIS		010878
CH	RONIC RFC COMMEN	IT: THE CHRONIC	C INHALATION RFC	IS CONSIDERED NOT VERIFIABLE	(09/20/90) BY THE	RfD/RfC WORK GROUP.		
BIS(2-CHLOR	OISOPROPYL	ETHER 03	9638-32-9					
NOAEL 35.8	MG/KG/DAY	MOUSE		RECREAGES HEMOSI ORIN	45	:-2	IRIS	010257
0	RAL: DIET	2 YEARS	ERYTHROCYTES	DECREASED HEMOGLOBIN		000	IKIS	010237
GE	NERAL COMMENT:	ALSO SEE HEAST	TABLE 3: CARCIN	OGENICITY.				
BIS(2-ETHYL	HEXYL) PHTH	ALATE / (DEH	IP) 000117-	81-7				
•							IRIS	010859
SU	BCHRONIC [RfD]	COMMENT: CONTAC	CT THE SUPERFUND CT THE SUPERFUND TABLE 3: CARCIN	HEALTH RISK TECHNICAL SUPPORT HEALTH RISK TECHNICAL SUPPORT OGENICITY.	CENTER: (513) 569 CENTER: (513) 569	?-7300. ?-7300.		
BISPHENOL A	A	000080-05	-7					
NOAEL 750	PDM	RAT					IRIS	005268
	RAL	13 WEEKS	WHOLE BODY	DECREASED WEIGHT	6E 10	E-1 00		005266

CHEMICAL LEVEL	<u>pose</u> Route exp	<u>SPECIES</u> Eriment length	TARGET	CRITICAL EFFECT	Subchroni [RfC] [RfD] <u>(mg/cu m) (mg/kg/da</u> UF UF	[RfC] [RfD]	REFERENCE 2			
BORON T	RIFLUORIDE	00763	7-07-2							
NOAEL	6 MG/CU M INHALATION: INTERMITTENT	rat 13 weeks	KIDNEY	NECROSIS	7E-3 300	7E-4 3000	010395			
BORON, ELEMENTAL 007440-42-8										
NOAEL	8.8 MG/KG/DAY ORAL: DIET	DOG 2 YEARS	TESTIS	LESIONS	9E-2 100	IRIS	005272			
	SUBCHRONIC [	RFD) COMMENT: THE	CHRONIC ORAL Rf	D WAS ADOPTED AS THE SUBC	HRONIC ORAL [RfD].					
LOAEL	4.5 MG/CU M INHALATION: INTERMITTENT	HUMAN	RESPIRATORY TRACT BRONCHUS	IRRITATION BRONCHITIS	2E-2 100	2E-2 100	00526 <del>9</del>			
	SUBCHRONIC [F CHRONIC [RfC]	RFC] COMMENT: THE CHR	SUBCHRONIC INHA	LATION [RfC] IS SPECIFICAL [RfC] IS SPECIFICALLY FO	LLY FOR ANHYDROUS BORAX. R ANHYDROUS BORAX.					
BROMINA	ATED DIBENZ GENERAL COMME		TE FOR QUANTITA	TIVE RISK ASSESSMENT.			005858			
BROMINA	ATED DIBENZ	OFURANS								
	GENERAL COMME	ENT: DATA INADEQUA	TE FOR QUANTITA	TIVE RISK ASSESSMENT.			005859			
BROMOA	CETONE GENERAL COMME	OOO59 ENT: DATA INADEQUA		TIVE RISK ASSESSMENT.			005860			
вкомос	HLOROETHA GENERAL COMME	<del>-</del>	TE FOR QUANTITA	TIVE RISK ASSESSMENT.			005861			

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Subchronic Chronic CHEMICAL DOSE ROUTE **SPECIES** [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF

BROMODICHLOROMETHANE 000075-27-4

LOAEL 17.9 MG/KG/DAY MOUSE

ORAL: GAVAGE 102 WEEKS KIDNEY CYTOMEGALY 2E-2 IRIS 005715

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

BROMOETHENE / (VINYL BROMIDE) 000593-60-2

LOAEL 9.7 PPM RAT

INHALATION: 24 MONTHS LIVER HYPERTROPHY 3E-3 IRIS 010929
INTERMITTENT LIVER BASOPHILIC FOCI 3000
LIVER EOSINOPHILIC FOCI

SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RFC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

BROMOFORM 000075-25-2

NOEL 17.9 MG/KG/DAY RAT

ORAL: GAVAGE 13 WEEKS LIVER EFFECTS 2E-1 IRIS 005722

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

IRIS 010961
CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (02/11/93) BY THE RFD/RFC WORK GROUP.

BROMOMETHANE 000074-83-9

IRIS 010861 IRIS 010860

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

BROMOPHENYL PHENYL ETHER, 4- 000101-55-3

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005864

<u>CHEMICAL</u> LEVEL	<u>DOSE</u> ROUTE EXPER	<u>SPECIES</u> IMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC] [Rf (mg/cu m) (mg/k	fD] [RfC]	ronic [RfD] <u>(mg/kg/day)</u> UF	REFERENCE
BROMOPI	ноѕ	002104-9	96-3					
NOAEL	5 MG/KG/DAY ORAL: DIET	RAT 3	BLOOD	DECREASED CHOLINESTERASE	•	5E-2	5E-3	005865
	ORAL: DIE	GENERATIONS		ACTIVITY		100	1000	003003
			LIVER	DECREASED CHOLINESTERASE ACTIVITY				
		D] COMMENT: BASE COMMENT: BASED O						
BROMOX	YNIL	001689-8	34-5					
NOEL	5 MG/KG/DAY ORAL: DIET	RAT 2 YEARS		NONE OBSERVED	;	2E-2	IRIS	005866
	SUBCHRONIC [R1	D] COMMENT: THE	CHRONIC ORAL RFD	WAS ADOPTED AS THE SUBCHRONIC		300		
BROMOY	YNIL OCTANO	ATE OO	1689-99-2					
	7.3 MG/KG/DAY	RAT	1003-33-2					
	ORAL: DIET	2 YEARS		NONE OBSERVED		2E-2 300	IRIS	005867
	SUBCHRONIC [R1	D] COMMENT: THE	CHRONIC ORAL Rf	WAS ADOPTED AS THE SUBCHRONIC				
BUSAN 7	7	031512-74	-0					
	GENERAL COMMEN	T: DATA INADEQUA	TE FOR QUANTITAT	TIVE RISK ASSESSMENT.				005868
BUSAN 9	0	002491-38	-5					
	GENERAL COMMEN	IT: DATA INADEQUA	TE FOR QUANTITA	TIVE RISK ASSESSMENT.				005869
BUTANOI	L, 1-	000071-36	<b>6-3</b>					
NOAEL	125 MG/KG/DAY	RAT	CENTRAL NERVO	IC HYPOLOTINITY		1E+0	IRIS	005870
	ORAL: GAVAGE	13 WEEKS	CENTRAL NERVO			100	1419	007010
			CENTRAL NERVO	US ATAXIA				

Subchronic Chronic CHEMICAL DOSE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF **BUTYL BENZYL PHTHALATE, N-**000085-68-7 NOAEL 159 MG/KG/DAY ORAL: DIET 26 WEEKS 2E+0 LIVER ALTERED WEIGHT IRIS 005616 100 GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. BUTYLATE 002008-41-5 NOEL 5 MG/KG/DAY DOG ORAL: CAPSULE 12 MONTHS LIVER INCREASED RELATIVE WEIGHT 5E-2 IRIS 005871 100 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. BUTYLCHLORIDE, T-000507-20-0 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005810 **BUTYROLACTONE, GAMMA-**000096-48-0 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005872 CACODYLIC ACID 000075-60-5 NOEL 9.2 MG/KG/DAY RAT ORAL: DIET 90 DAYS NONE OBSERVED 3E-2 3E-3 005873 3000 300

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ARSENIC BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.
CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO ARSENIC BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

CADMIUM 007440-43-9

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. IRIS 005280

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route expe	<u>SPECIES</u> Eriment Length	TARGET	CRITICAL EFFECT	Subchronic Chro [RfC] [RfD] [RfC] (mg/cu m) (mg/kg/day) (mg/cu m) ( UF UF UF	[RfD]	REFERENCE
CALCIUN	/ CYANIDE	00059	2-01-8				
	19.1 MG/KG/DAY		_ 0 . 0				
	ORAL: DIET	2 YEARS	WHOLE BODY THYROID NERVE	DECREASED WEIGHT EFFECTS MYELIN DEGENERATION	4E-2 500	IRIS	010258
	_	WAS ADOPTED	AS THE SUBCHRON	IC ORAL [RfD].	NG FOR DIFFERENCES IN MOLECULAR WEIGH	T. THE CHRO	NIC ORAL RFD
	CHRONIC [RTD]	COMMENT: CALCUL	ALED BY ANALOGY	TO FREE CYANIDE BY CORRECTING	FOR DIFFERENCES IN MOLECULAR WEIGHT.		
CAPROL	ACTAM	00010	5-60-2			IRIS	005284
NOAEL	50 MG/KG/DAY	RAT				IKIS	
	ORAL: DIET	90 DAYS	KIDNEY	EFFECTS	5E-1 100		005282
CAPTAF	OL	002425-0	6-1				
LOAEL	2 MG/KG/DAY ORAL: CAPSUL	DOG E 12 MONTHS	KIDNEY BLADDER	EFFECTS EFFECTS	2E-3 1000	IRIS	005874
	SUBCHRONIC [R GENERAL COMME	fd] comment: the nt: ALSO SEE HEA		D WAS ADOPTED AS THE SUBCHRONI INOGENICITY.	C ORAL [RfD].		
CAPTAN		000133-06	<b>-2</b>				
NOEL	12.5 MG/KG/DAY ORAL: DIET	RAT	WHOLE BODY	DECREASED WEIGHT	1.3E-1 100	IRIS	005875

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (05/27/92) BY THE RFD/RFC WORK GROUP.

CHRONIC [RfD] COMMENT: BASED ON A MULTI-GENERATION REPRODUCTION STUDY.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

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Subchronic Chronic REFERENCE [RfC] [RfC] [RfD] CHEMICAL <u>DOSE</u> ROUTE SPECIES [RfD] LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF HF

**CARBARYL** 000063-25-2

> NOAEL 9.6 MG/KG/DAY RAT

005876 1E-1 IRIS 2 YEARS TOXICITY ORAL: DIET KIDNEY 100 LIVER TOXICITY

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

IRIS 010882

CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (08/15/91) BY THE RFD/RFC WORK GROUP.

001563-66-2 CARBOFURAN

> NOEL 0.5 MG/KG/DAY DOG

CHOLINESTERASE INHIBITION 005877 5E-3 IRIS 1 YEAR **BLOOD** ORAL: DIET

100 **TESTIS EFFECTS** 

**EFFECTS UTERUS** 

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

000075-15-0 CARBON DISULFIDE

> NOEL 11 MG/KG/DAY RABBIT

1E-1 IRIS 010259 **FETUS** TOXICITY INHALATION:

100 INTERMITTENT

SUBCHRONIC (RfD) COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL (RfD).

CHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS DETERMINED FROM A TERATOLOGY STUDY WITH EXPOSURES BEFORE AND DURING THE ENTIRE GESTATION

PERIOD.

NOAEL 10 MG/CU M

RAT 010430 1E-2 1E-2 **FETUS** TOXICITY INHALATION: GESTATION

1000 1000 INTERMITTENT

SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].

000630-05-0 CARBON MONOXIDE

010493 CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS.

Subchronic Chronic CHEMICAL SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF

**CARBON TETRACHLORIDE** 000056-23-5

> IRIS 010862

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

**CHLORAL** 000075-87-6

> LOAEL 15.7 MG/KG/DAY MOUSE

ORAL: DRINKING 90 DAYS LIVER **EFFECTS** 2E-2 IRIS 005290 WATER 1000

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHLORDANE 000057-74-9

NOEL 0.055 MG/KG/DAY RAT

ORAL: DIET 130 WEEKS LIVER **HYPERTROPHY** 6E-5 IRIS 005296 1000

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

CHLORINE CYANIDE 000506-77-4

> NOAEL 25.3 MG/KG/DAY RAT

> > ORAL: DIET 2 YEARS WHOLE BODY 010261 DECREASED WEIGHT 5E-2 IRIS 500

THYROID **EFFECTS** 

NERVE MYELIN DEGENERATION

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHEMICAL LEVEL	<u>pose</u> Route exp	<u>SPECIES</u> Eriment Length	TARGET	CRITICAL EFFECT	[RfC]	ubchronic [RfD] mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
CHLORO	-1,3-BUTADIE	NE, 2- / (CHLO	ROPRENE) OO	0126-99-8				
NOAEL	32 PPM INHALATION	RAT 90 days	OLFACTORY EPITHELIUM	DEGENERATION	7E-2 30		7E-3 300	010515
	CHRONIC [RfD]	COMMENT: ALSO SE	E HEAST TABLE 2	E 2: ALTERNATE METHODS P: ALTERNATE METHODS SU RNATE METHODS SUBCHRON	BCHRONIC AND CHRONIC	C TOXICITY (	OTHER THAN CARCINOGEN	ICITY)
CHLORO	-M-CRESOL, I	P- 0000	59-50-7					
NOAEL	200 MG/KG/DAY ORAL: GAVAG	RAT 28 Days	WHOLE BODY	DECREASED WEIGHT GAIN		2E+0 100		005366
CHLORO	ACETALDEH		0107-20-0					005342
	GENERAL COMMI	INT: DATA INADEQUA	TE FOR QUANTITA	TIVE RISK ASSESSMENT.				005342
CHLORO	ACETIC ACID	0000	79-11-8					
LOAEL	30 MG/KG ORAL: GAVAG	RAT 13 WEEKS	HEART	MYOCARDITIS		2E-2 1000	2E-3 10000	005346
CHLORO	ANILINE, 2- GENERAL COMM	OOOO95 ENT: DATA INADEQUA		ATIVE RISK ASSESSMENT				005347
CHLORO	ANILINE, 3- GENERAL COMM	OOO108 ENT: DATA INADEQUA	- <del>-</del>	ATIVE RISK ASSESSMENT				005348
	ANILINE, 4- 12.5 MG/KG/DAY ORAL: DIET	000106 RAT 78 WEEKS	S-47-8 SPLEEN	PROLIFERATIVE LESIONS	:	4E-3 3000	IRIS	005349

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Subchronic Chronic <u>Dose</u> Route CHEMICAL SPECIES [RfC] [RfC] [RfD] REFERENCE [RfD] LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

CHLOROBENZENE 000108-90-7

> IRIS 010863

UF

UF

ÜF

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**CHLOROBENZILATE** 000510-15-6

> NOEL 5 MG/KG/DAY RABBIT

010260 13 DAYS 2E-2 IRIS ORAL: GAVAGE GASTRO-DECREASED STOOL QUANTITY 300

INTESTINAL SYSTEM WHOLE BODY DECREASED FOOD CONSUMPTION

WHOLE BODY DECREASED WEIGHT GAIN NERVOUS SYSTEM HYPERIRRITABILITY

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. BASED ON A TERATOLOGY STUDY WITH EXPOSURES DURING DAYS 7-19 OF GESTATION.

CHRONIC [RfD] COMMENT: BASED ON A TERATOLOGY STUDY WITH EXPOSURES DURING DAYS 7-19 OF GESTATION.

010931

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (02/11/93) BY THE RfD/RfC WORK GROUP.

CHLOROBENZOIC ACID, P-000074-11-3

NOAEL 26 MG/DAY RAT

5 MONTHS

ORAL: DIET NONE OBSERVED 2E+0 2E-1 005360 1000 100

CHLOROBENZOTRIFLUORIDE, 4-000098-56-6

NOAEL 15 MG/KG/DAY RAT

005364 ORAL: GAVAGE KIDNEY TUBULAR DEGENERATION 2E-1 2E-2 100 1000

SUBCHRONIC (RFD) COMMENT: BASED ON A 2-GENERATION REPRODUCTION STUDY WITH EXPOSURES 4 WEEKS PRIOR TO MATING, DURING GESTATION, AND FOR 90 DAYS POST-WEANING.

CHRONIC [RfD] COMMENT: BASED ON A 2-GENERATION REPRODUCTION STUDY WITH EXPOSURES 4 WEEKS PRIOR TO MATING, DURING GESTATION, AND FOR 90 DAYS POST-WEANING.

Subchronic Chronic REFERENCE CHEMICAL SPECIES [RfC] [RfD] [RfC] [RfD1 ROUTE LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF 000109-69-3 CHLOROBUTANE, 1-NOAEL 43 MG/KG/DAY RAT 4E-1 005808 103 WEEKS INCREASED MORTALITY ORAL: GAVAGE WHOLE BODY CENTRAL NERVOUS EFFECTS 100 SYSTEM **HEMATOLOGIC EFFECTS** BLOOD NOAEL 86 MG/KG/DAY RAT 005806 13 WEEKS WHOLE BODY DECREASED WEIGHT GAIN 9E-1 ORAL: GAVAGE 100 CENTRAL NERVOUS EFFECTS SYSTEM **HEMATOPOIESIS SPLEEN** CHLOROBUTANE, 2-000078-86-4 005809 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. CHLOROCYCLOPENTADIENE 041851-50-7 005297 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT CHLOROFORM 000067-66-3 LOAEL 12.9 MG/KG/DAY DOG 005372 1E-2 IRIS 7.5 YEARS LESIONS ORAL: CAPSULE LIVER 1000 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. CHLOROMETHANE / (METHYL CHLORIDE) 000074-87-3 010005 SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

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GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

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Subchronic Chronic CHEMICAL <u>Dose</u> Route SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

CHLORONITROBENZENE, M- 000121-73-3

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.

005879

CHLOROPHENOL, 2- 000095-57-8

NOAEL 50 PPM RAT

ORAL: DRINKING REPRODUCTION REPRODUCTIVE EFFECTS 5E-2 IRIS 010436 WATER

SUBCHRONIC [RfD] COMMENT: BASED ON A REPRODUCTIVE STUDY WITH EXPOSURE 10 WEEKS PRIOR TO AND DURING MATING, GESTATION AND WEANING. CHRONIC [RfD] COMMENT: BASED ON A REPRODUCTIVE STUDY WITH EXPOSURE 10 WEEKS PRIOR TO AND DURING MATING, GESTATION AND WEANING.

CHLOROPHENOL, 3- 000108-43-0

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005309

CHLOROPHENOL, 4- 000106-48-9

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005310

CHLOROPROPANE, 2- 000075-29-6

NOAEL 91.4 MG/KG/DAY RAT

INHALATION: 4 WEEKS LIVER EFFECTS 1E+0 1E-1 010444
INTERMITTENT 100 1000

CHLOROTOLUENE, M- 000108-41-8

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005880

CHLOROTOLUENE, O- 000095-49-8

NOAEL 20 MG/KG/DAY RAT

ORAL: GAVAGE 103 DAYS WHOLE BODY DECREASED WEIGHT GAIN 2E-1 IRIS 010167

CHLOROTOLUENE, P- 000106-43-4

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 010200

<u>CHEMICAL</u> LEVEL		<u>PECIES</u> Ment Length	TARGET	CRITICAL EFFECT		ochronic (RfD) g/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/cu UF UF	
CHLORPY	RIFOS	002921-	88-2					
NOEL	0.03 MG/KG/DAY ORAL: CAPSULE	HUMAN 20 days or 9 days	BL000	DECREASED CHOLINESTERASE		3E-3 10	IRIS	005881
	SUBCHRONIC [RfD]	COMMENT: THE	CHRONIC ORAL REC	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].			
	RIFOS METHYL		598-13-0					
NOAEL	1 MG/KG/DAY ORAL: DIET	RAT 3 GENERATIONS	REPRODUCTION	DECREASED FERTILITY		1E-2 100	1E-2 100	005882
		RAT 2 YEARS	LIVER	EFFECTS				
CHLOROT	ΓHALONIL	00189	7-45-6					
	1.5 MG/KG/DAY ORAL: DIET	DOG 2 YEARS	KIDNEY	EFFECTS		1.5E-2 100	IRIS	005883
	SUBCHRONIC [RfD] GENERAL COMMENT:			WAS ADOPTED AS THE SUBCHRONIC INOGENICITY.	ORAL [RfD].			
CHLORTH	HIOPHOS	060238	3-56-4					
	0.08 MG/KG/DAY ORAL: DIET	RAT 2 YEARS		NONE OBSERVED		8E-4 100	8E-4 100	005884
	SUBCHRONIC [RfD]	COMMENT: THE	CHRONIC ORAL [R	fd] was adopted as the subchron	IC ORAL [RfD].			
CHROMIL	JM(III)	016065-8	3-1					
NOEL	5 % (CR203) ORAL: DIET	RAT 840 days		NONE OBSERVED		1E+0 1000	IRIS	005731
	SUBCHRONIC [RfD]	COMMENT: THE	CHRONIC ORAL RE	NAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].			

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SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

CHRONIC [RfC] COMMENT: INHALATION ISSUES ARE UNDER REVIEW BY THE RFD/RFC WORK GROUP.

Subchronic Chronic CHEMICAL DOSE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF

CHROMIUM(VI) 018540-29-9

> NOAEL 2.4 MG/KG/DAY RAT

ORAL: DRINKING 1 YEAR 005522 NONE OBSERVED 2E-2 IRIS WATER 100

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

**CHRYSENE** 000218-01-9

> GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ALSO SEE HEAST TABLE 3: CARCINOGENICITY. 005885

COPPER 007440-50-8

> LOAEL 5.3 MG HUMAN

> > ORAL SINGLE DOSE GASTRO-IRRITATION 1.3 MG/L 1.3 MG/L 005374

INTESTINAL SYSTEM

CHRONIC [RfC] COMMENT: INHALATION ISSUES ARE UNDER REVIEW BY THE RFD/RFC WORK GROUP.

SUBCHRONIC [RfD] COMMENT: CURRENT DRINKING WATER STANDARD OF 1.3 MG/L. DWCD (1987) CONCLUDED TOXICITY DATA WERE INADEQUATE FOR

CALCULATION OF AN RfD FOR COPPER.

CHRONIC [RfD] COMMENT: CURRENT DRINKING WATER STANDARD OF 1.3 Mg/L. DWCD (1987) CONCLUDED TOXICITY DATA WERE INADEQUATE FOR CALCULATION

OF AN RfD FOR COPPER.

**COPPER CYANIDE** 000544-92-3

> NOAEL 5 MG/KG/DAY RAT

> > ORAL: GAVAGE 90 DAYS LIVER **HISTOPATHOLOGY** 5E-2 IRIS 010262

HISTOPATHOLOGY KIDNEY WHOLE BODY DECREASED WEIGHT ORGANS DECREASED WEIGHT

100

Subchronic Chronic CHEMICAL DOSE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET (mg/cu m) (mg/kg/day) CRITICAL EFFECT (mg/cu m) (mg/kg/day) UF UF LIF CRESOL, M- / (3-METHYLPHENOL) 000108-39-4 NOAEL 50 MG/KG/DAY RAT ORAL: GAVAGE 90 DAYS WHOLE BODY DECREASED WEIGHT GAIN 5E-1 IRIS 005380 NERVOUS SYSTEM NEUROTOXICITY 100 GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. IRIS 010888 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RFD/RFC WORK GROUP. CRESOL, O- / (2-METHYLPHENOL) 000095-48-7 NOAEL 50 MG/KG/DAY RAT ORAL: GAVAGE 90 DAYS WHOLE BODY DECREASED WEIGHT GAIN 5E-1 IRIS 005384 100 NERVOUS SYSTEM NEUROTOXICITY GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. IRIS 010889 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RFD/RFC WORK GROUP. CRESOL, P- / (4-METHYLPHENOL) 000106-44-5 NOAEL 5 MG/(KG-DAY) RABBIT CENTRAL NERVOUS HYPOACTIVITY 5E-3 010516 ORAL: GAVAGE **GESTATION** 5E-3 1000 1000 DAYS 6-18 SYSTEM RESPIRATORY DISTRESS SYSTEM WHOLE BODY MATERNAL DEATH SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. 010890 IRIS CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RFD/RFC WORK GROUP.

		HE	AST TABLE	: SUBCHRON	C AND CHRONIC TOXICIT	Y (OTHER THAN	CARCINOGENIC	ITY)	March 1994
CHEMICAL LEVEL	<u>DOSE</u> ROUTE		<u>CIES</u> Nt Length	TARGET C	RITICAL EFFECT	Subchro [RfC] [RfD] (mg/cu m) (mg/kg/ UF UF	[RfC]	[RfD]	REFERENCE
CUMENE			000098-82-	8					
NOAEL	154 MG/KG/ ORAL: GA		RAT 194 days	KIDNEY	INCREASED WEIGHT	4E 30		IRIS	005392
NOAEL	105.1 PPM INHALATI INTERMIT		RAT 4 WEEKS	CENTRAL NERVOUS SYSTEM NOSE	INVOLVEMENT IRRITATION	9E-2 1000	9E-3 10000		005908
CYANAZ	INE		021725-46	3-2					
NOEL	0.625 MG/K ORAL: DI		DOG 1 YEAR	WHOLE BODY BLOOD BLOOD	DECREASED WEIGHT INCREASED PLATELET COUNT ALTERED CLINICAL CHEMISTRY PARAMETERS	2E 30	:-3 0	2E-3 300	010411
	CHRONIC [	RfD] COM	MENT: THE CHRO		WAS ADOPTED AS THE SUBCHRONI WITHDRAWN FROM IRIS (07/01/92 GENICITY.		CURRENT NUMBER SUB-	JECT TO CH	ANGE.
CYANIDE	<b>.</b>		000057-12-	5					
NOAEL	10.8 MG/KG ORAL: DI	•	RAT 104 WEEKS	WHOLE BODY	DECREASED WEIGHT	2E	:-2	IRIS	005396

THYROID **EFFECTS** 500

NERVE MYELIN DEGENERATION

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: THE CASRN FOR CN- IS 000057-12-5; THE CASRN FOR HCN IS 000074-90-8.

**CYANOGEN** 000460-19-5

> NOAEL 21.6 MG/KG/DAY RAT

IRIS 010263 4E-2 ORAL: DIET 2 YEARS WHOLE BODY DECREASED WEIGHT

THYROID **EFFECTS** 500

NERVE MYELIN DEGENERATION

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

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005398

Subchronic Chronic

CHEMICAL DOSE SPECIES [RfC] [RfC] [RfD] REFERENCE
LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

JTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT <u>(mg/cu m) (mg/kg/day)</u> <u>(mg/kg/day)</u> UF UF UF UF UF

CYANOGEN BROMIDE 000506-68-3

NOAEL 44 MG/KG/DAY RAT

ORAL: DIET 2 YEARS WHOLE BODY DECREASED WEIGHT 9E-2 IRIS 010264

THYROID EFFECTS 500

NERVE MYELIN DEGENERATION

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. THE CHRONIC ORAL RFD

WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CYCLOATE 001134-23-2

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005886

CYCLOHEXANOL 000108-93-0

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005887

CYCLOHEXYLAMINE 000108-91-8

IRIS 005430

NOAEL 30 MG/KG/DAY RAT

ORAL: DIET 90 DAYS WHOLE BODY DECREASED WEIGHT GAIN 3E-1

TESTIS DECREASED WEIGHT 100

CYCLOPENTADIENE 000542-92-7

010494

GENERAL COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

DACTHAL 001861-32-1

NOEL 50 MG/KG/DAY RAT

ORAL: DIET 2 YEARS KIDNEY INCREASED WEIGHT 5E-1 IRIS 005888

ADRENAL INCREASED RELATIVE WEIGHT 100

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

	<u>ecies</u> Ent length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
DALAPON	000075-99	-0				
NOEL 8.45 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	KIDNEY	INCREASED RELATIVE WEIGHT	3E-2 300	IRIS	005889
SUBCHRONIC [RfD]			BY TO DALAPON SODIUM BY CORRECT: RONIC ORAL [RfD].	ING FOR DIFFERENCES IN MC	DECULAR WEIGHT. THE CH	RONIC ORAL
DDT 0	00050-29-3					
NOEL 0.05 MG/KG/DAY ORAL: DIET	RAT 27 WEEKS	LIVER	LESIONS	5E-4 100	IRIS	005408
SUBCHRONIC [RfD] GENERAL COMMENT:	COMMENT: THE ALSO SEE HEAS	CHRONIC ORAL RED T TABLE 3: CARCI	WAS ADOPTED AS THE SUBCHRONIC (NOGENICITY.	ORAL [RfD].		
DECABROMODIPHENYL E	THER 0	01163-19-5				
NOEL 1 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	LIVER	INCREASED WEIGHT	1E-2	IRIS	005891
				100		
SUBCHRONIC [RfD] GENERAL COMMENT:			WAS ADOPTED AS THE SUBCHRONIC NOGENICITY.	ORAL [RfD].		
DI-N-OCTYL PHTHALATE		17-84-0				
LOAEL 175 MG/KG/DAY ORAL: DIET	RAT 7-12 MONTHS	KIDNEY LIVER LIVER LIVER	INCREASED WEIGHT INCREASED WEIGHT INCREASED SGOT ACTIVITY INCREASED SGPT ACTIVITY	2E-2 1000	2E-2 1000	0102 <i>7</i> 5
DIAZINON	000333-41-	-5				
NOAEL 0.09 MG/KG/DAY	RAT		DEGREEOFE AUGUSTUS ATTENDED	or i	Or 1	005000
ORAL: DIET	35-42 DAYS	8L000	DECREASED CHOLINESTERASE ACTIVITY	9E-4 100	9E-4 100	005892

<u>CHEMICAL</u> LEVEL	DOSE ROUTE EXPE	<u>SPECIES</u> Eriment Length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
DIBENZOF	FURAN General Comme	OOO132 NT: DATA INADEQU		ATIVE RISK ASSESSMENT.			005409
	BENZENE, 1,	4- 0001	06-37-6				
NOAEL	ORAL: GAVAGE		LIVER LIVER	INCREASED RELATIVE WEIGHT ALTERED ENZYME ACTIVITIES	1E-1 100	IRIS	005893
DIBROMO	CHLOROMET	THANE (	000124-48-1				
NOEL	21.4 MG/KG/DAY ORAL: GAVAGE		LIVER	LESIONS	2E-1 100	IRIS	005894
	GENERAL COMME	NT: ALSO SEE HEA	ST TABLE 3: CAR	CINOGENICITY.			
	ETHANE, 1,2		06-93-4				
LOAEL	88 PPB INHALATION: INTERMITTENT	HUMAN	SPERM	EFFECTS	8E-4 300	2E-4 1000	010854
	CHRONIC [RfC]		REVIEW, CURRENT	TION [RfC] WAS MODIFIED TO ESTIM NUMBER SUBJECT TO CHANGE. CINOGENICITY.	MATE THE SUBCHRONIC INHAL	ATION [RfC].	
DIBUTYL	PHTHALATE	0000	84-74-2				
NOAÉL	125 MG/KG/DAY ORAL: DIET	RAT 52 WEEKS	WHOLE BODY	INCREASED MORTALITY	1E+0 100	IRIS	005622
						IRIS	010892

CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (07/26/90) BY THE RFD/RFC WORK GROUP.

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Subchronic Chronic DOSE ROUTE CHEMICAL SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

DICAMBA

001918-00-9

NOAEL 3 MG/KG/DAY
ORAL: GAVAGE

RABBIT GESTATION

FETUS

US DECREASED WEIGHT
US INCREASED POST-IMPLANTATION

3E-2 100 IRIS 010945

DAYS 6-18 FETUS INCREAS LOSSES

FETUS DAM INCREASED MORTALITY DECREASED WEIGHT GAIN

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

**DICHLOROBENZENE, 1,2-**

000095-50-1

IRIS

010864

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

GENERAL COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

DICHLOROBENZENE, 1,3-

INTERMITTENT

005414

CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (06/23/92) BY THE RfD/RfC WORK GROUP.

**DICHLOROBENZENE, 1,4-**

000106-46-7

000541-73-1

NOAEL 75 MG/CU M

M RAT

INHALATION: MULT!

MULTI-GENERA LIVER TION

INCREASED WEIGHT IN MALE

2.5E+0

IRIS

010840

SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RfC WAS MODIFIED TO DERIVE THE SUBCHRONIC INHALATION [RfC].

**PARENTS** 

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

**DICHLOROBUTENES** 

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.

005415

UF

UF

UF

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Subchronic Chronic CHEMICAL

SPECIES DOSE [RfC] [RfD] [RfC] [RfD] REFERENCE ROUTE LEVEL EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

DICHLORODIFLUOROMETHANE 000075-71-8

IRIS 005498 NOAEL 90 MG/KG/DAY DOG

ORAL: DIET 90 DAYS NONE OBSERVED 9E-1 005496 100

SUBCHRONIC [Rfc] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

DICHLOROETHANE, 1,1-000075-34-3

NOEL 115 MG/KG/DAY RAT

> INHALATION: 13 WEEKS NONE OBSERVED 1E+0 1E-1 005790

INTERMITTENT 100 1000

SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION.

CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

DICHLOROETHANE, 1,2-000107-06-2

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

DICHLOROETHYLENE, 1,1-000075-35-4

LOAEL 9 MG/KG/DAY RAT

> ORAL: DRINKING 2 YEARS 005419 LIVER LESIONS 9E-3 IRIS

WATER 1000

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfD] COMMENT: UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

Subchronic Chronic [RfC] [RfD] REFERENCE CHEMICAL DOSE SPECIES [RfC] [RfD] LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF DICHLOROETHYLENE, 1,2- (MIXED ISOMERS) 000540-59-0 LOAEL 50 PPM 9E-3 9E-3 010509 ORAL: DRINKING 2 YEARS LIVER LESIONS 1000 1000 WATER SUBCHRONIC (RfD) COMMENT: VALUES DERIVED FOR 1,1-DICHLOROETHYLENE WERE ADOPTED FOR 1,2- DICHLOROETHYLENE MIXED ISOMERS BASED ON ANALOGY. CHRONIC [RfD] COMMENT: VALUES DERIVED FOR 1,1-DICHLOROETHYLENE (000075-35-4) WERE ADOPTED FOR 1,2-DICHLOROETHYLENE MIXED ISOMERS BASED ON ANALOGY. DICHLOROETHYLENE, 1,2-C-000156-59-2 NOAEL 32 MG/KG/DAY RAT 90 DAYS DECREASED HEMATOCRIT 1E-1 1E-2 005420 ORAL: GAVAGE BLOOD 300 3000 BLOOD DECREASED HEMOGLOBIN CHRONIC [RfD] COMMENT: UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE. DICHLOROETHYLENE, 1,2-T-000156-60-5 NOAEL 17 MG/KG/DAY MOUSE 2E-1 IRIS 005895 ORAL: DRINKING 90 DAYS BLOOD INCREASED ALKALINE PHOSPHATASE WATER 100 DICHLOROPHENOL, 2,3-000576-24-9 005315 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT DICHLOROPHENOL, 2,4-000120-83-2 RAT NOEL 3 PPM 005314 3E-3 IRIS ORAL: DRINKING 2 IMMUNE SYSTEM ALTERED IMMUNE FUNCTION WATER **GENERATIONS** 100 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. CHRONIC [RfD] COMMENT: BASED ON A 2-GENERATION REPRODUCTION STUDY WITH EXPOSURES BEFORE AND DURING GESTATION, PARTURITION, AND WEANING OF PUPS.

DICHLOROPHENOL, 2,5- 000583-78-8

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT

005316

Subchronic Chronic CHEMICAL [RfC] [RfD] [RfC] [RfD] REFERENCE ROUTE LEVEL EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF ÜF DICHLOROPHENOL, 2,6-000087-65-0 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005317 DICHLOROPHENOL, 3,4-000095-77-2 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005318 DICHLOROPHENOL, 3.5-000591-35-5 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005319 DICHLOROPHENOXY ACETIC ACID, 2,4- 000094-75-7 NOAEL 1 MG/KG/DAY RAT ORAL: DIET 91 DAYS BLOOD 1E-2 TOXICITY IRIS 010265 LIVER TOXICITY 100 KIDNEY TOXICITY SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. DICHLOROPHENOXY) BUTYRIC ACID, 4-(2,4- / (2,4-DB) 000094-82-6 NOAEL 8 MG/KG/DAY DOG ORAL: DIET 90 DAYS CARDIOVASCULAR HEMORRHAGE 8E-2 IRIS 005890 SYSTEM 100 WHOLE BODY INCREASED MORTALITY DICHLOROPROPANE, 1,1-000078-99-9 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005897 DICHLOROPROPANE, 1,2-000078-87-5 NOAEL 69.3 MG/CU MRAT INHALATION: 13 WEEKS NASAL MUCOSA HYPERPLASIA 1.3E-2 IRIS 005898 INTERMITTENT 100

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

Subchronic Chronic CHEMICAL [RfC] [RfD] REFERENCE **SPECIES** [RfC] [RfD] DOSE LEVEL ROUTE EXPERIMENT LENGTH (mg/cu m) (mg/kg/day) TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) DICHLOROPROPANE, 1,3-000142-28-9 005899 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. DICHLOROPROPANE, 2,2-000594-20-7 005900 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. DICHLOROPROPENE, 1,3- / (TELONE II) 000542-75-6 NOEL 3 MG/KG/DAY RAT IRIS 005901 90 DAYS 3E-3 ORAL: DIET **ORGANS** INCREASED WEIGHT 1000 GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. NOAEL 5 PPM MOUSE IRIS 010351 INHALATION: 2 YEARS NASAL MUCOSA **HYPERTROPHY** 2E-2 INTERMITTENT **NASAL MUCOSA** HYPERPLASIA SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RFC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC]. **DICHLORPROP** 000120-36-5 005896 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. DICYCLOPENTADIENE 000077-73-6 NOEL 32 MG/KG/DAY RAT 005425 3E-2 ORAL: DIET 3 NONE OBSERVED 3E-1

SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) SUBCHRONIC [RfD] COMMENT: BASED ON A 3-GENERATION REPRODUCTION STUDY.

CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

CHRONIC [RfD] COMMENT: BASED ON A 3-GENERATION REPRODUCTION STUDY.

GENERATIONS

1000

100

CHEMICAL LEVEL	<u>dose</u> Route expe	<u>species</u> Riment Length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
DIELDRIN		000060-57-1					
NOAEL	0.005 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	LIVER	LESIONS	5E-5 100	IRIS	005429
		FD] COMMENT: THE NT: ALSO SEE HEAS		D WAS ADOPTED AS THE SUBCHRONI INOGENICITY.	C ORAL [RfD].		
DIETHYL	PHTHALATE	00008	4-66-2				
NOAEL	750 MG/KG/DAY ORAL: DIET	RAT 16 WEEKS	WHOLE BODY ORGANS	DECREASED GROWTH DECREASED WEIGHT	8E+0 100	IRIS	005620
DIETHYL	=	IYL PHOSPHATE NT: DATA INADEQUA		1-45-5 TIVE RISK ASSESSMENT			005922
DIETHYL	ANILINE, N,N- GENERAL COMME			TIVE RISK ASSESSMENT.			005903
DIETHYLI	ENE GLYCOL	MONOBUTYL ET	HER 00011	2-34-5			
NOAEL	18 PPM	RAT		NONE OBSERVED	2E-1 100	2E-2 1000	005482
	CHRONIC [RfC]	COMMENT: UNDER R	EVIEW.				
DIETHYLL	ENE GLYCOL	MONOETHYL ET	HFR 00011	1-90-0			
	200 MG/KG/DAY ORAL: DRINKI WATER	RAT	KIDNEY	HISTOPATHOLOGY		2E+0 100	005478
	CHRONIC [RfD]	COMMENT: BASED O	N A 3-GENERATIO	N REPRODUCTION STUDY.			
NOEL	500 MG/KG/DAY ORAL: DIET	RAT 90 days	KIDNEY Testis	IMPAIRED FUNCTION INCREASED WEIGHT	5E+0 100		005476

CHEMICAL LEVEL	<u>Dose</u> Route		<u>ecies</u> Ent Length	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) Uf	Chro [RfC] (mg/cu m) UF	onic [RfD] (mg/kg/day) UF	REFERENCE
DIETHYL	FORMAN	IIDE	0006	17-84-5						
NOEL	0.546 MG/ ORAL: 6		YS/WEEK 73 WEEKS	RAT	NONE OBSERVED		1.1E-2 100		1.1E-2 100	010437
DIETHYL	HYDRAZ	NE, 1,2	- 001	615-80-1						
	GENERAL	COMMENT:	DATA INADEQU	IATE FOR QUANT	ITATIVE RISK ASSESSMENT. AL	SO SEE HEAST TABLE	3: CARCINOGE	ENICITY.		005921
DIMETH	OATE		000060-	·51-5						
NOEL	0.05 MG/K ORAL: D		RAT 2 YEARS	BRAIN	DECREASED CHOLINESTER ACTIVITY	RASE	2E-4 300		IRIS	005923
	SUBCHRO	IIC [RfD]	COMMENT: THE	CHRONIC ORAL	RFD WAS ADOPTED AS THE SUBG	CHRONIC ORAL [RfD].				
DIMETH	YLANILIN	E, N,N-	0001	21-69-7						
LOAE	L 22.32 MG/ ORAL: (		MOUSE 13 WEEKS	SPLEEN	EFFECTS		2E-2 1000		IRIS	005924
DIMETH	YLFORMA	MIDE, N	N,N- 00	00068-12-2						
NOAE	L 96 MG/KG/ ORAL: [		RAT 119 DAYS	LIVER	EFFECTS		1E+0 100		1E-1 1000	005925
LOAE	L 22 MG/CU	M	HUMAN							
	INHALAT INTERMI			LIVER GASTRO INTESTINAL	EFFECTS EFFECTS SYSTEM	3E-2 300		IRIS		010352
	SUBCHRO	IIC [RfC]	COMMENT: THE	E CHRONIC INHA	LATION RFC WAS ADOPTED AS TI	HE SUBCHRONIC INHALA	ATION [RFC].			
DIMETH	YLPHENO	L, 2,3-	000!	526-75-0						
					ITATIVE RISK ASSESSMENT					005926

CHEMICAL LEVEL		<u>PECIES</u> Ment Length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] <u>(mg/cu m) (mg/kg/day)</u> UF UF	REFERENCE
DIMETHY	LPHENOL, 2,4-	00010	5-67-9				
NOAEL	50 MG/KG/DAY ORAL: GAVAGE	MOUSE 90 DAYS	NERVOUS SYSTEM BLOOD	EFFECTS ALTERATIONS	2E-1 300	IRIS	010266
DIMETHY	LPHENOL, 2,5- GENERAL COMMENT:		5-87-4 TE FOR QUANTITAT	IVE RISK ASSESSMENT.			005928
	LPHENOL, 2,6-	00057	6-26-1				
NOEL	O.6 MG/KG/DAY ORAL	RAT 8 MONTHS	WHOLE BODY ORGANS, MAJOR	INCREASED WEIGHT LESIONS	6E-3 100	IRIS	005431
	YLPHENOL, 3,4-		5-65-8				
NOEL	1.4 MG/KG/DAY ORAL	RAT 8 MONTHS	WHOLE BODY ORGANS, MAJOR CARDIOVASCULAR SYSTEM	DECREASED WEIGHT LESIONS ALTERED BLOOD PRESSURE	1E-2 100	IRIS	005437
DIMETHY	LPHTHALATE	0001	31-11-3				
NOEL	1000 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	KIDNEY	EFFECTS	1E+1 100	1E+1 100	010267
	CHRONIC RFC COM	MENT: THE CHRON	IC INHALATION RF	C IS CONSIDERED NOT VERIFIABLE	(07/26/90) BY THE RfD/RfC	IRIS WORK GROUP.	010894
DIMETHY	LTEREPHTHAL	ATE 000	0120-61-6				
LOAEL	125 MG/KG/DAY ORAL: DIET	RAT 103 WEEKS	KIDNEY	INFLAMMATION	1E-1 1000	IRIS	005930

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHEMICAL LEVEL		<u>SPECIES</u> IMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC] [RfD] [RfC]	ronic [RfD] (mg/kg/day) Uf	REFERENCE
DIMETHY	LUREA, N,N- GENERAL COMMENT		18-94-7 NTE FOR QUANT	ITATIVE RISK ASSESSMENT			005931
DINITRO-	O-CRESOL, 4,6		34-52-1 RONIC INHALAT	ION [RfC] IS CONSIDERED NOT VER	IFIABLE (02/11/93) BY THE RFD/RFC WOR	K GROUP.	010470
DINITRO-	P-CRESOL, 2,6 GENERAL COMMENT		9-93-8 Ate for quant	ITATIVE RISK ASSESSMENT			005934
	BENZENE, 1,2- 0.4 Mg/kg/day ORAL: DRINKING WATER	RAT	8-29-0 Spleen	INCREASED WEIGHT	4E-3 100	4E-4 1000	010201
				GY TO 1,3-DINITROBENZENE. TO 1,3-DINITROBENZENE.			
	BENZENE, 1,3- 0.4 mg/kg/day ORAL: DRINKING WATER	RAT	9-65-0 SPLEEN	INCREASED WEIGHT	1E-3 100	IRIS	010471
	BENZENE, 1,4- 0.4 MG/KG/DAY ORAL: DRINKING WATER	RAT	O-25-4 SPLEEN	INCREASED WEIGHT	4E-3 100	4E-4 1000	010202
	SUBCHRONIC [Rf			GY TO 1,3-DINITROBENZENE. TO 1,3-DINITROBENZENE.			
DINITROF	PHENOL, 2,3- GENERAL COMMEN	OOOO6:		ITATIVE RISK ASSESSMENT			005936

Subchronic Chronic REFERENCE CHEMICAL DOSE SPECIES [RfC] [RfD] [RfC] [RfD] LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF DINITROPHENOL, 2,4-000051-28-5 LOAEL 2 MG/KG/DAY HUMAN ORAL EYE CATARACT 2E-3 IRIS 010438 1000 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. IRIS 010895 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (06/13/91) BY THE RFD/RFC WORK GROUP. DINITROPHENOL, 2,5-000329-71-5 005937 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT DINITROPHENOL, 2,6-000573-56-8 005938 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT DINITROPHENOL, 3,5-000586-11-8 005939 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT DINITROTOLUENE, 2.3-000602-01-7 005940 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 000121-14-2 DINITROTOLUENE, 2,4-NOAEL 0.2 MG/KG/DAY DOG 2E-3 IRIS 005941 ORAL: GELATIN UP TO 2 CENTRAL NERVOUS NEUROTOXICITY CAPSULE YEARS SYSTEM 100 ERYTHROCYTES **HEINZ BODIES** BILIARY TRACT HYPERPLASIA SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD IS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. 010896 IRIS CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (12/20/90) BY THE RFD/RFC WORK GROUP. DINITROTOLUENE, 2.5-000619-15-8 005942 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT

1-39

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route		<u>ECIES</u> Ent Length	TARGET C	RITICAL EFFI	ест	(RfC) (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chron [RfC] (mg/cu m) (1 UF	[RfD]	REFERENCE
DINITRO	OLUENE	. 2,6-	00060	06-20-2							
MOAFI	4 MG/KG/D	AY	DOG								
NONEL	ORAL: I		13 WEEKS	WHOLE BODY CENTRAL NERVOUS SYSTEM BLOOD BLOOD BILE DUCT KIDNEY	MORTALITY NEUROTOXIC HEINZ BODI METHEMOGLO HYPERPLASI HISTOPATHO	ES BINEMIA A		1E-2 300		1E-3 3000	005943
				REVIEW, CURRENT NUM ST TABLE 3: CARCING		TO CHANGE.					
DINITRO	OLUENE	, 3,4-	00061	0-39-9							
	GENERAL	COMMENT:	DATA INADEQU	ATE FOR QUANTITATIV	Æ RISK ASSE	SSMENT					005944
DINOSEB			000088-85	-7							
	4 40 440 46		RAT	-,							
LONEL	1 MG/KG/D ORAL: I		29 WEEKS	FETUS	DECREASED	WEIGHT		1E-3 1000		IRIS	005945
				CHRONIC ORAL RFD WAS							
DIPHENY	LAMINE,	N,N-	00012	22-39-4							
NOEL	2.5 MG/KG	/DAY	DOG								
	ORAL: I	TET	2 YEARS	WHOLE BODY LIVER KIDNEY	DECREASED INCREASED INCREASED			2.5E-2 100		IRIS	005946
	SUBCHRO	IC [RfD]	COMMENT: THE	CHRONIC ORAL RFD	AS ADOPTED	AS THE SUBCHRONI	C ORAL [RfD]	•			
DIRECT L				399-55-7							2050/7
	GENERAL	COMMENT:	DATA INADEQU	ATE FOR QUANTITATIV	VE RISK ASSE	SSMENT					005947

CHEMICAL LEVEL	<u>Dose</u> Route ex	<u>Species</u> (Periment Length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
DISULFO	TON	000298	3-04-4				
LOAEL	. 0.04 MG/KG/D						
	ORAL: DIET	2 YEARS	EYE Blood	DEGENERATION CHOLINESTERASE INHIBITION	4E-5 1000	IRIS	010412
	SUBCHRONIC	[RfD] COMMENT: T	HE CHRONIC ORAL RFD	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].		
ENDOSU	LFAN	00011	5-29-7				
NOAEL	. 15 PPM ORAL: DIET	RAT 2 YEARS	HILO E BODY	DECREAGED LIFTOUT ALL	<b></b>		
	ORAL: DIEI	Z TEARS	WHOLE BODY KIDNEY BLOOD VESSEL	DECREASED WEIGHT GAIN GLOMERULONEPHROSIS ANEURYSMS	6 <b>E-3</b> 100	6E-3 100	010926
NOAEL	. 10 PPM	DOG					
	ORAL: DIET	1 YEAR	WHOLE BODY	DECREASED WEIGHT GAIN			010938
	SUBCHRONIC CHRONIC [Rf	[RfD] COMMENT: T D] COMMENT: WITH STUDIES.	HE CHRONIC ORAL [Rf[ DRAWN FROM IRIS (12,	D) IS ADOPTED AS THE SUBCHORNI (01/92). UNDER REVIEW, CURRENT	C ORAL [RfD]. NUMBER SUBJECT TO CHANGE	. BASED ON CO-CRITICAL	RAT AND DOG
ENDOTH	ALL	000145	5-73-3				
NOEL	2 MG/KG/DAY ORAL: DIET	DOG 2 YEARS	STOMACH SMALL INTESTINE	INCREASED WEIGHT INCREASED WEIGHT	2E-2 100	IRIS	005948
	SUBCHRONIC	[RfD] COMMENT: T	HE CHRONIC ORAL RFD	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].		
ENDRIN		000072-20	0-8				
NOEL	0.025 MG/KG/E ORAL: DIET						
	OKAL: UIEI	2 YEARS	CENTRAL NERVOUS		3E-4 100	IRIS	005445
			LIVER	LESIONS			
	SUBCHRONIC	[RfD] COMMENT: T	HE CHRONIC ORAL RFD	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].		

March 1994

Subchronic Chronic CHEMICAL DOSE **SPECIES** [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

EPICHLOROHYDRIN 000106-89-8

LOAEL 37.8 MG/CU M RAT

INHALATION: 136 WEEKS KIDNEY LESIONS 2E-3 010440
INTERMITTENT 1000 1000

SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE-TO-ROUTE EXTRAPOLATION. THE CHRONIC ORAL [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. CHRONIC [RfD] COMMENT: WITHDRAWN FROM IRIS (04/01/92).
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

NOAEL 19 MG/CU M RAT

INHALATION: 90 DAYS NASAL LESIONS 1E-2 IRIS 010492
INTERMITTENT EPITHELIUM 100

EPTC 000759-94-4

NOEL 2.5 MG/KG/DAY RAT

ORAL: DIET 2 HEART DEGENERATIVE CARDIOMYOPATHY 2.5E-2 IRIS 005959
GENERATIONS 100

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].
CHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS DETERMINED FROM A 2-GENERATION REPRODUCTION STUDY.

ETHOPROP 013194-48-4

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005951

ETHOXYETHANOL ACETATE, 2- 000111-15-9

010507

SUBCHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)
CHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

ETHOXYETHANOL ACRYLATE, 2- 000106-74-1

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005953

ETHOXYETHANOL DODECANOATE, 2- 000106-13-8

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005956

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route ex	SPECIES PERIMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chron [RfC] (mg/ci m) (m UF	[RfD]	REFERENCE
ETHOXY			068554-00-7	, ATIVE RISK ASSESSMENT					005055
	GENERAL COM	TENT: DATA INADEWO	AIR FOR QUARITIE	(IIAE KISK VSSESSMEN)					005955
ETHOXY	ETHANOL, 2	- 0001	10-80-5						
LOAEL	357 MG/KG/DAY								
	ORAL: GAVA	GE 103 WEEKS	WHOLE BODY	DECREASED WEIGHT				4E-1 1000	005470
MOEI	50 UL/KG/DAY	RAT							
HOLL	ORAL: GAVA		FETUS	SKELETAL MALFORMATIONS		5E-1			005468
						100			
	SUBCHRONIC	[RfD] COMMENT: BAS	ED ON A REPRODUC	TION STUDY WITH EXPOSURES DURI	NG DAYS 1-21	OF GESTATION.	•		
NOACI	380 MG/CU M	DADOLT							
NUAEL	INHALATION		BLOOD	ALTERED HEMATOLOGY	2E+0		IRIS		010441
	INTERMITTE	NT .			30				
FTHOYVI	ETHYL METH	ACRYLATE, 2-	002370-63-6	n					
LIIIOXII		•		ATIVE RISK ASSESSMENT					005954
ETHYL A	CETATE	000141	-78-6						
NOEL	900 MG/KG/DAY								
	ORAL: GAVA	GE 90 DAYS	WHOLE BODY	INCREASED MORTALITY DECREASED WEIGHT		9E+0 100		IRIS	005957
ETHYL BI	FNZENE	000100	-41-4						
	mi valli v L	000100	च।~ <b>च</b>					IRIS	010867
							IRIS		010866
		_							

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

Subchronic Chronic [RfC] CHEMICAL DOSE SPECIES [RfC] [RfD] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH (mg/cu m) (mg/kg/day) **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) UF UF **ETHYL CHLORIDE** 000075-00-3 NOAEL 1504 PPM MOUSE 010371 INHALATION: 10 DAYS **FETUS DEVELOPMENTAL TOXICITY** 1E+1 IRIS INTERMITTENT 300 SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RFC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC]. CHRONIC [RFC] COMMENT: BASED ON A DEVELOPMENTAL STUDY WITH EXPOSURES DURING DAYS 6-15 OF GESTATION. **ETHYL ETHER** 000060-29-7 NOAEL 500 MG/KG/DAY RAT 010396 90 DAYS **EFFECTS** 2E+0 IRIS ORAL: GAVAGE LIVER 300 ETHYL METHACRYLATE 000097-63-2 NOEL 7.5 MG/KG/DAY RAT 005961 ORAL: DRINKING 9E-2 9E-2 2 YEARS KIDNEY INCREASED RELATIVE WEIGHT 100 100 WATER CHRONIC [RfD] COMMENT: CALCULATED FROM METHYL METHACRYLATE DATA BY MULTIPLYING BY THE RATIO OF THE MOLECULAR WEIGHTS (114.5/100.13). ETHYL-O-XYLENE, 4-000934-80-5 010472 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ETHYLANILINE, N-000103-69-5 005958 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT ETHYLENE CYANOHYDRIN 000109-78-4 NOEL 30 MG/KG/DAY RAT 3E-1 005780 ORAL: DRINKING 90 DAYS **HEART** DECREASED WEIGHT 3E-1 100 100 WATER BRAIN DECREASED WEIGHT

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route expe	<u>species</u> Riment Length	TARGET	CRITICAL EFFECT	Subchror [RfC] [RfD] (mg/cu m) (mg/kg/c UF UF	[RfC] [RfD]	REFERENCE Y)
ETHYLEN	IE DIAMINE	00010	7-15-3				
NOAEL	22.6 MG/KG/DAY ORAL: DIET	RAT 3 MONTHS	HEART BLOOD	DECREASED WEIGHT HEMATOLOGIC CHANGES	2E- 100		005796
	CHRONIC RFC CO	MMENT: THE CHRON	IC INHALATION	RFC IS CONSIDERED NOT VERIFIA	BLE (12/18/90) BY THE R	IRIS fD/RfC WORK GROUP.	010898
ETHYLEN	IE GLYCOL	00010	7-21-1				
NOEL	200 MG/KG/DAY	RAT				IRIS	005454
	ORAL: DIET		FETUS	FETOTOXICITY	2E+ 100		005452
	SUBCHRONIC (Rf	D) COMMENT: BASE	D ON A REPROD	OUCTION STUDY WITH EXPOSURES DUI	RING DAYS 6-15 OF GESTA	TION.	
		NOBUTYL ETH	ER 0001	11-76-2			
NOAEL	121 MG/CU M INHALATION: INTERMITTENT	RAT 13 WEEKS	BL000	ALTERED HEMATOLOGY	2E-1 100	2E-2 1000	010353
	CHRONIC [RfC]	COMMENT: UNDER R	EVIEW, CURREN	T NUMBER SUBJECT TO CHANGE.			
ETHYLEN	IE THIOUREA	00009	6-45-7				
LOAEL	0.25 MG/KG/DAY ORAL: DIET	RAT 24 MONTHS	THYROID	HYPERPLASIA	8E-300		010397
		D] COMMENT: THE		RFD WAS ADOPTED AS THE SUBCHRON	NIC ORAL [RfD].		44445
	CHRONIC [RfC]	COMMENT: THE CHR	ONIC INHALATI	ON [RfC] IS CONSIDERED NOT VER	FIABLE (08/12/92) BY T	HE RFD/RFC WORK GROUP.	010899
ETHYLTO	LUENE, M-	000620	D-14-4				
	GENERAL COMMEN	IT: DATA INADEQUA	TE FOR QUANTI	TATIVE RISK ASSESSMENT			005963

	<u>ECIES</u> Ent Length	TARGET C	RITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
ETHYLTOLUENE, O- GENERAL COMMENT:	000611 DATA INADEQUA	=	/E RISK ASSESSMENT			005962
ETHYLTOLUENE, P- GENERAL COMMENT:	000622 DATA INADEQUA		/E RISK ASSESSMENT			005964
FLUORANTHENE NOAEL 125 MG/KG/DAY ORAL: GAVAGE	OOO2O6- MOUSE 90 DAYS	44-0 KIDNEY LIVER BLOOD	NEPHROPATHY WEIGHT CHANGES HEMATOLOGICAL CHANGES	4E-1 300	IRIS	010168
FLUORENE NOAEL 125 MG/KG/DAY ORAL: GAVAGE	000086-73 MOUSE 13 WEEKS	-7 ERYTHROCYTES	DECREASED COUNTS	4E-1 300	IRIS	010169
FLUORINE / (SOLUBLE FLU NOAEL 0.06 MG/KG/DAY ORAL: DRINKING WATER	JORIDE) 00 HUMAN	7782-41-4 тоотн	FLUOROSIS	6E-2 1	IRIS	005965
FLURIDONE NOEL 200 PPM ORAL: DIET	059756-60 RAT 2 YEARS	-4 KIDNEY TESTIS WHOLE BODY ORGANS	GLOMERULONEPHRITIS ATROPHY DECREASED WEIGHT DECREASED WEIGHT	8E-2 100	IRIS	005966

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

<u>CHEMICAL</u> LEVEL		PECIES MENT LENGTH	TARGET	CRITICAL EFFECT	[RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
FOLPET		000133-07-3						
NOEL	10 MG/KG/DAY ORAL: CAPSULE	DOG 1 YEAR	WHOLE BODY BLOOD	ALTERED WEIGHT GAIN ALTERED CHEMISTRY		1E-1 100	IRIS	005967
	SUBCHRONIC [RfD] GENERAL COMMENT:	COMMENT: THE (	CHRONIC ORAL RFD TABLE 3: CARCIN	WAS ADOPTED AS THE SUBCHRONIC OGENICITY.	ORAL [RfD]	•		
FORMALI	DEHYDE	000050	-00-0					
NOAEL	15 MG/KG/DAY	RAT						
	ORAL: WATER	2 YEARS	GASTRO- INTESTINAL TRAC	LESIONS T		2E - 1 100	IRIS	010398
	GENERAL COMMENT:	ALSO SEE HEAST	TABLE 3: CARCIN	OGENICITY.				
FORMALI	DEHYDE CYANO	HYDRIN (	000107-16-4					
	GENERAL COMMENT:	DATA INADEQUAT	E FOR QUANTITATI	VE RISK ASSESSMENT				005782
FORMIC A	ACID 200 mg/kg/day	000064-18	3-6					
	ORAL: WATER	MULTI- GENERATION	MHOLE BODY	DECREASED GROWTH		2E+0 100	2E+0 100	010268
	CHRONIC [RfD] CO	MMENT: BASED ON CHANGE.	A MULTI-GENERAT	ION STUDY. WITHDRAWN FROM IRIS	(12/01/90)	. UNDER REVIE	W, CURRENT NUMBER SUBJ	ECT TO
FURAN		000110-00-9						
NOAEL	1.4 MG/KG/DAY ORAL: GAVAGE	MOUSE 13 WEEKS	LIVER	LESIONS		1E-2 100	IRIS	005462

Subchronic Chronic CHEMICAL [RfC] [RfC] [RfD] REFERENCE SPECIES [RfD] LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) **FURFURAL** 000098-01-1 LOAEL 7.9 MG/KG/DAY RAT ORAL: GAVAGE 13 WEEKS LIVER **HEPATOTOXICITY** 3E-2 IRIS 005466 300 SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) **GLYCIDALDEHYDE** 000765-34-4 NOAEL 29 MG/CU M RAT 1E-2 4E-3 1E-3 IRIS 005968 INHALATION: 12 WEEKS WHOLE BODY DECREASED WEIGHT INTERMITTENT 300 300 3000 KIDNEY **EFFECTS** SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.5. CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.5. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. HEPTACHLOR 000076-44-8 NOEL 0.15 MG/KG/DAY RAT 005506 5E-4 IRIS ORAL: DIET 2 YEARS LIVER INCREASED WEIGHT 300 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY HEPTACHLOR EPOXIDE 001024-57-3 LOAEL 0.0125 MG/KG/DAY DOG IRIS 010399 1.3E-5 ORAL: DIET **60 WEEKS** LIVER INCREASED RELATIVE WEIGHT 1000 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

HEPTANE, N- 000142-82-5

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT

005969

Subchronic Chronic CHEMICAL SPECIES [RfC] [RfD] (RfC) [RfD] REFERENCE ROUTE LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) **HEXABROMOBENZENE** 000087-82-1 NOAEL 2 MG/KG/DAY RAT ORAL: DIET 12 WEEKS LIVER INDUCED CARBOXYLESTERASE 2E-2 IRIS 005970 ACTIVITY 100 **HEXACHLOROBENZENE** 000118-74-1 IRIS 010868 IRIS 010900 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (11/15/90) BY THE RFD/RFC WORK GROUP. SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. **HEXACHLOROBUTADIENE** 000087-68-3 LOAEL 0.5 MG/KG/DAY MOUSE ORAL: DIET 13 WEEKS **RENAL TUBULES** REGENERATION 2E-4 010927 1000 CHRONIC [RfD] COMMENT: WITHDRAWN FROM IRIS (05/01/93). UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE. SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. HEXACHLOROCYCLOHEXANE, DELTA- 000319-86-8 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ALSO SEE HEAST TABLE 3: CARCINOGENICITY. 010495

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

HEXACHLOROCYCLOHEXANE, EPSILON- 006108-10-7

010496

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route ex	<u>SPECIES</u> (PERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchron [RfC] [RfD] (mg/cu m) (mg/kg/de UF UF	[RfC] [RfD]	REFERENCE
HEXACH	LOROCYCLO	DHEXANE, GAMM	1A- 000058-8	9-9			
NOAEL	0.33 MG/KG/D/	NY RAT					
	ORAL: DIET	12 WEEKS	LIVER KIDNEY	TOXICITY TOXICITY	3E-3 100	IRIS	005537
	GENERAL COM	MENT: ALSO SEE HEA	ST TABLE 3: CARCI	NOGENICITY.			
	CHRONIC [Rf	C) COMMENT: THE CH	RONIC INHALATION	[RfC] IS CONSIDERED NOT VERIF	IABLE (05/27/92) BY TH	E RfD/RfC WORK GROUP.	010903
		PENTADIENE	000077-47-4				
NOAEL	7.1 MG/KG/DAY	rat 13 weeks	FORESTOMACH	LESIONS	7E-2	! IRIS	005299
		is weeks	1 OKES I GENERI	LLSTONS	100	ikis	003299
NOAEL	0.15 PPM INHALATION	RAT: 13 WEEKS	NASAL CAVITY	SQUAMOUS METAPLASIA	7E-4	7E-5	010445
	INTERMITTE			GOMIOGO FIETA EROTA	100	1000	010445
LIEVACLU	000571144	·					
	LOROETHAN 1 mg/kg/day	NE OOO RAT	067-72-1				
HONEL	ORAL: DIET		KIDNEY	DEGENERATION	1E-2 100	IRIS	005518
	GENERAL COM	MENT: ALSO SEE HEA	ST TABLE 3: CARCI	NOGENICITY.			
							010904
	CONCORIC [KT	C) COMMENT: THE CH	KONIC INHALATION	[RfC] IS CONSIDERED NOT VERIF	IABLE (11/05/92), BY 1	HE RTD/RTC WORK GROUP.	
HEXACHI	LOROPHENE	0000	70-30-4				
LOAEL	0.75 MG/KG/D/						
	ORAL: DIET	13 WEEKS	NERVOUS SYSTEM	EFFECTS	3E-3 300	IRIS	005972
HEXAME	THYLENE DI		0124-09-4				
	GENERAL COM	MENT: DATA INADEQU	ATE FOR QUANTITAT	IVE RISK ASSESSMENT			005973

CHEMICAL LEVEL		<u>CIES</u> NT LENGTH	TARGET	CRITICAL EFFECT	Sui (RfC) (mg/cu m) (m UF		Chronic (RfC) (RfD) <u>mg/cu m) (mg/kg/day)</u> UF UF	REFERENCE
HEXANE,	N-	000110-54	-3					
LOAEL !	570 MG/KG/DAY ORAL	RAT	NERVOUS SYSTEM TESTIS	NEUROPATHY ATROPHY		6E-1 1000	6E-2 10000	005974
LOAEL	73 MG/CU M INHALATION: INTERMITTENT	HUMAN	NERVOUS SYSTEM	NEUROTOXICITY	2E-1 300		IRIS	010273
	SUBCHRONIC [RfC]	COMMENT: THE	CHRONIC INHALATIO	N RFC WAS ADOPTED AS THE	SUBCHRONIC INHALATI	ON [RfC].		
HEXANON	JF 2-	000591-7	8-6					
112/1/11011	•			VE RISK ASSESSMENT				005976
HYDROGE	N SULFIDE	00778	3-06-4					
	3.1 MG/KG/DAY ORAL: FOOD	PIG 105 DAYS	GASTRO- INTESTINAL SYST	DISTURBANCE EM		3E-2 100	IRIS	010269
NOAEL 4	42 MG/CU M INHALATION: INTERMITTENT	MOUSE 13 WEEKS	NASAL MUCOSA	INFLAMMATION	9E-3 100		IRIS	010354
HYDROQL	JINONE	000123	-31-9					
NOAEL 4	4.29 MG/KG/DAY ORAL	HUMAN 3-5 MONTHS	BL000	HEMATOLOGICAL EFFECTS		4E-1 10	4E-2 100	005526
	CHRONIC RFC COMME	NT: THE CHRON	IC INHALATION RFC	IS CONSIDERED NOT VERIFI	ABLE (10/01/90) BY	THE RfD/RfC I	IRIS WORK GROUP.	010905
IRON		07439-89-6 Data inadequa	TE FOR QUANTITATI	VE RISK ASSESSMENT				005527

March 1994

<u>CHEMICAL</u> LEVEL		<u>ecies</u> Ent length	TARGET	CRITICAL EFFECT	[RfC] <u>(mg/cu m)</u> UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
ISOBUTY	L ALCOHOL	00007	<b>78-83-1</b>					
	316 MG/KG/DAY ORAL: GAVAGE	RAT 13 WEEKS	NERVOUS SYSTEM			3E+0 100	IRIS	005977
ISOPHOR	ONE	000078-5	59-1					
NOEL	150 MG/KG/DAY ORAL: CAPSULE	DOG 90 DAYS	KIDNEY	LESIONS		2E+0 100	IRIS	005910
	GENERAL COMMENT:	ALSO SEE HEAS	T TABLE 3: CARCI	NOGENICITY.				
	CHRONIC RFC COMM	ENT: THE CHRON	IIC INHALATION RE	C IS CONSIDERED NOT VERIFIABLE	(11/15/90)	BY THE RfD/Rf	IRIS FC WORK GROUP.	010906
ISOPROPA	AI IN	033820-5	3-0					
	15 MG/KG/DAY	RAT	<b>J</b> - <b>U</b>					
	ORAL: DIET	90 DAYS	BLOOD ORGANS, UNSPECIFIED	HEMATOLIGICAL EFFECTS ALTERED WEIGHTS		1.5E-1 100	IRIS	005978
LACTONI	TRILE	000078-9	7-7					
	GENERAL COMMENT:	DATA INADEQUA	TE FOR QUANTITAT	IVE RISK ASSESSMENT				005783
LEAD	C	07439-92-1						,
	CHRONIC [RfC] CON CHRONIC [RfD] CON GENERAL COMMENT:	4MENT: REFER T	O IRIS	CHNICAL INFORMATION, SECTION V	ON NATIONAL	AMBIENT AIR	QUALITY STANDARDS.	010447
LEAD ALK	(VI C							

### LEAD ALKYLS

010448

CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS. CHRONIC [RfD] COMMENT: REFER TO IRIS

GENERAL COMMENT: DIMETHYLETHYL LEAD; METHYLTRIETHYL LEAD; TETRABUTYL LEAD; TETRAETHYL LEAD; TETRAMETHYL LEAD; TETRAPROPYL LEAD; TRIMETHYL LEAD; TRIMETHYL LEAD; TRIMETHYL LEAD; TRIMETHYL LEAD; TRIPROPYL LEAD

<u>CHEMICAL</u> LEVEL		<u>PECIES</u> Ment Length	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
LINURON LOAEL	0.625 MG/KG/DAY ORAL: DIET	000330-55- DOG 2 YEARS	BL000	HEMATOLOGICAL EFFECTS		2E-3 300	IRIS	005990
MALATHI	GENERAL COMMENT:	OO0121-7	T TABLE 3: CA	RFD WAS ADOPTED AS THE SUBCHRO RCINOGENICITY.	NIC ORAL [RfD].			
NOEL	0.23 MG/KG/DAY ORAL: CAPSULE SUBCHRONIC [RfD]	HUMAN 47 DAYS COMMENT: THE	BLOOD CHRONIC ORAL	HEMATOLOGICAL EFFECTS RFD WAS ADOPTED AS THE SUBCHRO	NIC ORAL [RfD].	2E-2 10	IRIS	005991
	CHRONIC [RFC] CO	00010						
NOVEL	10 MG/KG/DAY ORAL: DIET SUBCHRONIC (RfD)	RAT 2 YEARS COMMENT: THE	KIDNEY	LESIONS RFD WAS ADOPTED AS THE SUBCHRO	NIC ODAL IDFOI	1E-1 100	IRIS	005992
	YDRAZIDE	00012		KID WAS ASSITED AS THE SOSSIAN	are once this.			
	ORAL: DIET	28 MONTHS	KIDNEY CHRONIC ORAL	ALTERED FUNCTION  R FD WAS ADOPTED AS THE SUBCHRO	NIC ORAL [RfD].	5E-1 1000	IRIS	005993
MALONON	NITRILE 0.21 MG/KG/DAY	000109-	77-3					
	ORAL: GAVAGE	120 DAYS	LIVER Spleen	EFFECTS EFFECTS		2E-4 1000	2E-5 10000	005994

CHEMICAL LEVEL	<u>pose</u> Route exp	<u>SPECIES</u> ERIMENT LENGTH	TARGET C	RITICAL EFFECT	[RfC] (mg/cu m) ( UF	Subchronic [RfD] <u>(mg/kg/day)</u> UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day UF UF	REFERENCE )
MANCO	7FB	008018-0	1.7					
	2.9 MG/KG/DAY	RAT	1-7					
	ORAL: DIET	90 WEEKS	THYROID	GOITROGENIC EFFECTS		3E-2 100	3E-2 100	005995
MANEB		012427-38-	2					
	5 MG/KG/DAY	MONKEY	<b>-</b>					
	ORAL: DIET	6 MONTHS	THYROID	INCREASED WEIGHT		5E-2 100	IRIS	005996
MANGA	HECE	007420.0	)C E					
_	NESE .0.005 MG/KG/DA	007439-9 Y Human	6-0					
NOVEL	ORAL: DRINKI WATER		CENTRAL NERVOUS SYSTEM	EFFECTS		5E-3 1	IRIS	010850
	SUBCHRONIC (R	fD] COMMENT: THE DIETARY INFO	CHRONIC ORAL WATER RMATION.	R RFD WAS ADOPTED AS THE SUBCH	RONIC ORAL W	ATER [RfD].	SEE IRIS FOR SPECIFIC	
NOAFI	0.14 MG/KG/DAY	HUMAN						
NONEL	ORAL: DIET	CHRONIC	CENTRAL NERVOUS	EFFECTS		1.4E-1 1	IRIS	010851
	SUBCHRONIC [R	fD] COMMENT: THE	CHRONIC ORAL FOOD	RFD WAS ADOPTED AS THE SUBCHR	ONIC ORAL FO	OD TREDI. SE	F IRIS FOR SPECIFIC D	IFTARY
		INFORMATION.				on fully of		
	SUBCHRONIC [R	fC] COMMENT: A SU	BCHRONIC [RfC] HAS	NOT BEEN DERIVED FOR MANGANE	SE.		IRIS	010959
MEPHOS	FOLAN	000950-	10-7					
	0.09 MG/KG/DAY	RAT	10-7					
	ORAL: DIET	17 WEEKS	LIVER	ALTERED WEIGHT		9E-4	9E-5	005997
			KIDNEY BLOOD	ALTERED WEIGHT		100	1000	
			BLOOD	DECREASED CHOLINESTERASE ACTIVITY				
			ERYTHROCYTES	DECREASED CHOLINESTERASE				
			BRAIN	ACTIVITY DECREASED CHOLINESTERASE ACTIVITY				

UF

UF

UF

Subchronic Chronic CHEMICAL [RfC] [RfC] [RfD] REFERENCE [RfD] DOSE ROUTE SPECIES EXPERIMENT LENGTH **TARGET** (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) LEVEL CRITICAL EFFECT

MERCURY, ELEMENTAL VAPOR 007439-97-6

SUBCHRONIC [RFC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

MERCURY, INORGANIC 007439-97-6

RAT

ORAL: PARENTERAL KIDNEY EFFECTS 3E-4 3E-4 005800

CHRONIC [RfD] COMMENT: UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.

SUBCHRONIC [RfC] COMMENT: THIS VALUE IS SPECIFICALLY FOR ELEMENTAL MERCURY.

CHRONIC [RfC] COMMENT: THIS VALUE IS SPECIFICALLY FOR ELEMENTAL MERCURY. UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.

MERPHOS 000150-50-5

NOEL 0.1 MG/KG/DAY HEN

ORAL: CAPSULE 3 MONTHS NERVOUS SYSTEM ATAXIA 3E-4 IRIS 005998

NERVOUS SYSTEM DELAYED NEUROTOXICITY 300

WHOLE BODY DECREASED WEIGHT

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (06/25/92) BY THE RfD/RfC WORK GROUP.

SUBCHRONIC (RfD) CPMMENT: THE CHRONIC ORAL RFD WAS MODIFIED TO ESTIMATE THE SUBCHRONIC ORAL [RfD].

MERPHOS OXIDE 000078-48-8

NOEL 0.1 MG/KG/DAY HEN

ORAL: CAPSULE 3 MONTHS NERVOUS SYSTEM ATAXIA 3E-4 IRIS 005999

NERVOUS SYSTEM DELAYED NEUROTOXICITY 300

WHOLE BODY DECREASED WEIGHT

010908

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (06/25/92) BY THE RfD/RfC WORK GROUP. SUBCHRONIC [RfD] CPMMENT: THE CHRONIC ORAL RfD WAS MODIFIED TO ESTIMATE THE SUBCHRONIC ORAL [RfD].

010907

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route ex	<u>SPECIES</u> XPERIMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	subchronic (RfD) mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
METHAC	RYLONITRII	LE 00012	26-98-7					
NOAEL.	3.2 PPM	DOG						
	INHALATION INTERMITTE		LIVER LIVER CENTRAL NERVOL SYSTEM BRAIN	INCREASED SGOT INCREASED SGPT US LOSS OF HINDLIMB MOTOR CONT	TROL	1E-3 300	IRIS	005812
	SUBCHRONIC CHRONIC [R1	[RfD] COMMENT: BASE	D ON ROUTE TO RO E HEAST TABLE 2:	: ALTERNATE METHODS - SUBCHRON!				
METHAN	OL	000067-5	6-1					
NOEL	500 MG/KG/DA	Y RAT						
	ORAL: GAVA	AGE 90 DAYS	BLOOD BLOOD BRAIN	INCREASED ALKALINE PHOSPHAT INCREASED SGPT DECREASED WEIGHT	TASE	5E+0 100	IRIS	010271
METHOM	IYL	016752-7	7-5					
NOEL	2.5 MG/KG/DA ORAL: DIE1		KIDNEY	LESIONS		2.5E-2 100	IRIS	005802
	SUBCHRONIC	[RfD] COMMENT: THE	CHRONIC ORAL Rfi	WAS ADOPTED AS THE SUBCHRONIC	C ORAL [RfD].			
METHOX	YCHLOR	00007	2-43-5					
NOEL	5.01 MG/KG/D		- / -					
	ORAL: GAVA		REPRODUCTION	LOSS OF LITTERS		5E-3 1000	IRIS	010357
	SUBCHRONIC	[RfD] COMMENT: THE	CHRONIC ORAL Rfi	WAS ADOPTED AS THE SUBCHRONIC	C ORAL [RfD].			
	CHRONIC Rf0	C COMMENT: THE CHRON	IC INHALATION R	fC IS CONSIDERED NOT VERIFIABLE	E (11/07/91) B	Y THE RfD/R	IRIS FC WORK GROUP.	010909

Subchronic

Chronic

CHEMICAL LEVEL

DOSE SPECIES ROUTE EXPERIMENT LENGTH

CRITICAL EFFECT

[RfC] [RfD] (mg/cu m) (mg/kg/day) [RfC] [RfD]

(mg/cv m) (mg/kg/day) UF

METHOXYETHANOL ACETATE, 2-000110-49-6

010497

REFERENCE

March 1994

CHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) SUBCHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

**METHOXYETHANOL, 2-**

000109-86-4

TARGET

NOAEL 93 MG/CU M

RABBIT

13 WEEKS

TESTICLE **EFFECTS**  2E-1

IRIS

010372

INHALATION: INTERMITTENT

SUBCHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). CHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

METHYL ACETATE

000079-20-9

LIVER

LIVER

NOEL 1156 MG/KG/DAY

RAT

ORAL: GAVAGE 90 DAYS INCREASED ALKALINE PHOSPHATASE

INCREASED SGPT

1E+1 100

010002 1E+0

1000

CHRONIC [RfD] COMMENT: CALCULATED FROM DATA OBTAINED WITH METHANOL BY MULTIPLYING BY THE MOLECULAR WEIGHT RATIO (74.08/32.04).

**METHYL ACRYLATE** 

000096-33-3

010498

CHRONIC [RfD] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) GENERAL COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

METHYL CHLOROCARBONATE

000079-22-1

CHRONIC [RfD] COMMENT: WITHDRAWN FROM IRIS (05/01/89).

GENERAL COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

March 1994

Subchronic Chronic CHEMICAL [RfC] REFERENCE DOSE SPECIES [RfC] [RfD] [RfD] LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF UF UF

METHYL ETHYL KETONE

000078-93-3

NOAEL 1711 MG/KG/DAY

RAT ORAL: DRINKING MULTI-FETUS WATER **GENERATION** 

DECREASED BIRTH WEIGHT

2E+0 1000 IRIS 010853

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS MODIFIED TO ESTIMATE THE SUBCHRONIC ORAL [RfD].

GENERAL COMMENT: MULTI-GENERATION DEVELOPMENTAL STUDY PERFORMED WHT THE SURROGATE 2-BUTANOL, A METABOLITE OF METHYL ETHYL KETONE.

NOAEL 1010 PPM

MOUSE

INHALATION: 10 DAYS **FETUS** 

KIDNEY

DECREASED BIRTH WEIGHT

1E+0 3000

IRIS

010845

INTERMITTENT

SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RfC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].

METHYL ETHYL KETONE PEROXIDE 001338-23-4

CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/22/93) BY THE RfD/RfC WORK GROUP.

010948

METHYL ISOBUTYL KETONE 000108-10-1

NOAEL 250 MG/(KG-DAY)

ORAL: GAVAGE

RAT

13 WEEKS

WHOLE BODY **LETHARGY** LIVER

INCREASED RELATIVE WEIGHT IN FEMALES

8E-1 300

8E-2 3000

010949

LIVER

INCREASED ABSOLUTE WEIGHT IN

FEMALES

KIDNEY INCREASED RELATIVE WEIGHT IN **FEMALES** 

> INCREASED ABSOLUTE WEIGHT IN FEMALES

INCREASED URINARY PROTEIN KIDNEY

LEVELS IN FEMALES

SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) CHRONIC [Rfc] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) CHRONIC [RfD] COMMENT: WITHDRAWN FROM IRIS (03/01/91), UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.

METHYL ISOCYANATE

000624-83-9

IRIS

010013

CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (12/18/90) BY THE RFD/RFC WORK GROUP.

<u>CHEMICAL</u> LEVEL		<u>PECIES</u> MENT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
	MERCURY 0.003 Mg Hg/kg/DA		67-92- <b>6</b>					AAFF/7
	ORAL		CENTRAL NERVOL SYSTEM	JS EFFECTS		3E-4 10	IRIS	005547
	SUBCHRONIC [RfD] CHRONIC [RfD] CO	COMMENT: THE MMENT: THE ADI	CHRONIC ORAL REG MINISTERED DOSE V	WAS ADOPTED AS THE SUBCHRONIC WAS CALCULATED FROM 200 NG HG/M	ORAL [RfD].			
METHYL	METHACRYLAT	E 00	0080-62-6					
NOEL	7.5 MG/KG/DAY ORAL: WATER	RAT 24 MONTHS	KIDNEY	INCREASED RELATIVE WEIGHT		8E-2 100	<b>8</b> E-2 100	010014
METHYL	PARATHION	0002	98-00-0				IRIS	010015
NOAEL	2.5 PPM	RAT		OUG THEOTERICE THURSTEIN		2E-3	IKIS	010846
	ORAL: DIET	90 DAYS	ERYTHROCYTES	CHOLINESTERASE INHIBITION		100		010040
METHYL	STYRENE (MIXE	D ISOMERS)	025013-15-4					
	CHRONIC [RfD] CO			HEAST TABLE 2: ALTERNATE METHO	DS SUBCHRO	ONIC AND CHR	ONIC TOXICITY (OTHER T	010500 HAN
	GENERAL COMMENT	CARCINOGENI : ALSO SEE HEA	ST TABLE 2: ALTE	RNATE METHODS SUBCHRONIC AND	CHRONIC TOX	ICITY (OTHER	THAN CARCINOGENICITY)	
METHYL	STYRENE, ALPH	1A 000	0098-83-9					
	GENERAL COMMENT	: ALSO SEE HEA	ST TABLE 2: ALTE	RNATE METHODS SUBCHRONIC AND	CHRONIC TOX	ICITY (OTHER	THAN CARCINOGENICITY)	010499

CHEMICAL LEVEL		PECIES MENT LENGTH	TARGET	CRITICAL EFFECT	Subchroni [RfC] [RfD] (mg/cu m) (mg/kg/da UF UF	[RfC] [RfD]	REFERENCE
METHYL-	4-CHLOROPHEN	IOXY) BUTYF	RIC ACID, 4-	2- 000094-81-5			
NOEL	12 MG/KG/DAY ORAL: DIET	RAT 13 WEEKS	171/50		a= a	****	24222
	ORAL: DIEI	12 MEEK2	LIVER KIDNEY	EFFECTS EFFECTS	1E-1 100	IRIS	010008
		DOG					
	ORAL: DIET	13 WEEKS	LIVER KIDNEY	EFFECTS EFFECTS			
METHYL-	4-CHLOROPHEN	IOXY) PROPI	ONIC ACID.	2-(2- 000093-65-2			
	3 MG/KG/DAY	RAT		_ (_ 000000			
	ORAL: DIET	90 DAYS	KIDNEY	ALTERED WEIGHT	1E-2 300	IRIS	010009
METHYL-	4-CHLOROPHEN	IOXYACETIC	ACID, 2- (	000094-74-6			
NOEL	0.15 MG/KG/DAY	DOG	·				
	ORAL: DIET	52 WEEKS	KIDNEY LIVER	EFFECTS EFFECTS	5E-4 300	IRIS	010007
	SUBCHRONIC [RfD]	COMMENT: THE	CHRONIC ORAL	RFD WAS ADOPTED AS THE SUBCHRON	IC ORAL [RfD].		
METHYLO	CYCLOHEXANE	000	108-87-2				
NOAEL	287 MG/CU M	RAT	WIR HEY	MINERAL ITATION	75.0	75.0	040/74
	INHALATION: INTERMITTENT	1 YEAR	KIDNEY	MINERALIZATION PAPILLARY HYPERPLASIA	3E+0 100	3E+0 100	010431
METHYLI	ENE BROMIDE	0000	74-95-3				
	GENERAL COMMENT:	ALSO SEE HEA	ST TABLE 2: AL	TERNATE METHODS SUBCHRONIC A	ND CHRONIC TOXICITY (OT	HER THAN CARCINOGENICITY	010501

CHEMICAL LEVEL		SPECIES IMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	ochronic [RfD] [Rf <u>a/kg/day) (mg/</u> Uf U	cu m) (mg/kg/day)	REFERENCE
METHYLE	NE CHLORIDE	/ (DICHLOROM	METHANE)	000075-09-2				
NOAEL	5.85 MG/KG/DAY ORAL: DRINKING WATER	RAT 24 MONTHS	LIVER	TOXICITY		6E-2 100	IRIS	005553
	SUBCHRONIC (RfC	) COMMENT: THE	CHRONIC ORAL RE	D WAS ADOPTED AS THE SUBCH	RONIC ORAL [RfD].			
NOAEL	694.8 MG/CU M INHALATION: INTERMITTENT	RAT 2 YEARS	LIVER	TOXICITY	3E+0 100		E+0 00	005552
	GENERAL COMMENT	T: ALSO SEE HEAS	ST TABLE 3: CARC	INOGENICITY.				
	ENE-BIS(2-CHL)	OROANILINE),	4,4'- 000101	-14-4				
LONEL	ORAL	9 YEARS	LIVER BLADDER	EFFECTS EFFECTS		7E-4 10000	7E-4 10000	010413
	CHRONIC [RfC]	COMMENT: THE CHI	RONIC INHALATION	[RfC] IS CONSIDERED NOT V	/ERIFIABLE 02/10/93)	BY THE RfD/RfC	WORK GROUP.	010933
METHYLI	ENEDIPHENYL I	SOCYANATE,	4,4- / (DIPEHN	NYLMETHANE DIISOCY	ANATE) 000101	1-68-8		
NOAEL	0.2 MG/CU M INHALATION: INTERMITTENT	RAT 24 MONTHS	NASAL CAVITY	LESIONS	2E-5 300		E-5 00	010449
	CHRONIC [RfC]	COMMENT: UNDER I	REVIEW, CURRENT	NUMBER SUBJECT TO CHANGE.				
METOLA	CHLOR	051218	3-45-2					
NOAEL	300 PPM ORAL: DIET	RAT 2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		1.5E-1 100	1.5E-1 100	010950
	SUBCHRONIC [Rf	D] COMMENT: THE	CHRONIC ORAL [R	TELEFORM THE SUCCESSION OF THE	CHRONIC ORAL [RfD].			

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<u>CHEMICAL</u> LEVEL		<u>SPECIES</u> IMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chronic (RfC) [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
METRIBU	ZIN	021087-6	4-9					
NOAEL	100 PPM	DOG						
	ORAL: DIET	2 YEARS	LIVER KIDNEY WHOLE BODY WHOLE BODY	EFFECTS EFFECTS MORTALITY DECREASED WEIGHT		2.5E-2 100	IRIS	010928
	SUBCHRONIC [Rf	D] COMMENT: THE	CHRONIC ORAL RE	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].			
MIREX		002385-85-	5					
	0.07 MG/KG/DAY	RAT						
	ORAL: DIET	2 YEARS	LIVER LIVER LIVER THYROID	CYTOMEGALY FATTY METAMORPHOSIS ANGIECTASIS CYSTIC FOLLICLES		2E - 4 300	IRIS	010841
	SUBCHRONIC [Rfi GENERAL COMMENT	D] COMMENT: THE T: ALSO SEE HEA	CHRONIC ORAL RFE ST TABLE 3: CARC	D WAS ADOPTED AS THE SUBCHRONIC INOGENICITY.	ORAL [RfD].			
MOLINAT	E	002212-6	7-1					
NOEL	0.2 MG/KG/DAY ORAL: GAVAGE	RAT	REPRODUCTIVE SYSTEM	TOXICITY		2E-3 100	IRIS	010017
	SUBCHRONIC [Rf0 CHRONIC [RfD] (	)] COMMENT: THE COMMENT: BASED	CHRONIC ORAL RFE ON A REPRODUCTION	WAS ADOPTED AS THE SUBCHRONIC STUDY.	ORAL [RfD].			
MOLYBDE	ENUM .	007439	9-98-7					
LOAEL	0.14 MG/KG/DAY	HUMAN						
	ORAL: WATER, DIET		URINE JOINTS BLOOD	INCREASED URIC ACID PAIN, SWELLING DECREASED COPPER LEVELS		5E-3 30	IRIS	010489

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RFD].

Subchronic Chronic CHEMICAL [RfC] [RfC] (RfD) REFERENCE DOSE ROUTE SPECIES [RfD] LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) MONOCHLORAMINE 010599-90-3 NOAEL 9.5 MG/KG/DAY PAT 010517 2 YEARS WHOLE BODY 1E-1 IRIS ORAL: DRINKING WEIGHT CHANGES 100 WATER LIVER WEIGHT CHANGES WEIGHT CHANGES KIDNEY SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. **NAPHTHALENE** 000091-20-3 CHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. NAPHTHOQUINONE, 1,4-000130-15-4 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 010020 **NICKEL (METALLIC)** 007440-02-0 NOAEL 100 PPM RAT IRIS 005579 2 YEARS WHOLE BODY DECREASED WEIGHT 2E-2 ORAL: DIET ORGANS, MAJOR DECREASED WEIGHT 300 SUBCHRONIC (RfD) COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. 000557-19-7 **NICKEL CYANIDE** 010953 CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/20/93) BY THE RfD/RfC WORK GROUP. **NICOTINONITRILE** 000100-54-9 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005584 **NITRIC OXIDE** 010102-43-9 CHRONIC (RfC) COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS. 010451

March 1994

Subchronic Chronic CHEMICAL [RfC] [RfD] [RfC] [RfD] REFERENCE DOSE SPECIES LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF

**NITRITE** 

014797-65-0

NOEL 10 PPM HUMAN

ORAL: WATER **BLOOD METHEMOGLOBINEMIA**  1E-1 10

010021

IRIS

SUBCHRONIC [RfD] COMMENT: THIS VALUE IS BASED ON NITRATE (NITROGEN) DATA FROM THE MOST SENSITIVE HUMAN POPULATION (INFANTS). THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RFD].

CHRONIC [RfD] COMMENT: THIS VALUE IS BASED ON NITRATE (NITROGEN) DATA FROM THE MOST SENSITIVE HUMAN POPULATION (INFANTS).

**NITROANILINE, 2-**

000088-74-4

CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RFD] IS CONSIDERED NOT VERIFIABLE (06/23/92) BY THE RFD/RFC WORK GROUP.

010936

LOAEL 9.8 MG/CU M

INHALATION:

4 WEEKS INTERMITTENT

BLOOD **HEMATOLOGICAL EFFECTS**  2E-3 1000 2E-4 10000 010935

NITROANILINE, M-

000099-09-2

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT

010400

NITROANILINE, P-

000100-01-6

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT

010024

March 1994

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route i	<u>spec</u> Experimen	CIES IT LENGTH	TARGET	CRITICAL EFFECT	[RfC] (mg/cu m) (i UF	ubchronic [RfD] mg/kg/day) UF	Chronic [RfC] [R (mg/cu m) (mg/k UF U	
NITROBE	NZENE		000098-9	95-3					
_	25 MG/CU M		MOUSE						
25/142	INHALATIC INTERMITT		90 DAYS	BLOOD ADRENAL KIDNEY LIVER	HEMATOLOGICAL EFFECTS LESIONS LESIONS LESIONS		5E-3 1000	IR	RIS 005589
			RAT						
	INHALATIO INTERMITT		90 DAYS	BLOOD ADRENAL KIDNEY LIVER	HEMATOLOGICAL EFFECTS LESIONS LESIONS LESIONS				
	SUBCHRONIC CHRONIC [R	[RfD] C	OMMENT: BASE ENT: ALSO SE	D ON ROUTE TO RO E HEAST TABLE 2: L RFD, WHILE STI	2: ALTERNATE METHODS - SUBCHROUTE EXTRAPOLATION. ALTERNATE METHODS SUBCHRONIC / LL AVAILABLE ON IRIS, IS BEING	AND CHRONICTOX	CICITY (OTHE	ER THAN CARCINOGE	NICITY)
NITROFU	RANTOIN		000067	-20-9					
	300 PPM		MOUSE						
	ORAL: DIE	E <b>T</b>	13 WEEKS	TESTIS	DAMAGE		7E-1 100		-2 005593 000
NITROGE	N DIOXIDE	:	010102	-44-0					
	10 PPM	-	HUMAN						
	ORAL: WAT	ER		BLOOD	METHEMOGLOBINEMIA		1E+0 1	18	ers 010402
			ORAL RFD WAS	ADOPTED AS THE	ON NITRATE (NITROGEN) DATA FROM SUBCHRONIC ORAL [RfD]. NITRATE (NITROGEN) DATA FROM TI			-	•
	CHRONIC [R	RFC] COMM	ENT: REFER T	O APPENDIX A: TÉ	CHNICAL INFORMATION, SECTION V	ON NATIONAL A	MBIENT AIR	QUALITY STANDARD	010912 9 <b>s.</b>

CHRONIC (RfC) COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS.

**NITROGEN OXIDES** 

010170

Subchronic Chronic CHEMICAL LEVEL DOSE ROUTE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE EXPERIMENT LENGTH **TARGET** (mg/cu m) (mg/kg/day) CRITICAL EFFECT (mg/cu m) (mg/kg/day) UF

NITROMETHANE 000075-52-5

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 010026

**NITROPHENOLS** 

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005594

NITROPROPANE, 2- 000079-46-9

LOAEL 78 MG/CU M RAT

INHALATION: 22 MONTHS LIVER LESIONS 2E-2 IRIS 010374
INTERMITTENT 1000

1000

SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RFC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RFC].

NITROSODIPHENYLAMINE, P- 000156-10-5

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 010027

NITROTOLUENE, M- 000099-08-1

LOAEL 200 MG/KG/DAY RAT

ORAL: GAVAGE 6 MONTHS SPLEEN LESIONS 1E-1 1E-2 010029
1000 10000

SUBCHRONIC [RfD] COMMENT: BASED ON DATA OBTAINED WITH O-NITROTOLUENE. CHRONIC [RfD] COMMENT: BASED ON DATA OBTAINED WITH O-NITROTOLUENE.

NITROTOLUENE, O- 000088-72-2

LOAEL 200 MG/KG/DAY RAT

ORAL: GAVAGE 6 MONTHS SPLEEN LESIONS 1E-1 1E-2 010028

Subchronic Chronic CHEMICAL DOSE SPECIES [RfC] [RfC] [RfD] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF NITROTOLUENE, P-000099-99-0 LOAEL 200 MG/KG/DAY RAT ORAL: GAVAGE 6 MONTHS SPLEEN LESIONS 1E-1 1E-2 010030 1000 10000 SUBCHRONIC [RfD] COMMENT: BASED ON DATA OBTAINED WITH O-NITROTOLUENE. CHRONIC [RfD] COMMENT: BASED ON DATA OBTAINED WITH O-NITROTOLUENE. OCTABROMODIPHENYL ETHER 032536-52-0 NOAEL 2.5 MG/KG/DAY ORAL: GAVAGE 90 DAYS LIVER HISTOLOGICAL CHANGES 3E-2 010032 IRIS 100 OCTAMETHYLPYROPHOSPHORAMIDE 000152-16-9 NOAEL 0.02 MG/KG/DAY HUMAN ORAL AT LEAST 30 BLOOD DECREASED CHOLINESTERASE 2E-3 010031 2E-3 DAYS ACTIVITY 10 10 OSMIUM TETROXIDE 020816-12-0 010954 CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/22/93) BY THE RfD/RfC WORK GROUP. OZONE 010028-15-6 CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS. 010171 PARALDEHYDE 000123-63-7 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 010033 **PARATHION** 000056-38-2 NOAEL 0.064 MG/KG/DAY HUMAN CHOLINESTERASE DECREASED CHOLINESTERASE 6E-3 005598 ORAL 6E-3 ACTIVITY

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

March 1994 Subchronic Chronic <u>Dose</u> Route [RfC] REFERENCE CHEMICAL **SPECIES** [RfC] [RfD] [RfD] LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF PARTICULATE MATTER CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS. 010034 **PEBULATE** 001114-71-2 NOEL 5 MG/KG/DAY RAT 010036 ORAL: DIET SUBCHRONIC BLOOD INCREASED CLOTTING TIME 5E-2 5E-2 100 100 PENDIMETHALIN 040487-42-1 NOEL 12.5 MG/KG/DAY DOG 010037 ORAL: CAPSULE 2 YEARS LIVER **EFFECTS** 4E-2 IRIS 300 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. PENTABROMODIPHENYL ETHER 032534-81-9 NOAEL 1.8 MG/KG/DAY RAT 010038 2E-2 IRIS ORAL: GAVAGE 90 DAYS LIVER ALTERED ENZYME ACTIVITIES 100 PENTACHLOROBENZENE 000608-93-5 LOAEL 8.3 MG/KG/DAY RAT IRIS 010039 8E-3 ORAL: DIET 100 DAYS LIVER TOXICITY 1000 KIDNEY TOXICITY PENTACHLOROCYCLOPENTADIENE 025329-35-5 005302 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. **PENTACHLORONITROBENZENE** 000082-68-8 NOEL 0.75 MG/KG/DAY DOG 010040 3E-3 IRIS ORAL: DIET 2 YEARS LIVER TOXICITY 300

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

March 1994

REFERENCE

Subchronic Chronic CHEMICAL DOSE SPECIES

[RfC] [RfD] [RfC] [RfD] ROUTE EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day)

PENTACHLOROPHENOL 000087-86-5

NOEL 3 MG/KG/DAY RAT

> ORAL: GAVAGE 62 DAYS FETUS **FETOTOXICITY** 3E-2 IRIS 005600 100

SUBCHRONIC [RfD] COMMENT: BASED ON A TERATOLOGY STUDY WITH EXPOSURE 62 DAYS PRIOR TO MATING AND THROUGHOUT GESTATION AND LACTATION.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

PENTACHLOROPROPENE, 1,1,2,3,3,- 001600-37-9

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 010041

PENTANE, N-000109-66-0

> GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005603

PHENANTHRENE 000085-01-8

> GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005604

**PHENOL** 000108-95-2

> NOAEL 60 MG/KG/DAY RAT

> > ORAL: GAVAGE FETUS DECREASED WEIGHT 6E-1 IRIS 005824 100

SUBCHRONIC [RfD] COMMENT: BASED ON A DEVELOPMENTAL STUDY WITH EXPOSURES DURING DAYS 6-15 OF GESTATION. THE ORAL CHRONIC RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfD] COMMENT: BASED ON A DEVELOPMENTAL STUDY WITH EXPOSURES DURING DAYS 6-15 OF GESTATION.

IRIS 010913

CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (02/22/90) BY THE RFD/RFC WORK GROUP.

PHENYLENEDIAMINE, M-000108-45-2

> NOEL 6 MG/KG/DAY RAT

> > 010044

ORAL 90 DAYS LIVER LESIONS 6E-2 IRIS 100

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CHEMICAL LEVEL

DOSE ROUTE

**SPECIES** EXPERIMENT LENGTH

TARGET

CRITICAL EFFECT

Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day)

TIF

Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day)

UF

REFERENCE

PHENYLENEDIAMINE, O-

000095-54-5

000106-50-3

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

010042

PHENYLENEDIAMINE, P-

NOAEL 18.7 MG/KG/DAY

RAT

2 YEARS

WHOLE BODY **EFFECTS** 

1.9E-1 100

010043

PHENYLMERCURIC ACETATE

ORAL: DIET

000062-38-4

NOAEL 0.0084 MG/KG/DAY ORAL: DIET

2 YEARS

KIDNEY

DAMAGE

8E-5 100

IRIS

010277

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. CALCULATED BY ANALOGY TO MERCURY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO MERCURY BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

**PHORATE** 

000298-02-2

NOAEL 0.033 MG/KG/DAY

ORAL: DIET

13 WEEKS CHOLINESTERASE INHIBITION 2E-4 200

2E-4 200

010403

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

**PHOSGENE** 

000075-44-5

IRIS

010045

CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (10/01/90) BY THE RFD/RFC WORK GROUP.

<u>CHEMICAL</u> LEVEL		<u>PECIES</u> Ment Length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
PHOSPHI	NE	007803-5	1-2				
NOEL	0.026 MG/KG/DAY ORAL: DIET	RAT 2 YEARS			3E-4 100	IRIS	010174
	SUBCHRONIC [RfD]	COMMENT: THE	CHRONIC ORAL R	FD WAS ADOPTED AS THE SUBCH	RONIC ORAL [RfD].		
NOAEL	1.4 MG/CU M INHALATION: INTERMITTENT	RAT 24 WKS	KIDNEY	RENAL EFFECTS	3E-4 1000	3E-5 10000	010173
PHOSPHO	DRUS, WHITE	0077	/23-14-0				
	GENERAL COMMENT:	FORMERLY LIS	TED AS PHOSPHORU	JS (INORGANIC COMPOUNDS).		IRIS	010452
РНОТОСЬ	HEMICAL OXIDA	ANTS					
			ATE FOR QUANTITA	ATIVE RISK ASSESSMENT.			010172
PHTHALI	C ACID, M- GENERAL COMMENT:	OOO12 DATA INADEQU		ATIVE RISK ASSESSMENT			010047
PHTHALIC	C ACID, O- GENERAL COMMENT:	380000 DATA INADEQU		ATIVE RISK ASSESSMENT			010046
PHTHALI	C ACID, P-	000100	-21-0				
NOEL	142 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	BLADDER	HYPERPLASIA	1E+0 100	1E+0 100	010048

March 1994

<u>CHEMICAL</u> LEVEL	DOSE ROUTE EXPE	<u>SPECIES</u> Riment Length	TARGET	CRITICAL EFFECT	Subchronid (RfC) (RfD) (mg/cu m) (mg/kg/day UF UF	[RfC] [RfD]	REFERENCE
PHTHALI	C ANHYDRIDI	0000	85-44-9				
LOAEL	1562 MG/KG/DAY ORAL: DIET	MOUSE 104 WEEKS	LUNG KIDNEY	HISTOPATHOLOGY HISTOPATHOLOGY	2E+0 1000	IRIS	010049
	SUBCHRONIC [R	FD] COMMENT: THE	CHRONIC ORAL RE	D WAS ADOPTED AS THE SUBCHR	ONIC ORAL [RfD].		
LOAEL	0.1 MG/CU M INHALATION: INTERMITTENT	HUMAN 12 YEARS	NOSE Lungs	RHINITIS Bronchitis	1.2E-1 300	1.2E-1 300	010847
	SUBCHRONIC [R	FC] COMMENT: THE	CHRONIC INHALAT	ION [RfC] WAS ADOPTED AS TH	E SUBCHRONIC INHALATION [R	fC].	
POLYBRO	MINATED BIP	HENYLS					
LOAEL	0.07 MG/KG/DAY ORAL: GAVAGE	RAT 25 Weeks	LIVER LIVER	INCREASED WEIGHT LESIONS	7E-5 1000	7E-6 10000	010050
	GENERAL COMME	IT: ALSO SEE HEAS	ST TABLE 3: CARC	INOGENICITY.			
POTASSI	JM CYANIDE	0001	51-50-8				
NOAEL	27 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	WHOLE BODY THYROID NERVE	DECREASED WEIGHT EFFECTS Myelin Degeneration	5E-2 500	IRIS	010278

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. THE ORAL CHRONIC RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

<u>CHEMICAL</u> LEVEL	<u>pose</u> Route		<u>PECIES</u> IENT LENGTH	TARGET	CRITICAL EFFECT	(Rfc) (mg/cu m) (n Uf	ubchronic (RfD) ng/kg/day) UF	Chroni [RfC] (mg/cu m) (m UF	[RfD]	REFERENCE
POTASSI			NIDE	000506-61-6						
NOAEL	82.7 MG/K ORAL: D		RAT 2 YEARS	WHOLE BODY THYROID NERVE	DECREASED WEIGHT EFFECTS MYELIN DEGENERATION		2E-1 500		IRIS	010279
			WAS ADOP	TED AS THE SUBCHRON	GY TO FREE CYANIDE BY CORRECT IC ORAL [Rf0]. TO FREE CYANIDE BY CORRECTING				THE ORAL	CHRONIC RfD
PROFLUR	ALIN		02639	9-36-0						
NOEL 3	3 MG/KG/DA ORAL: [		RAT Subchron	ıc	NONE OBSERVED		6E-3 500		6E-3 500	010051
PRONAM	IDE		02395	0-58-5						
NOEL	7.5 MG/KG ORAL: [	•	DOG 2 YEARS		NONE OBSERVED		7.5E-2 100		IRIS	010280
	SUBCHRO	NIC [RfD]	COMMENT:	THE CHRONIC ORAL RE	D WAS ADOPTED AS THE SUBCHRO	NIC ORAL [RfD].				
PROPACE	HLOR		0019	18-16-7						
NOEL	13.3 MG/K ORAL: I		RAT 90 days	WHOLE BODY	DECREASED WEIGHT GAIN		1.3E-1 100		IRIS	0101 <i>7</i> 5
PROPAZII	NE		000139	9-40-2						
NOEL	5 MG/KG/D ORAL: I		RAT 2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		2E-2 300		IRIS	010052
	SUBCHRO	NIC [RfD]	COMMENT:	THE CHRONIC ORAL RE	D WAS ADOPTED AS THE SUBCHRO	NIC ORAL [RfD].				
PROPION	ITRILE		00010	7-12-0						
		COMMENT:			TIVE RISK ASSESSMENT					010053

		HEAD! IADEL	i. Subunituit	IIC AND CHRONIC TOX	ICH I (OTHER THAIF O	Anomogenion I,	Mai Cil 1774
<u>CHEMICAL</u> LEVEL	<u>dose</u> Route expe	<u>SPECIES</u> RIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronio [RfC] [RfD] (mg/cu m) (mg/kg/da) UF UF	[RfC] [RfD]	REFERENCE
PROPYL	ALCOHOL, N- GENERAL COMME		71-23-8 ATE FOR QUANTITATI	VE RISK ASSESSMENT			005627
PROPYL	ENE GLYCOL	0000!	57-55-6				
NOEL	L 50000 PPM ORAL: DIET	DOG 2 YEARS	ERYTHROCYTES BLOOD BLOOD	DECREASED COUNT DECREASED HEMATOCRIT DECREASED HEMOGLOBIN		2E+1 100	005631
NOE	ORAL: DIET	RAT 20 WEEKS	KIDNEY	LESIONS	3E+1 100		005629
						IRIS	010914
	CHRONIC RTC C	OPMENT: THE CHRO	NIC INHALATION RFC	C IS CONSIDERED NOT VERIFI	ABLE (04/25/91) BY THE RfC	O/RTC WORK GROUP.	
PROPYL	ENE GLYCOL N	MONOETHYL ET	HER 001569-	-02-4			
NOEI	L 680 MG/KG/DAY ORAL: DRINKI WATER	RAT NG 30 DAYS	WHOLE BODY	DECREASED WEIGHT GAIN	7E+0 100	7E-1 1000	005488
	CHRONIC RFC C	OMMENT: THE CHRO	NIC INHALATION RFC	C IS CONSIDERED NOT VERIFI	ABLE (04/25/91) BY THE Rf0	IRIS D/RfC WORK GROUP.	010915
	LENE GLYCOL N L 947 mg/kg/day	AONOMETHYL I RAT	ETHER 00010	7-98-2			
NOC	ORAL: GAVAGE		LIVER KIDNEY	HISTOPATHOLOGY HISTOPATHOLOGY	7E+0 100	7E-1 1000	005486
NOAI	EL 1000 PPM INHALATION:	RAT, RABBIT 13 WEEKS	CENTRAL NERVOUS	S EFFECTS	2E+1 30	IRIS	010276
	INTERMITTENT		SYSTEM		30		

-			L	4	994
	a	rc	n	•	v

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route exp	<u>SPECIES</u> ERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
PROPYLE	NE OXIDE	000075	-56-9				
LOAEL	71 MG/CU M INHALATION: INTERMITTEN	RAT 2 YEARS T	EPITHELIUM	UNSPECIFIED	3E-2 100	IRIS	010375
		RFC] COMMENT: THE ENT: ALSO SEE HEAS		TION RFC WAS ADOPTED AS THE SUBCICINOGENICITY.	CHRONIC INHALATION [RFC].		
PYRENE		000129-00-0	0				
NOAEL	75 MG/KG/DAY ORAL: GAVAG	MOUSE E 13 WKS	KIDNEY	EFFECTS	3E-1 300	IRIS	010176
PYRIDINE		000110-86-	1				
NOAEL	1 MG/KG/DAY ORAL: GAVAG	RAT E 90 DAYS	LIVER LIVER	INCREASED WEIGHT INCREASED RELATIVE WEIGHT	1E-2 100	IRIS	010055
RDX / (CY	(CLONITE)	000121-	82-4				
NOEL	0.3 MG/KG/DAY ORAL	RAT 105 WEEKS	PROSTATE PROSTATE	INFLAMMATION HEMOSIDEROSIS	3E-3 100	IRIS	010056
		RfD] COMMENT: THE ENT: ALSO SEE HEAS		TO WAS ADOPTED AS THE SUBCHRONIC CINOGENICITY.	ORAL [RfD].		
RONNEL		000299-84-	3				
NOAEL	5 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	LIVER	EFFECTS	5E-2 100	5E-2 100	010057

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CHEMICAL LEVEL	<u>Dose</u> Route		<u>CIES</u> Ent Length	TARGET	CRITICAL EFFECT	[RfC] [Rf (mg/cu m) (mg/k	fD] [RfC]	hronic [RfD] <u>(mg/kg/day)</u> UF	REFERENCE
SELENIO L NOAEL	JS ACID 0.046 MG/K ORAL: DI		OO778 HUMAN	B3-00-8 WHOLE BODY	SELENOSIS, CLINICAL		5E-3 3	IRIS	010504
	SUBCHRON I	C [RfD]	COMMENT:	THE CHRONIC ORAL I	RFD WAS ADOPTED AS THE SUBCHR	ONIC ORAL [RfD].			
SELENIUN NOAEL	0.853 MG/D ORAL: DI	ET	007782- HUMAN	WHOLE BODY	SELENOSIS, CLINICAL		5E-3 3	IRIS	010404
	SUBCHRONI	IC [RfD]	COMMENT:	THE CHRONIC ORAL I	RFD WAS ADOPTED AS THE SUBCHI	ONIC ORAL [RfD].			
SELENOU NOAEL	REA 0.072 MG/K ORAL: DI		OOO63 HUMAN	O-10-4 WHOLE BODY	SELENOSIS		5E-3 15	5E-3 15	010473
	CHRONIC	[RfD] CO	MENT: WIT	HDRAWN FROM IRIS	(05/01/91). UNDER REVIEW, CU	RENT NUMBER SUBJECT T	O CHANGE.		
SILVER LOAEL	0.014 MG/K IV		007440-2 Human 2-9 Year:		ARGYRIA		5E-3 3	IRIS	010453

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. CHRONIC [RfD] COMMENT: BASED ON A TOTAL IV DOSE OF 1 GRAM.

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Subchronic Chronic CHEMICAL DOSE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) ÜF UF SILVER CYANIDE 000506-64-9 NOAEL 55.7 MG/KG/DAY RAT ORAL: DIET 2 YEARS WHOLE BODY DECREASED WEIGHT 1E-1 IRIS 010283 THYROID **EFFECTS** 500 NERVE MYELIN DEGENERATION

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. THE ORAL CHRONIC RFD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

SIMAZINE 000122-34-9

NOAEL 0.52 MG/(KG-DAY) RAT

ORAL: DIET 2 YEARS WHOLE BODY DECREASED WEIGHT GAIN 5E-3 IRIS 010955
BLOOD HEMATOLOGICAL EFFECTS 100

SUBCHRONIC [RfD] COMMENT: THE ORAL CHRONIC RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY

SODIUM CYANIDE 000143-33-9

NOAEL 20.4 MG/KG/DAY RAT

ORAL: DIET CENTRAL NERVOUS EFFECTS 4E-2 IRIS 005640 SYSTEM 500

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. THE ORAL CHRONIC RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

SODIUM DIETHYLDITHIOCARBAMATE 000148-18-5

NOEL 30 MG/KG/DAY RAT

ORAL 90 DAYS EYE CATARACTS 3E-1 IRIS 005644
WHOLE BODY DECREASED WEIGHT 100

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

SODIUM METAVANADATE 013718-26-8

NOAEL 10 PPM RAT

ORAL: DRINKING 3 MONTHS KIDNEY IMPAIRED FUNCTION 1E-2 1E-3 005735
WATER 100 1000

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Subchronic Chronic CHEMICAL **SPECIES** [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) STRONTIUM, STABLE 007440-24-6 NOAEL 190 MG/KG/DAY RAT, YOUNG ORAL: DRINKING 20 DAYS BONE RACHITIC CHANGES 6E-1 IRIS 010842 WATER 300 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. **STRYCHNINE** 000057-24-9 LOAEL 2.5 MG/KG/DAY RAT ORAL: GAVAGE 28 DAYS UNSPECIFIED TOXICITY 3E-3 IRIS 010285 UNSPECIFIED **HISTOPATHOLOGY** 1000 GENERAL COMMENT: THE LOAEL IS ALSO THE FEL. STYRENE 000100-42-5 IRIS 010059 SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. NOAEL 22 PPM HUMAN INHALATION: CENTRAL **EFFECTS** 3E+0 IRIS 010511 OCCUPATIONAL **NERVOUS SYSTEM** 10

CHRONIC [RfC] COMMENT: THE MEAN DURATION OF EXPOSURE FOR 50 WORKERS WAS 8.6 YEARS. AIR EXPOSURE CONCENTRATIONS WERE ESTIMATED FROM THE

SUMMATION OF THE PRINCIPLE URINARY METABOLITES OF STYRENE, MANDELIC ACID AND PHENYLGLYOXYLIC ACID. SEE IRIS FOR MORE INFORMATION.

INFORMATION.

GENERAL COMMENT: ALSO SEE TABLE 3: CARCINOGENICITY.

SUCCINONITRILE 000110-61-2

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005585

SULFUR DIOXIDE 007446-09-5

CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS. 010505

CHEMICAL LEVEL

ROUTE

SPECIES EXPERIMENT LENGTH

TARGET

CRITICAL EFFECT

[RfC] [RfD] (mg/cu m) (mg/kg/day)

Subchronic

[RfC] (mg/cu m) (mg/kg/day)

Chronic

[RfD] REFERENCE

**SULFUR OXIDES** 

CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS.

010035

**SULFURIC ACID** 

007664-93-9

NOAEL 0.066-0.098 MG/CU M HUMAN

INHALATION

RESPIRATORY SYSTEM

RESPIRATORY EFFECTS

005647

CHRONIC [RfC] COMMENT: REPORTED EFFECTS OCCURRED AT PORTAL OF ENTRY. ESTIMATES OF MG/DAY REFERENCE DOSES ARE INAPPROPRIATE BECAUSE EFFECTS AT PORTAL OF ENTRY DEPEND ON CONCENTRATION IN AIR. AN ACCEPTABLE AIR CONCENTRATION OF 0.07 MG/CU M WAS ESTIMATED BY CARSON ET AL (1981) FROM AVAILABLE DATA.

**TEMEPHOS** 

003383-96-8

NOAEL 200 PPM

RAT

ORAL: DIET 99 DAYS 2E-1 100

2E-2 010060 1000

**TERBUFOS** 

013071-79-9

NOAEL 0.0025 MG/KG/DAY

ORAL: DIET

6 MONTHS

DOG

CHOLINESTERASE INHIBITION

2.5E-5 100

2.5E-5 100

TEREPHTHALIC ACID

000100-21-0

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.

010474

010408

**TETRACHLOROAZOXYBENZENE** 

ORAL: DIET

021232-47-3

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT

010064

TETRACHLOROBENZENE, 1,2,4,5- 000095-94-3

NOAEL 0.34 MG/KG/DAY

RAT 13 WEEKS

KIDNEY

LESIONS

3E-3

100

IRIS

010286

<u>CHEMICAL</u> LEVEL		<u>SPECIES</u> IMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
TETRACH	ILOROCYCLOPI GENERAL COMMENT		000695-77-2 TE FOR QUANTITAT	2 TIVE RISK ASSESSMENT			005303
	ILOROETHANE, 89.3 Mg/kg/day oral: gavage	, 1,1,1,2- OG RAT 103 WEEKS	00630-20-6 LIVER KIDNEY	LESIONS LESIONS	3E-2 3000	IRIS	010407
		COMMENT: THE		D WAS ADOPTED AS THE SUBCHRONINGENICITY.	IIC ORAL [RfD].		
	1LOROETHYLEN 14 MG/KG/DAY ORAL	HOUSE 6 WEEKS	0127-18-4 LIVER	HEPATOTOXICITY	1E-1 100	IRIS	005650
TETRACH	ILOROHYDRAZ	OBENZENE	071753-42-9				010065
TETRACH	ILOROPHENOL, GENERAL COMMENT		04901-51-3 TE FOR QUANTITA	TIVE RISK ASSESSMENT			005324
	ILOROPHENOL, 25 mg/kg/day oral: gavage	, 2,3,4,6- 00 RAT 90 DAYS	00058-90-2 LIVER LIVER	INCREASED WEIGHT CENTRILOBULAR HYPERTROPH	3E-1 100	IRIS	005323
TETRACH	ILOROPHENOL, GENERAL COMMENT		00935-95-5 ATE FOR QUANTITA	TIVE RISK ASSESSMENT			005325
TETRACH	ILOROPROPENI GENERAL COMMENT			TIVE RISK ASSESSMENT			010066

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route exp	<u>species</u> Eriment Length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] [RfC] (mg/cu m) (mg/kg/day) (mg/cu UF UF UF	Chronic [RfD] REFERENCE <u>J m) (mg/kg/day)</u> UF
TETRACE	HLOROVINPH	OS / (STIROPHO	S) 000	0961-11-5		
NOEL	3.1 MG/KG/DAY ORAL: DIET	DOG 2 YEARS	LIVER KIDNEY WHOLE BODY	INCREASED WEIGHT INCREASED WEIGHT	3E-2 100	IRIS 010067
	SUBCHRONIC [F GENERAL COMME	fD] COMMENT: THE NT: ALSO SEE HEA	CHRONIC ORAL RES ST TABLE 3: CARC	D WAS ADOPTED AS THE SUBCHRONINGENICITY.	IC ORAL [RfD].	
TETRAET	HYL DITHIOP	YROPHOSPHAT	E 003689-24	-5		
NOEL	0.5 MG/KG/DAY ORAL: DIET	RAT 3 MONTHS	ERYTHROCYTES BLOOD	DECREASED CHOLINESTERASE ACTIVITY DECREASED CHOLINESTERASE ACTIVITY	5E-3 100	IRIS 010287
THALLIC		OO1314-COMMENT: THE CHI		IS CONSIDERED NOT VERIFIABLE	(07/22/93) BY THE RfD/RfC WORK GR	ROUP. 010956
	M (I) ACETAT	E 00056	63-68-8			
NOAEL	0.26 MG/KG/DAY ORAL	RAT 90 days	LIVER BLOOD HAIR	INCREASED SGOT INCREASED SERUM LDH ALOPECIA	9E-4 300	IRIS 005664
	SUBCHRONIC [R CHRONIC [RfD]	fD] COMMENT: CALC COMMENT: CALCULA	CULATED BY ANALOG ATED BY ANALOGY T	Y TO THALLIUM (I) SULFATE BY O THALLIUM (I) SULFATE BY COR	CORRECTING FOR MOLECULAR WEIGHT DER	IFFERENCES. ERENCES.
THALLIUI	M (I) CARBON	IATE 006	533-73-9			
	0.23 MG/KG/DAY ORAL	RAT 90 DAYS	LIVER BLOOD HAIR	INCREASED SGOT INCREASED SERUM LDH ALOPECIA	8E-4 300	IRIS 005668

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO THALLIUM (1) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO THALLIUM (1) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES.

March 1994

Subchronic Chronic CHEMICAL DOSE ROUTE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) THALLIUM (I) CHLORIDE 007791-12-0 NOAEL 0.23 MG/KG/DAY RAT ORAL 90 DAYS LIVER INCREASED SGOT 8E-4 IRIS 005672 BLOOD INCREASED SERUM LDH 300 HAIR ALOPECIA SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO THALLIUM (1) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO THALLIUM (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES. THALLIUM (I) NITRATE 010102-45-1 NOAEL 0.26 MG/KG/DAY RAT ORAL 90 DAYS LIVER INCREASED SGOT 9E-4 IRIS 005676 BLOOD INCREASED SERUM LDH 300 HAIR ALOPECIA SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO THALLIUM (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO THALLIUM (1) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES. THALLIUM (I) SULFATE 007446-18-6 NOAEL 0.25 MG/KG/DAY RAT ORAL 90 DAYS LIVER INCREASED SGOT 8E-4 IRIS 005682 BLOOD INCREASED SERUM LDH 300 HAIR **ALOPECIA** 

THALLIUM (IN SOLUBLE SALTS)

CHRONIC [RfD] COMMENT: REFER TO IRIS FOR OTHER THALLIUM SALTS.

010458

THALLIUM SELENITE 012039-52-0

010957

CHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS WITHDRAWN FROM IRIS (08/01/93). THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/22/93) BY THE RfD/RfC WORK GROUP.

Subchronic Chronic CHEMICAL DOSE ROUTE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF UF THIOCYANOMETHYLTHIO)BENZOTHIAZOLE, 2-( 021564-17-0 NOEL 25 MG/KG/DAY RAT ORAL: DIET SUBCHRONIC STOMACH LESIONS 3E-1 3E-2 010068 100 1000 SUBCHRONIC [RfD] COMMENT: BASED ON CONFIDENTIAL BUSINESS INFORMATION DATA CHRONIC [RfD] COMMENT: BASED ON CONFIDENTIAL BUSINESS INFORMATION DATA **THIOFANOX** 013196-18-4 NOAEL 0.025 MG/KG/DAY DOG ORAL 8 DAYS CHOLINESTERASE DECREASED CHOLINESTERASE 3E-4 010069 3E-4 ACTIVITY 100 100 SUBCHRONIC [RfD] COMMENT: BASED ON CONFIDENTIAL BUSINESS INFORMATION DATA. CHRONIC [RfD] COMMENT: BASED ON CONFIDENTIAL BUSINESS INFORMATION DATA. THIRAM 000137-26-8 010459 IRIS NOAEL 0.61 MG/KG/DAY FERRET ORAL 24 WEEKS REPRODUCTION 6E-3 010070 IMPAIRED 100 TIN AND COMPOUNDS NOAEL 2000 PPM RAT 005688 ORAL: DIET 2 YEARS LIVER LESIONS 6E-1 6E-1 100 100 KIDNEY LESIONS

SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO STANNOUS CHLORIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT. CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO STANNOUS CHLORIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

	<u>CIES</u> NT LENGTH TARGET	CRITICAL EFFECT	Subchro [RfC] [RfD] (mg/cu m) (mg/kg/ UF UF	[RfC] [RfD]	REFERENCE
TOLUENE	000108-88-3				
NOAEL 223 MG/KG/DAY ORAL: GAVAGE	RAT 13 WEEKS LIVER KIDNE	ALTERED WEIGHT ALTERED WEIGHT	2E- 100		010469
				IRIS	010844
SUBCHRONIC [Rfc]	COMMENT: CONTACT THE	SUPERFUND HEALTH RISK TECHNICAL SUPPORT	CENTER: (513) 569	-7300.	
TOLUENE-2,5-DIAMINE	000095-70-5				
NOAEL 56 MG/KG/DAY	RAT				
ORAL: DIET	78 WEEKS		6E- 100		010073
SUBCHRONIC [RfD] (CHRONIC [RfD] COM	COMMENT: DETERMINED   MENT: DETERMINED FROM	FROM DATA OBTAINED WITH THE SULFATE SALT I DATA OBTAINED WITH THE SULFATE SALT.	r <b>.</b>		
TOLUENE-2,6-DIAMINE	000823-40-5				
NOAEL 16 MG/KG/DAY	RAT				
ORAL: DIET	2 YEARS		2E- 100		010074
SUBCHRONIC [RfD] C CHRONIC [RfD] COM	COMMENT: DETERMINED   MENT: DETERMINED FROM	ROM DATA OBTAINED WITH THE DIHYDROCHLOR DATA OBTAINED WITH THE DIHYDROCHLORIDE	RIDE.		
TOLUENEDIAMINE, 2,3-	002687-25-4				
GENERAL COMMENT:	DATA INADEQUATE FOR	NUANTITATIVE RISK ASSESSMENT			010071
TOLUENEDIAMINE, 3,4-	000496-72-0	·			
		NUANTITATIVE RISK ASSESSMENT			010072
TOLUIDINE, M-	000108-44-1				
*		NUANTITATIVE RISK ASSESSMENT			010075

<u>CHEMICAL</u> LEVEL		<u>CCIES</u> NT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chroni [RfC] (mg/cu m) (mg UF	[RfD]	REFERENCE
TRIALLA	TE	002303-17-	-5						
NOAEL	1.275 MG/KG/DAY ORAL: DIET	DOG 24 MONTHS	SPLEEN LIVER	EFFECTS EFFECTS		1.3E-2 100		IRIS	010076
	SUBCHRONIC [RfD]	COMMENT: THE	CHRONIC ORAL RE	WAS ADOPTED AS THE SUBCHRONIC	ORAL [RfD].				
	IOBENZENE, 1,2,	4- 0006 RAT	15-54-3						
WORLE	ORAL: DIET	45 OR 90 DAYS	LIVER LIVER	ALTERED WEIGHT ENZYME INDUCTION		5E-2 100		IRIS	010077
TRICHLO	RO-1,2,2-TRIFLU	OROETHANE	, 1,1,2- 000	0076-13-1					
NOEL	2000 PPM INHALATION: INTERMITTENT	RAT 24 MONTHS	WHOLE BODY	DECREASED WEIGHT	3E+1 100	3E+0 100	3E+1 100	IRIS	010460 010376
	SUBCHRONIC [RfD]	COMMENT: BASE	ON ROUTE TO R	OUTE EXTRAPOLATION USING AN ABS	ORPTION FACTO	OR OF 0.2.			
	RO-2'-HYDROXY		HER, 2,2,4'-	003380-34-5					
NOEL	500 MG/KG/DAY ORAL	rat 4 weeks	WHOLE BODY	DECREASED WEIGHT		4E+0 100			005492
TRICHLO	ROBENZENE, 1,2	,4- 000°	120-82-1						
NOAEL	100 PPM ORAL: DRINKING WATER	RAT	ADRENAL	INCREASED WEIGHT		1E-2 1000		IRIS	010506
	SUBCHRONIC [RfD] COMMENT: BASED ON A MULTIGENERATION REPRODUCTION STUDY.								
NOAEL	104 PPM INHALATION	RAT, RABBIT, DOG, MONKEY 6 AND 26	LIVER	NON-ADVERSE WEIGHT CHANGES	2E+0 100		2E-1 1000		010958
		WEEKS			100		1000		

Subchronic
MICAL DOSE SPECIES FRACT FRACT

ic Chronic

March 1994

REFERENCE

CHEMICAL DOSE SPECIES [RfC] [RfC] [RfD] (RfC) [RfD]
LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) (mg/kg/day)

UF UF UF UF UF

TRICHLOROCYCLOPENTADIENE 077323-84-3

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005304

TRICHLOROETHANE, 1,1,1- 000071-55-6

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

TRICHLOROETHANE, 1,1,2- 000079-00-5

NOAEL 3.9 MG/KG/DAY MOUSE

ORAL: DRINKING 90 DAYS BLOOD CLINICAL CHEMISTRY ALTERATIONS 4E-2 IRIS 005702 WATER

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

TRICHLOROFLUOROMETHANE 000075-69-4

IRIS 005502

LOAEL 1000 MG/KG/DAY RAT

ORAL 6 WEEKS WHOLE BODY INCREASED MORTALITY 7E-1 005500

SUBCHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)
CHRONIC [RfC] COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

TRICHLOROPHENOL, 2.3.4- 015950-66-0

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005330

TRICHLOROPHENOL, 2,3,5- 000933-78-8

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005331

TRICHLOROPHENOL, 2,3,6- 000933-75-5

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005332

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Subchronic Chronic CHEMICAL SPECIES [RfC] [RfD1 [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF TRICHLOROPHENOL, 2.4.5-000095-95-4 NOEL 1000 PPM RAT ORAL: DIET 98 DAYS LIVER **HEPATOTOXICITY** 1E+0 IRIS 005329 KIDNEY **EFFECTS** 100 IRIS 010919 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (04/24/91) BY THE RFD/RFC WORK GROUP. TRICHLOROPHENOL, 2.4.6-000088-06-2 IRIS 010461 CHRONIC RFC COMMENT: THE CHRONIC INHALATION RFC IS CONSIDERED NOT VERIFIABLE (04/24/91) BY THE RFD/RFC WORK GROUP. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY. TRICHLOROPHENOL, 3,4,5-000609-19-8 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005333 TRICHLOROPHENOXY) PROPIONIC ACID, 2(2,4,5-000093-72-1 NOEL 0.75 MG/KG/DAY DOG ORAL: DIET 2 YEARS LIVER **HISTOPATHOLOGY** 8E-3 IRIS 010284 100 SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]. TRICHLOROPHENOXYACETIC ACID, 2,4,5- 000093-76-5 IRIS 010178 NOEL 10 MG/KG/DAY RAT ORAL: DIET 90 DAYS KIDNEY WEIGHT EFFECTS 1E-1 010179 LIVER WEIGHT EFFECTS 100 TRICHLOROPROPANE, 1,1,1-007789-89-1 GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005705

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route ex	<u>SPECIES</u> PERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
TRICHLO	ROPROPANE	, 1,1,2- 000	)598-77-6				
NOEL	100 MG/L	RAT					
	ORAL: DRINK WATER	KING 13 WEEKS	LIVER KIDNEY THYROID	HISTOPATHOLOGY HISTOPATHOLOGY HISTOPATHOLOGY	5E-2 300	IRIS	005708
TRICHLO	ROPROPANE	E, 1,2,2- 003	3175-23-3				
	GENERAL COM	MENT: DATA INADEQUA	TE FOR QUANTITA	TIVE RISK ASSESSMENT			005706
TRICHLO	ROPROPANE	E. 1.2.3- 000	0096-18-4				
	8 MG/KG/DAY	RAT	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				
	ORAL	120 DAYS	WHOLE BODY LIVER	TOXICITY LESIONS	6E-2 100	IRIS	005714
			KIDNEY	LESIONS	100		
			ERYTHROCYTES BLOOD	DECREASED COUNT DECREASED HEMATOCRIT			
			BLOOD	DECREASED HEMOGLOBIN			
	GENERAL COMM	IENT: ALSO SEE HEAS	ST TABLE 3: CAR	CINOGENICITY.			
TRICHLO	ROPROPENE	. 1.2.3- 000	096-19-5				
	18 MG/CU M	DOG					
	INHALATION: INTERMITTEN		EYE	IRRITATION	5E-3 100	5E-3 100	010078
				OUTE EXTRAPOLATION USING AN A E EXTRAPOLATION USING AN ABSO			
TDIOLUC	DOTO: 11515						
	ROTOLUENE 0.5 PPM	• • •	077-46-5				
LOAEL	ORAL: DIET	RAT 28 days	LIVER	LESIONS	5E-5		005335
			KIDNEY THYROID	LESIONS LESIONS	1000		
			INIKUIP	FE910M9			

CHEMICAL LEVEL		<u>ecies</u> Ent length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu_m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
TRICHLO	ROTOLUENE, AL	PHA,2,6- 0	02014-83-7				
LOAEL	0.5 PPM ORAL: DIET	RAT 28 DAYS	LIVER KIDNEY THYROID	LESIONS LESIONS LESIONS	5E-5 1000		005339
TRIFLUR	ALIN	001582-09	9-8				
	0.75 MG/KG/DAY	DOG					
	ORAL: DIET	12 MONTHS	LIVER BLOOD	INCREASED WEIGHT METHEMOGLOBINEMIA	7.5E-3 100	IRIS	010080
	SUBCHRONIC [RfD] GENERAL COMMENT:			D WAS ADOPTED AS THE SUBCHRONIC INOGENICITY.	ORAL [RfD].		
TRIMETH	IYLBENZENES						
		DATA INADEQUA	TE FOR QUANTITA	TIVE RISK ASSESSMENT			005727
TRINITRO	DBENZENE, 1,3,5	- 00009	99-35-4				
NOAEL	0.51 MG/KG/DAY	RAT					
	ORAL: WATER	16 WEEKS	SPLEEN	INCREASED WEIGHT	5E-4 1000	IRIS	010081
				A OBTAINED WITH 1,3-DINITROBENZ BTAINED WITH 1,3-DINITROBENZENE			
TRINITRO	PHENOLS						
	GENERAL COMMENT:	DATA INADEQUA	TE FOR QUANTITA	TIVE RISK ASSESSMENT			010082
TRINITRO	PHENYLMETHY	LNITRAMINE	000479-45-	8			
LOAEL	125 MG/KG/DAY	RABBIT			<b>.</b>	<b></b>	
	ORAL: GAVAGE	9 MONTHS	LIVER KIDNEY	HISTOPATHOLOGICAL EFFECTS HISTOPATHOLOGICAL EFFECTS	1E-1 1000	1E-2 10000	010377
			SPLEEN	HISTOPATHOLOGICAL EFFECTS			

					Alon I (Ollien Illian OA)	· · · · · · · · · · · · · · · · · · ·	ngi Cii 1774
<u>CHEMICAL</u> LEVEL		<u>ECIES</u> Ent length	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
TRINITROT	OLUENE, 2,4,6	- 000	118-96-7				
	.5 MG/KG/DAY	DOG	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				
	ORAL: GAVAGE		LIVER	EFFECTS	5E-4 1000	IRIS	010416
;	SUBCHRONIC [RfD] GENERAL COMMENT:	COMMENT: TI ALSO SEE HI	HE CHRONIC ORAL REAST TABLE 3: CAR	FD WAS ADOPTED AS THE SUBCH CINOGENICITY.	RONIC ORAL [RfD].		
URANIUM,	SOLUBLE SAL	TS					
			INTACT THE SUPERF	JND HEALTH RISK TECHNICAL S	SUPPORT CENTER: (513) 569-7300	•	
\/A\\A\\	_						
VANADIUM	-	007440	-62-2				
NOAEL 5	PPM ORAL: DRINKING WATER	RAT LIFETIME			7E-3 100	7E-3 100	005739
VANADIUM	1 PENTOXIDE	00	1314-62-1				
NOAEL 17	7.85 PPM	RAT					
	ORAL: DIET	LIFETIME			<b>9E-3</b> 100	IRIS	005743
!	SUBCHRONIC [RfD]	COMMENT: TI	HE CHRONIC ORAL R	FD WAS ADOPTED AS THE SUBCH	RONIC ORAL [RfD].		
VANADIUM	1 SULFATE	036	907-42-3				
NOAEL 2.	.24 MG/KG/DAY	RAT					
	ORAL: DRINKING WATER	LIFETIME			2E-2 100	2E-2 100	005747
VERNAM /	(VERNOLATE)	00	1929-77-7				
	MG/KG/DAY	RAT					
	ORAL: DIET		WHOLE BODY	DECREASED WEIGHT	1E-2 100	IRIS	010083

March 1994

Subchronic Chronic CHEMICAL DOSE SPECIES [RfC] [RfD] [RfC] [RfD] REFERENCE LEVEL ROUTE EXPERIMENT LENGTH **TARGET** CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF UF VINYL ACETATE 000108-05-4 NOAEL 100 MG/KG/DAY RAT ORAL: WATER 2 YEARS WHOLE BODY ALTERED WEIGHT 1F+0 1E+0 010417 KIDNEY ALTERED WEIGHT 100 100 NOAEL 176 MG/CU M MOUSE INHALATION: 104 WEEKS NASAL CAVITY LESIONS 2E-1 IRIS 010418 INTERMITTENT

SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RFC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].

VINYL CHLORIDE 000075-01-4

SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

VINYL-1-CYCLOHEXENE, 4- 000100-40-3

GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 010084

WARFARIN 000081-81-2

LOAEL 2 MG/DAY HUMAN

ORAL BLOOD INCREASED PROTHROMBIN TIME 3E-4 IRIS 010409

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

March 1994

CHEMICAL DOSE LEVEL

ROUTE

SPECIES EXPERIMENT LENGTH

TARGET

CRITICAL EFFECT

[RfC] [RfD] (mg/cu m) (mg/kg/day)

Subchronic

[RfC] [RfD] (mg/cu m) (mg/kg/day)

Chronic

REFERENCE

XYLENE, M-

000108-38-3

NOAEL 250 MG/KG ORAL: GAVAGE

103 WEEKS

CENTRAL NERVOUS HYPERACTIVITY

SYSTEM

WHOLE BODY DECREASED WEIGHT WHOLE BODY

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

2E+0 100

005755

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RfD/RfC WORK GROUP.

010920

**XYLENE, MIXTURE** 

001330-20-7

IRIS

010872

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RfD/RfC WORK GROUP.

010921

XYLENE, O-

000095-47-6

NOEL 250 MG/KG

RAT ORAL: GAVAGE

103 WEEKS CENTRAL NERVOUS HYPERACTIVITY

SYSTEM

WHOLE BODY DECREASED WEIGHT 2E+0 100

005751

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RfD/RfC WORK GROUP.

010922

XYLENE, P-

000106-42-3

SUBCHRONIC [RfD] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

010923

CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RfD/RfC WORK GROUP.

	SPECIES IMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE
ZINC (METALLIC)	007440-	36-6				
LOAEL 1.0 MG/KG/DAY ORAL: DIET SUPPLEMENT	HUMAN 10 WEEKS	BL000	DECREASED BLOOD ENZYME	3E-1 3	IRIS	010937
CHRONIC [RfD] C	COMMENT: THE CH	RONIC ORAL RFD W	AS ADOPTED AS THE SUBCHRONIC OR	AL [RfD].		
ZINC CYANIDE  NOAEL 24.3 MG/KG/DAY  ORAL: DIET	000557-2 rat 2 years	21-1 WHOLE BODY THYROID	DECREASED WEIGHT EFFECTS	5E-2 500	IRIS	010289
		NERVE	MYELIN DEGENERATION			
			GY TO FREE CYANIDE BY CORRECTING F			
ZINC PHOSPHIDE	001314	-84-7				
LOAEL 3.48 MG/KG/DAY ORAL: DIET	RAT 13 WEEKS	WHOLE BODY	DECREASED WEIGHT DECREASED FOOD INTAKE	3E-3 1000	IRIS	010290
ZINEB	012122-67-7					
LOAEL 25 MG/KG/DAY ORAL: DIET	RAT 2 YEARS	THYROID	HYPERPLASIA	5E-2 500	IRIS	010085

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].

March 1994

**ACENAPHTHENE** 

000083-32-9

010165 US EPA. 1989. MOUSE ORAL SUBCHRONIC STUDY WITH ACENAPHTHENE. STUDY CONDUCTED BY HAZELTON LABORATORIES, INC., FOR THE OFFICE OF SOLID WASTE, WASHINGTON, DC.

US EPA. 1989. RfD/RfC WORK GROUP.

**ACENAPTHYLENE** 

000208-96-8

005202 US EPA. 1987. HEALTH EFFECTS ASSESSMENT FOR ACENAPHTHYLENE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF EMERGENCY AND REMEDIAL RESPONSE, WASHINGTON, DC.

**ACEPHATE** 

030560-19-1

005833 CHEVRON CHEMICAL COMPANY. 1987. CONFIDENTIAL BUSINESS INFORMATION UNPUBLISHED DATA. MRID NO. 40504819

US EPA. 1984. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR ACEPHATE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON, DC.

US EPA. 1989. RfD/RfC WORK GROUP.

**ACETONE** 

000067-64-1

005204 US EPA. 1986. NINETY-DAY GAVAGE STUDY IN ALBINO RATS USING ACETONE. OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON, DC.

US EPA. 1988. UPDATED HEALTH EFFECTS ASSESSMENT FOR ACETONE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF EMERGENCY AND REMEDIAL RESPONSE, WASHINGTON, DC.

US EPA. 1986. RfD/RfC WORK GROUP.

**ACETONE CYANOHYDRIN** 

000075-86-5

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**CARBON MONOXIDE** 

000630-05-0

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CRESOL, O- / (2-METHYLPHENOL)

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**NICOTINONITRILE** 

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**NITRIC OXIDE** 

010102-43-9

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000100-01-6

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March 1994

## PROPYLENE GLYCOL MONOETHYL ETHER

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## HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOX!CITY (OTHER THAN CARCINOGENICITY)

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								SUDCIII OTTIC	CIII	Offic	
CHEMICAL	DOSE	SPECI	ES				[RfC]	[RfD]	[RfC]	[RfD]	REFERENCE
LEVEL	ROUTE	EXPERIMENT	<del></del>	TARGET	CRITICAL	EFFECT	(mg/cu m)	(mg/kg/day)	(mg/cu m)	(mg/kg/day)	
		270 20000					UF	UF	UF	UF	
ACETON	E CYANO	OHYDRIN	000	075-86-5			•				
NOEL	4.0 MG/K	G/DAY	RAT								
	INHALA	•	14 WEEKS	CENTRAL NERVOL	S EFFECTS	•	1E-1		1E-2		010432
	INTERM			SYSTEM			100		1000		
	SUBCHRO					-II, DOSE CONVERSIONS	ON HEAST)	. AN ERROR IN	THE UNCERT	AINTY FACTOR	THAT WAS
				IEED (1988) WAS							
	CHRONIC	[RfC] COMME	NT: 4E-2 MG,	/KG/DAY (SEE APF	PENDIX A-II	, DOSE CONVERSIONS ON	HEAST).				
	GENERAL	COMMENT: T	HE SUBCHRONI	AND CHRONIC IN	HALATION (	[RfC] VALUES WERE DERI	VED FROM MI	ETHODOLOGY THA	IT IS KOT C	JRRENT WITH	THE INTERIM
			INHALATION MI	THODOLOGY USED	BY THE RED	/Rfc WORK GROUP. ALSO	SEE TABLE	1: SUBCHRONIC	: AND CHRON	IC TOXICITY.	

**ACETONITRILE** 

000075-05-8

NOAEL 100 PPM

MOUSE

INHALATION:

92 DAYS

LIVER INCREASED RELATIVE WEIGHT 5E-1 300

C. babaania

5E-2 3000

Checosic

005208

SUBCHRONIC [RfC] COMMENT: 1E-1 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

CHRONIC [RfC] COMMENT: 1E-2 MG/KG/DAY. (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE. GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION [RFC] VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

**BARIUM** 

007440-39-3

NOEL 0.8 MG/CU M

INHALATION: INTERMITTENT

INTERMITTENT

4 MONTHS

**FETUS** 

**FETOTOXICITY** 

5E-3 100

5E-4 1000 005249

SUBCHRONIC [RfC] COMMENT: 1E-3 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). BASED ON A REPRODUCTION STUDY.

CHRONIC [RfC] COMMENT: 1E-4 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). BASED ON A REPRODUCTION STUDY.

GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION [RfC] VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

CHLORO-1,3-BUTADIENE / (CHLOROPRENE)

000126-99-8

NOAEL 10 PPM

RAT

2 YEARS INHALATION:

INTERMITTENT

HAIR WHOLE BODY ALOPECIA DECREASED WEIGHT GAIN 2E-2 100

2E-2 100

005878

SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION ASSUMING AN INHALATION ABSORPTION FACTOR OF 0.5.

CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION ASSUMING AN INHALATION ABSORPTION FACTOR OF 0.5.

GENERAL COMMENT: SEE ALSO HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

## HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

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Subchronic Chronic CHEMICAL DOSE SPECIES REFERENCE [RfC] [RfD] [RfC] [RfD] LEVEL ROUTE EXPERIMENT LENGTH TARGET CRITICAL EFFECT (mg/cu m) (mg/kg/day) (mg/cu m) (mg/kg/day) UF ÜF **CHLOROBENZENE** 000108-90-7 LOAEL 75 PPM RAT INHALATION: 120 DAYS LIVER 005353 **EFFECTS** 2E-2 INTERMITTENT KIDNEY **EFFECTS** 10000 CHRONIC (RfC) COMMENT: 5E-3 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE. GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION [RFC] VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY. CYCLOPENTADIENE 000542-92-7 NOEL 87.3 MG/KG/DAY RAT INHALATION: 194 DAYS LIVER LESIONS 3E+0 005401 INTERMITTENT KIDNEY LESIONS 100 SUBCHRONIC [RfC] COMMENT: 9E-1 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). GENERAL COMMENT: THE SUBCHRONIC INHALATION [RfC] VALUE WAS DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. **DICHLOROBENZENE, 1,2-**000095-50-1 NOEL 49 PPM RAT INHALATION: UP TO 7 WHOLE BODY DECREASED WEIGHT GAIN 2E+0 2E-1 005412 INTERMITTENT MONTHS 100 1000 CHRONIC [RfC] COMMENT: 4E-2 MG/KG/DAY (SEE APPENDIX A-11, DOSE CONVERSIONS ON HEAST). GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION (RFC) VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY. DICHLORODIFLUOROMETHANE 000075-71-8 LOAEL 482.3 MG/KG/DAY **GUINEA PIG** INHALATION: 005497 6 WEEKS LIVER LESIONS 2E+0 2E-1 INTERMITTENT 1000 10000

GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION [RFC] VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM

INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

SUBCHRONIC [RfC] COMMENT: 5E-1 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).
CHRONIC [RfC] COMMENT: 5E-2 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

# HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

Subchronic

Chronic

March 1994

<u>CHEMICAL</u>	<u>Dose</u> Route		CIES INT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	g/day) (mg/cu m) (mg/kg/	
DICHLOR	OETHAN	NE, 1,1-	0000	75-34-3				
NOEL	138 MG/KO INHALA INTERM	TION:	CAT 13 WEEKS	KIDNEY	DAMAGE	5E+0 100	5E-1 1000	005789
	CHRONIC	NIC [RFC] [RFC] COM COMMENT:	MENT: 1E-1 MG THE SUBCHRONI	G/KG/DAY (SEE (C AND CHRONI)	SEE APPENDIX A-II, DOSE CONVE APPENDIX A-II, DOSE CONVERSI C INHALATION [RFC] VALUES WEF SED BY THE RFD/RFC WORK GROUF	IONS ON HEAST). RE DERIVED FROM METHODOI	LOGY THAT IS NOT CURRENT W 1: CHRONIC AND SUBCHRONIC	ITH THE INTERIM

DICYCLOPENTADIENE 000077-73-6

LOAEL 1 PPM RAT

INHALATION: 90 DAYS LIVER DYSFUNCTION 2E-3 2E-4 005424
INTERMITTENT 1000 10600

SUBCHRONIC [RfC] COMMENT: 6E-4 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

CHRONIC [RfC] COMMENT: 6E-5 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION [RFC] VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

ETHOXYETHANOL ACETATE, 2- 000111-15-9

NOEL 30.1 MG/KG/DAY RAT

INHALATION: DAY 6-18 OF FETUS DECREASED OSSIFICATION 3E-1 3E-1 005952
INTERMITTENT GESTATION 100 100

SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. THE SUBCHRONIC ORAL [RfD]WAS BASED ON A REPRODUCTION STUDY WITH EXPOSURES DURING DAYS 6-18 OF GESTATION.

CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. THE CHRONIC ORAL [RfD] WAS BASED ON A REPRODUCTION STUDY WITH EXPOSURES DURING DAYS 6-18 OF GESTATION.

GENERAL COMMENT: NO PHARMACOKINETIC DATA WERE PROVIDED TO JUSTIFY ROUTE TO ROUTE EXTRAPOLATION FOR THE ORAL [RfD] VALUES. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY VALUES.

## HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

March 1994

CHEMICAL LEVEL		PECIES MENT LENGTH	TARGET	CRITICAL EFFECT	Subchron [RfC] [RfD] (mg/cu m) (mg/kg/da UF UF	[RfC] [RfD]	
FURFURA	<b>AL</b>	000098-0	1-1				
	20 PPM	HAMSTER					
	INHALATION: INTERMITTENT	13 WEEKS	NASAL CAVITY	OLFACTORY DEGENERATION	5E-1 100	5E~2 1060	005465
	SUBCHRONIC [RfC	COMMENT: 1E-	1 MG/KG/DAY (SEE	APPENDIX A-11, DOSE CONVERSION PENDIX A-11, DOSE CONVERSIONS	NS ON HEAST).		
	GENERAL COMMENT	: THE SUBCHRON	IC AND CHRONIC I	NHALATION [RfC] VALUES WERE DE BY THE RfD/RfC WORK GROUP. AL	RIVED FROM METHODOLOGY	THAT IS NOT CURRENT NONIC AND CHRONIC TOXIC	VITH THE INTERIM
METHAC	RYLONITRILE	0001	26-98-7				
NOEL	3.2 PPM	DOG					
	INHALATION: INTERMITTENT	90 DAYS	LIVER LIVER	INCREASED SGOT INCREASED SGPT	7E-3 300	7E-4 3000	005811
	SUBCHRONIC [RfC	COMMENT: 2E-	3 MG/KG/DAY (SEE	APPENDIX A-II, DOSE CONVERSIO	NS ON HEAST). THESE V	LUES DIFFER FROM THOS	IN THE 1987
	CHRONIC [RfC] C		G/KG/DAY (SEE AP	PENDIX A-II, DOSE CONVERSIONS	ON HEAST). THESE VALUE	S DIFFER FROM THOSE II	1 THE 1987
	GENERAL COMMENT	: THE SUBCHRON		NHALATION [RfC] VALUES WERE DE BY THE RfD/RfC WORK GROUP. AL	· - · · · · · · · · · · · · · · · ·		
METHOX	YETHANOL ACI	ETATE, 2-	000110-49-6				
NOAEL	10 PPM	RABBIT					
	INHALATION: INTERMITTENT	13 WEEKS	TESTIS	DEGENERATION	2E-7 100	2E-: 100	

SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.5. CALCULAYED FROM DATA OBTAINED WITH METHOXYETHANOL CONVERTED BY MULTIPLYING BY THE MOLECULAR WEIGHT RATIO (118.13/76.09).

CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.5. CALCULATED FROM DATA OBTAINED WITH METHOXYETHANOL CONVERTED BY MULTIPLYING BY THE MOLECULAR WEIGHT RATIO (118.13/76.09).

GENERAL COMMENT: NO PHARMACOKINETIC DATA WERE PROVIDED TO JUSTIFY ROUTE TO ROUTE EXTRAPOLATION FOR THE ORAL [RfD] VALUES.

# HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

March 1994

<u>CHEMICAL</u> LEVEL		PECIES MENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	REFERENCE		
METHOX	YETHANOL, 2-	0001	09-86-4						
	31 MG/CU M	RABBIT							
	INHALATION: INTERMITTENT	13 WEEKS	TESTICLE	EFFECTS	1E-2 100	1E-3 1000	010910		
	CHRONIC [RfD] CO	MMENT: BASED C	N ROUTE TO RO	PROUTE EXTRAPOLATION.  NUTE EXTRAPOLATION.  IC AND CHRONIC TOXICITY (OTI	MER THAN CARCINOGENICITY).				
METHYL	ACRYLATE	00009	6-33-3						
NOEL	15 PPM INHALATION: INTERMITTENT	RAT 2 YEARS		NONE OBSERVED	3E-2 100	3E-2 100	010003		
SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. GENERAL COMMENT: NO PHARMACOKINETIC DATA WERE PROVIDED TO JUSTIFY ROUTE TO ROUTE EXTRAPOLATION FOR THE ORAL [RfD] VALUES.									
METHYL	ISOBUTYL KETO	)NE 000	0108-10-1						
	50 PPM	RAT							
	INHALATION: INTERMITTENT	90 DAYS	LIVER KIDNEY	INCREASED WEIGHT EFFECTS	8E-1 100	8E-2 1000	005562		
	CHRONIC [RfC] CO	MMENT: 2E-2 MG	/KG/DAY (SEE	EE APPENDIX A-II, DOSE CONVI APPENDIX A-II, DOSE CONVERSI INHALATION [RFC] VALUES MEI		AT IS NOT CURRENT WITH	THE INTERIM		

INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

IRIS, EPA'S INTEGRATED RISK INFORMATION SYSTEM, IS UPDATED MONTHLY. FURTHER INFORMATION: RISK INFORMATION HOTLINE: (513) 569-7254.

# HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

March 1994

<u>CHEMICAL</u> LEVEL	<u>Dose</u> Route exi	<u>SPECIES</u> PERIMENT LENGTH	TARGET	CRITICAL EFFECT	[RfC]	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] [RfD] <u>(mg/cu_m) (mg/kg/day)</u> UF UF	REFERENCE
METHYL	STYRENE (M	IIXED ISOMERS)	025013-15-	4				
LOAEL	5.6 MG/KG/DAY INHALATION: INTERMITTEN	103 WEEKS	NASAL CAVITY	LESIONS		6E-3 1000	6E-3 1000	005567
	CHRONIC [RfD	] COMMENT: BASED (	ON ROUTE TO ROU'	ROUTE EXTRAPOLATION WITH TE EXTRAPOLATION WITH AN E PROVIDED TO JUSTIFY RO	ABSORPTION FACTOR OF	F 0.5.	ORAL [RfD] VALUES.	
LOAEL	11.2 MG/KG/DA INHALATION: INTERMITTEN	103 WEEKS	NASAL CAVITY	LESIONS	4E-2 1000		4E-2 1000	005566
	CHRONIC [RfC	] COMMENT: 1E-2 MG ENT: THE SUBCHRON	G/KG/DAY (SEE AI IC AND CHRONIC	E APPENDIX A-II, DOSE CO PPENDIX A-II, DOSE CONVE INHALATION [RfC] VALUES ( D BY THE RfD/RfC WORK GR	RSIONS ON HEAST). WERE DERIVED FROM ME	THODOLOGY THAT	IS NOT CURRENT WITH	THE INTERIM
METHYL	STYRENE, A	I PHA OOC	0098-83-9					
	970 MG/CU M	RAT	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,					
	INHALATION: INTERMITTEN		LIVER KIDNEY	INCREASED WEIGHT INCREASED WEIGHT		7E-1 100	7E-2 1000	010016
	CHRONIC [RfD	COMMENT: BASED (	ON ROUTE TO ROU'	ROUTE EXTRAPOLATION USING TE EXTRAPOLATION USING AL E PROVIDED TO JUSTIFY RO	N ABSORPTION FACTOR (	DF 0.5.	ORAL [RfD] VALUES.	
METHYLE	NE BROMID	E 0000	74-95-3					
NOAEL	11 MG/KG/DAY INHALATION: INTERMITTEN		BLOOD	INCREASED CARBOXYHE	MOGLOBIN	1E-1 100	1E-2 1000	010011
	CHRONIC [RfD	] COMMENT: BASED (	ON ROUTE TO ROU'	ROUTE EXTRAPOLATION, INC TE EXTRAPOLATION, INCLUD E PROVIDED TO JUSTIFY RO	ING AN ABSORPTION FA	CTOR OF 0.5.	_	

## HEAST TABLE 2: ALTERNATE METHODS - SUBCHRONIC AND CHRONIC TOX!CITY (OTHER THAN CARCINOGENICITY)

March 1994

<u>CHEMICAL</u> LEVEL	<u>pose</u> Route e	<u>SPECIES</u> XPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] [RfD] (mg/cu m) (mg/kg/day) UF UF	Chronic (RfC) (RfD) (mg/cu m) (mg/kg/day) UF UF	REFERENCE
NITROBE	NZENE	000098	3-95-3				
LOAEL	25 MG/CU M INHALATION INTERMITTE		BLOOD ADRENAL KIDNEY LIVER	HEMATOLOGICAL EFFECTS LESIONS LESIONS LESIONS	2E-2 1000	2E-3 10000	010518
	INHALATIO INTERMITT		BLOOD ADRENAL KIDNEY LIVER	HEMATOLOGICAL EFFECTS LESIONS LESIONS LESIONS			

SUBCHRONIC [RfC] COMMENT: 6E-4 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

CHRONIC [RfC] COMMENT: 6E-4 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION (RTC) VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM
INHALATION METHODOLOGY USED BY THE RTD/RTC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

TRICHLOROBENZENE, 1,2,4- 000120-82-1

GENERAL COMMENT: INFORMATION REMOVED FROM THIS TABLE. SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

TRICHLOROFLUOROMETHANE 000075-69-4

LOAEL 1940 MG/KG/DAY DOG

INHALATION: 90 DAYS KIDNEY INCREASED BUN 7E+0 7E-1 005501
CONTINUOUS LUNG INFLAMMATION 1000 10000

SUBCHRONIC [RfC] COMMENT: 2E+O MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

CHRONIC [RfC] COMMENT: 2E-1 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).

GENERAL COMMENT: THE SUBCHRONIC AND CHRONIC INHALATION [RFC] VALUES WERE DERIVED FROM METHODOLOGY THAT IS NOT CURRENT WITH THE INTERIM

INHALATION METHODOLOGY USED BY THE RFD/RFC WORK GROUP. ALSO SEE TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

# REFERENCES FOR HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

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## **ACETONE CYANOHYDRIN**

## 000075-86-5

010432 BLANK TL AND DC THAKE. 1984. THREE-MONTH INHALATION TOXICITY OF ACETONE CYANOHYDRIN IN MALE AND FEMALE SPRAGUE-DAWLEY RATS. MONSANTO REPORT NOP. MSL-4423. TSCA 8(D) SUBMISSION 878216397 (OTS 0510325).

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## **ACETONITRILE**

### 000075-05-8

005208 COATE WB. 1983. 90-DAY SUBCHRONIC TOXICITY STUDY OF ACETOMITRILE IN 86C3F1 MICE. FINAL REPORT (REVISED).

US EPA. 1987. HEALTH EFFECTS ASSESSMENT FOR ACETONITRILE.PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF EMERGENCY AND REMEDIAL RESPONSE, WASHINGTON, DC.

## **BARIUM**

#### 007440-39-3

005249 TARASENKO N, O PROMIN AND A SILAYEV. 1977. BARIUM COMPOUNDS AS INDUSTRIAL POISONS (AN EXPERIMENTAL STUDY). J HYG EPIDEM MICROBIOL IMMUNOL. 21: 361.

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## CHLORO-1,3-BUTADIENE / (CHLOROPRENE)

### 000126-99-8

005878 DU PONT DE NEMOURS AND COMPANY, INC. 1985. 2-YEAR INHALATION CARCINOGENICITY STUDY OF CHLOROPRENE IN RATS. EI DU PONT DE NEMOURS AND CO., INC., WILMINGTON, DE.

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### CHLOROBENZENE

### 000108-90-7

005353 DILLEY, JV. 1977. TOXIC EVALUATION OF INHALED CHLOROBENZENE. NIOSH, DHEW, CINCINNATI, OH, CONTRACT 210-76-0126. CITED IN US EPA, 1985.

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## CYCLOPENTADIENE

### 000542-92-7

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# REFERENCES FOR HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

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## DICHLOROBENZENE, 1,2-

### 000095-50-1

005412 HOLLINGSWORTH RL, VK ROWE, F OYEN, TR TORKELSON AND EM ADAMS. 1958. TOXICITY OF O-DICHLOROBENZENE. AM MED ASSOC ARCH IND HEALTH. 17(1): 180-187.

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### **DICHLORODIFLUOROMETHANE**

#### 000075-71-8

005497 PRENDERGAST JA, RA JONES, LJ JENKINS AND J SIEGAL. 1967. EFFECTS ON EXPERIMENTAL ANIMALS OF LONG-TERM INHALATION OF TRICHLOROETHYLENE, CARBON TETRACHLORIDE, 1,1,1-TRICHLOROETHANE, DICHLORODIFLUOROMETHANE AND 1,1-DICHLOROETHYLENE. TOXICOL APPL PHARMACOL. 10: 270-289.

US EPA. 1987. HEALTH EFFECTS ASSESSMENT FOR FULLY HALOGENATED METHANES. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF EMERGENCY AND REMEDIAL RESPONSE, WASHINGTON. DC.

## DICHLOROETHANE, 1,1-

## 000075-34-3

005789 HOFMANN HT, H BIRNSTIEL AND P JOBST. 1971. ON THE INHALATION TOXICITY OF 1,1- AND 1,2-DICHLOROETHANE, ARCH TOXIKOL. 27: 248-265.

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### DICYCLOPENTADIENE

### 000077-73-6

DODD DE, LC LONGO AND DL EISLER. 1982. DICYCLOPENTADIENE VAPOR NINETY-DAY INHALATION STUDY ON RATS AND MICE. BUSHY RUN RESEARCH CENTER, EXPORT, PA, TSCA 8E SUBMISSION BY EXXON CHEM AMER DOC 1D 88-8300464, ODD DOC 1D 88HQ-0283-0364, MICROFICHE NO. OTS 204864.

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### ETHOXYETHANOL ACETATE, 2-

## 000111-15-9

005952 UNION CARBIDE. 1984. A TERATOGENIC EVALUATION OF CELLOSOLVE ACETATE IN FISHER 344 RATS AND NEW ZEALAND WHITE RABBITS FOLLOWING INHALATION EXPOSURE. BUSHY RUN RESEARCH CENTER, EXPORT, PA, OCTOBER 1984. FYI-AX-1184-0360.

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## **FURFURAL**

## 000098-01-1

005465 FERON VJ, A KRUYSSE AND HC DREEF VANDER MEULEN. 1979. REPEATED EXPOSURE TO FURFURAL VAPOR: 13 WEEK STUDY IN SYRIAN GOLDEN HAMSTERS. ZENTRASE. BAKTEVIOL PAVASITEN KD INFECTION SKV HYG ABT 1 ORIG REIHE B. 168(5-6): 442-451.

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# REFERENCES FOR HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

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### METHACRYLONITRILE

### 000126-98-7

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### METHOXYETHANOL, 2-

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### METHYL ACRYLATE

### 000096-33-3

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# REFERENCES FOR HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

March 1994

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000098-83-9

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000074-95-3

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010518 CIIT (CHEMICAL INDUSTRY INSTITUTE OF TOXICOLOGY). 1984. NINETY- DAY INHALATION STUDY OF NITROBENZENE IN F-344 RATS,CD RATS AND B6C3F1 MICE WITH COVER LETTER DATED 6/24/84 AND EPA RESPONSE DATED 8/06/84, UNPUBLISHED STUDY. FYI-OTS-0784-0333 AND COMPUTER PRINT-OUT OF PATHOLOGY FINDING.

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# REFERENCES FOR HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

March 1994

## **TRICHLOROFLUOROMETHANE**

## 000075-69-4

005501 JENKINS, LJ, RA JONES, RA COON AND J SIEGAL. 1970. REPEATED AND CONTINUOUS EXPOSURES OF LABORATORY ANIMALS TO TRICHLOROFLUORMETHANE. TOXICOL APPL PHARMACOL. 16: 133-142.

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[SLOPE FACTOR] **CUNIT RISKI EXPERIMENT LENGTH** 

**CEPA** ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL ROUTE SPECIES TARGET CANCER GROUP] (mg/kg/day)<sup>-1</sup>(mg/kg/day)<sup>-1</sup> (ug/L)-1 (ug/cu m)-1

**ACEPHATE** 030560-19-1

> IRIS IRIS IRIS 010086

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**ACROLEIN** 000107-02-8

> IRIS 005001

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**ACRYLAMIDE** 000079-06-1

> ORAL: DRINKING 2 YEARS IRIS IRIS 4.5E+0 IRIS

IRIS 010087 WATER RAT MAHMARY **TUMORS** 

> THYROID TUMORS UTERUS **TUMORS** ORAL CAVITY TUMORS CENTRAL MERVOUS TUMORS

SYSTEM

INHALATION (SLOPE) COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**ACRYLONITRILE** 000107-13-1

> IRIS IRIS IRIS 005004

> INHALATION: IRIS 2.4E-1 IRIS 005003

**OCCLIPATIONAL** HE BEAM LUNG **TUMORS** 

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON NEAST.

**ALACHLOR** 015972-60-8

> ORAL: DIET MULTIPLE SITES TUMORS 8E-2 2.3E-6 010180

ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

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[SLOPE FACTOR] **CUNIT RISK!** EXPERIMENT LENGTH **IEPA** ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL ROUTE SPECIES TARGET CANCER GROUP] (mg/kg/day)<sup>-1</sup>(mg/kg/day)<sup>-1</sup> (vg/L)<sup>-1</sup> (ug/cu m)-1 **ALDRIN** 000309-00-2 ORAL: DIET IRIS IRIS 1.7E+1 IRIS IRIS 005006 MOUSE LIVER CARCINONA INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. INHALATION (UNIT RISK) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **ALLYL CHLORIDE** 

000107-05-1

IRIS

010181

GENERAL COMMENT: FOR RCRA ACTIVITIES ONLY, CONTACT THE HEALTH ASSESSMENT SECTION (202) 260-4761 FOR RCRA APPROVED NUMERIC ASSESSMENT OF THIS

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

ANILINE

000062-53-3

IRIS

IRIS

IRIS

010088

IRIS GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**ARAMITE** 

000140-57-8

ORAL: DIET 104 WKS

**TUMORS** 

IRIS 2.5E-2

IRIS

IRIS

010206

INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-11, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**TUMORS** 

ARSENIC, INORGANIC

007440-38-2

IRIS

010925

INHALATION: **OCCUPATIONAL** 

HUMAN

RAT

RESPIRATORY

SYSTEM

LIVER

IRIS

5.0E+1

IRIS

005007

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FAC ORAL (mg/kg/day) <sup>-1</sup> (mg/	INHALATION	(UNIT R ORAL (Ug/L) <sup>-1</sup>	ISK] INHALATION (ug/cu m)-1	REFERENCE
ASBESTO	os	001332-21-4								
					IRIS IRIS				IRIS	005010 005919
ATRAZIN	E	001912-24-9								
	ORAL: DIET	2 YEARS RAT	MANNARY GLAND MANNARY GLAND MANNARY GLAND MANNARY GLAND	ADENOMA FIBROADENOM ADENOCARCIN CARCINOSARC	OHA	2.22E-1		6.3E-6		010380
	ORAL (SLOPE) GENERAL COMME	COMMENT: UNDER REVIE NT: ALSO SEE HEAST 1	EV, NUMBER SUBJECT T ABLE 1: SUBCHRONIC	O CHANGE. AND CHRONIC T	OXICITY (	OTHER THAN CARCI	(NOGENICITY)			
AZOBENZ	ZENE	000103-33-	3							
	ORAL: DIET	2 YEARS RAT	ABDOMINAL CAVITY	SARCOMA	IRIS	IRIS	1.1E-1	IRIS	IRIS	010089
	INHALATION [S	LOPE] COMMENT: SEE /	PPENDIX A-II, DOSE	CONVERSIONS O	N HEAST.					
BENZENE		000071-43-2								
	INHALATION: OCCUPATIONAL	HUHAN	BLOOD	LEUKENIA	IRIS	IRIS	2.9E-2	IRIS	IRIS	005011
	INHALATION [S	COMMENT: BASED ON RO LOPE] COMMENT: SEE A IT: ALSO SEE HEAST I	PPENDIX A-II, DOSE	CONVERSIONS O		OTHER THAN CARCI	NOGENICITY)			
BENZIDIN	E	000092-87-5								
	GENERAL COMME	IT: ALSO SEE HEAST 1	ABLE 1: SUBCHRONIC	AND CHRONIC TO	IRIS OXICITY (	IRIS OTHER THAN CARCI	IRIS NOGENICITY).	IRIS	IRIS	005014
BENZOTR	RICHLORIDE	000098-0	7-7							
					IRIS	IRIS		IRIS		010092

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CHENICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL INHALATION (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	[UNIT RIS ORAL (ug/L) <sup>-1</sup>	SK) INHALATION (ug/cu m)-1	REFERENCE
BENZO[A	]ANTHRACEN	IE 0000	56-55-3		IRIS				010182
BENZO[A	]PYRENE	000050-	32-8		IRIS	IRIS	IRIS		010508
BENZO[B	FLUORANTH	ENE 0002	205-99-2						
	GENERAL COMME	NT: ALSO SEE HEAST	TABLE 1: SUBC	HRONIC AND CHRONIC TO	IRIS KICITY (	OTHER THAN CARCINOGENICITY	).		010183
BENZO[K	FLUORANTH	ENE 000	207-08-9						
	GENERAL COMME	NT: FOR RCRA ACTIV	/ITIES ONLY, CO	NTACT THE MEALTH ASSE	IRIS SSMENT S	ECTION (202) 260-4761 FOR (	RCRA AFPROVED NUI	MERIC ASSESSMI	010090 ENT OF THIS
BENZYL (	CHLORIDE	000100	44-7						
					IRIS	IRIS	IRIS		010093
BERYLLIU	M	007440-41	-7						
	GENERAL COMME	NT: ALSO SEE HEAST	TABLE 1: SUBC	RONIC AND CHRONIC TO	IRIS XICITY (	IRIS OTHER THAN CARCINOGENICITY	IRIS )_		005018
	INHALATION: OCCUPATIONAL	HUMAN	LUNG	TUMORS	IRIS	8.4E+0		IRIS	005017
	INHALATION [S	LOPE) COMMENT: SEE	APPENDIX A-II,	, DOSE CONVERSIONS ON	HEAST.				
BIS(2-CH	LOROETHYL)	ETHER 000	111-44-4						
	ORAL	560 DAYS Mouse	LIVER	TUMORS	IRIS	IRIS 1.1E+0	IRIS	IRIS	005076
	INHALATION [SI	LOPE] COMMENT: BAS	SED ON ROUTE TO	ROUTE EXTRAPOLATION.	SEE APP	ENDIX A-II, DOSE CONVERSION	IS ON HEAST.		

CHEMICAL	ROUTE	CPERIMENT LENG SPECIES	TH TARGET	CANCER	[EPA GROUP]	(SLOPE FAI ORAL (mg/kg/dey) <sup>-1</sup> (mg	INHALATION	(UNIT II ORAL (ug/L) <sup>-1</sup>	ISK] INHALATION (ug/cu m)-1	REFERENCE
BIS(2-CHI	LOROISOPROP	YL) ETHER	039638-32-9							
	ORAL: GAVAGE	2 YEARS Mouse	LIVER LUNG	TUMORS TUMORS	С	7E-2	3.5E-2	<b>2E-6</b>	1E-5	005079

INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION (50% RESPIRATORY ABSORPTION), SEE APPENDIX A-II: DOSE CONVERSIONS ON MEAST. GENERAL COMMENT: FORMERLY LISTED AS BIS(2-CHLORO-1-METHYL(ETHYL)ETHER). ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

BIS(2-ETHYLHEXYL) PHTHALATE / (DEHP) 000117-81-7

IRIS IRIS IRIS 005120

GENERAL CONNENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**BIS(CHLOROMETHYL) ETHER** 000542-88-1

> INHALATION: 10-100 DAYS IRIS IRIS 2.2E+2 IRIS IRIS 005077

INTERMITTENT RAT RESPIRATORY **TUMORS** 

SYSTEM

ORAL [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION.

INHALATION (SLOPE) COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

**BROMODICHLOROMETHANE** 000075-27-4

> IRIS IRIS IRIS 005148

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

BROMOETHENE / (VINYL BROMIDE) 000593-60-2

INHALATION: 2 YEARS **B2** 1.1E-1 3.2E-5 010094

INTERMITTENT RAT LIVER

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

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[SLOPE FACTOR] **CUNIT RISKI** EXPERIMENT LENGTH REFERENCE ORAL INHALATION ORAL INHALATION **IEPA** CHEMICAL ROUTE TARGET GROUP] (mg/kg/day)-1 (mg/kg/day)-1 (ug/L)-1 SPECIES CANCER (ug/cu m)-1 **BROMOFORM** 000075-25-2 2 YEARS IRIS 3.9E-3 IRIS IRIS 005150 ORAL: GAVAGE IRIS RAT INTESTINE, LARGE ADENOMATOUS POLYP **ADENOCARCI NOMA** INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE MEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **BUTADIENE, 1.3-**000106-99-0 1.8E+0 IRIS 010477 INHALATION: IRIS INTERMITTENT MOUSE MULTIPLE SITES **TUMORS** INHALATION (SLOPE) COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON MEAST. **BUTYL BENZYL PHTHALATE, N-**000085-68-7 005122 IRIS GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) **CADMIUM** 007440-43-9 IRIS 005019 IRIS ORAL ISLOPEI COMMENT: THERE IS INADEQUATE EVIDENCE FOR THE CARCINOGENICITY OF THIS COMPOUND BY THE ORAL ROUTE. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **CAPTAFOL** 002425-06-1 8.6E-3 2.4E-7 010095 ORAL: DIET MOUSE LYMPHATIC SYSTEM LYMPHOSARCOMA ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **CAPTAN** 000133-06-2 010184 1.0E-7 **B2** 3.5E-3 ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

						(SLOPE FACT	OR]	(UNIT R	[SK]	
CHEMICAL	ROUTE	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	ORAL (mg/kg/day) <sup>-1</sup> (mg/	INHALATION kg/day) <sup>-1</sup>	ORAL (ug/L) <sup>-1</sup>	INHALATION (ug/cu m)-1	REFERENCE
CARBAZO	OLE	000086-74-8	3							
	ORAL: DIET	% WEEKS WOUSE	LIVER	TUMORS	<b>B2</b>	<b>2</b> E-2		5.7E-7		010096
CARBON	TETRACHLORII	DE 00005	6-23-5							
0,4,12014	ORAL: DIET		LIVER	TUMORS	IRIS	IRIS	5. <b>3</b> E-2	IRIS	IRIS	005022
CHLORAI	INHALATION [UNI GENERAL COMMENT	PE] COMMENT: SEE A T RISK] COMMENT: B : ALSO SEE HEAST T  000118-75-2 82 WEEKS	ASED ON ROUTE TO RO ABLE 1: SUBCHRONIC	OUTE EXTRAPOLA AND CHRONIC TO	TION. IN			0x OF 0.4.		010097
CHLORD	A NE	MOUSE 000057-74-	LIVER LUNG	TUMORS TUMORS						
CHLORD	ORAL: DIET	000037-74-	9		IRIS	IRIS	1.3E+0	IRIS	IRIS	005024
		MOUSE	LIVER	CARCINOMA						
		PE] COMMENT: BASED : ALSO SEE HEAST 1						ON HEAST.		
CHLORO	-2-METHYLANIL	INE HYDROCHLO	ORIDE, 4- 00	03165-93-3						
	ORAL: DIET	18 MONTHS MOUSE	CARDIOVASCULAR SYSTEM CARDIOVASCULAR	NEMANGIOMA HEMANGIOSAR	B2 CONA	4.6E-1		1.3E-5		010419
			SYSTEM							

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[SLOPE FACTOR] **CUNIT RISK** EXPERIMENT LENGTH **IEPA** ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL ROUTE SPECIES TARGET CANCER (ug/L)<sup>-1</sup> GROUP] (ma/ka/dev)<sup>-1</sup>(ma/ka/dev)<sup>-1</sup> (ug/cu m)-1 CHLORO-2-METHYLANILINE, 4-000095-69-2 18 MONTHS ORAL: DIET **B2** 5.8E-1 010098 1.6E-5 MOUSE CARD IOVASCULAR HEMANGIONA SYSTEM CARD IOVASCULAR **HEMANGIOSARCOMA** SYSTEM

ORAL [SLOPE] COMMENT: BASED ON VASCULAR TUMORS IN NICE TREATED WITH 4- CHLORO-2-METHYLANILINE HYDROCHLORIDE.

CHLOROBENZILATE

000510-15-6

ORAL: GAVAGE. **82 VEEKS** DIET MOUSE

**B2 HEPATOMA** 

2.7E-1

IRIS

2.7E-1

7.8E-6 7.8E-5 010848

INHALATION [SLOPE] COMMENT: ABSORBANCE BY THE INHALATION ROUTE WAS ASSUMED TO EQUAL ORAL ABSORPTION SINCE THERE WERE NO PHARMACOKIMETIC DATA TO THE CONTRARY. SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST.

INHALATION LUNIT RISKI COMMENT: ABSORBANCE BY THE INHALATION ROUTE WAS ASSUMED TO EQUAL ORAL ABSORPTION SINCE THERE WERE NO PHARMACOKINETIC DATA TO THE CONTRARY.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**CHLOROFORM** 

000067-66-3

IRIS

IRIS

005036

ORAL: GAVAGE

78 WEEKS

LIVER

IRIS

C

8.1E-2

IRIS

MOUSE

LIVER

CARCINOMA

005035

INHALATION (SLOPE) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-11, DOSE CONVERSIONS ON HEAST.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**CHLOROMETHANE** 

000074-87-3

INHALATION:

INTERMITTENT

24 MONTHS MOUSE

KIDNEY

**TUMORS** 

1.3E-2

6.3E-3

3.7E-7 1.8E-6 005038

ORAL [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENCIITY).

CHENICAL	ROUTE EXP	ERIMENT LENGT SPECIES	<u>H</u> Target	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL INHALAT: (mg/kg/dey) <sup>-1</sup> (mg/kg/dey) <sup>-1</sup>		RISK) INHALATION REFERENCE (ug/cu m)-1
CHLORON	METHYL METHY		000107-30-2		IRIS			005081
		FOR RCRA AC POUND.	CTIVITIES ONLY, CONTACT	THE HEALTH ASS	ESSMENT	SECTION (202) 260-4761 FG	R RCRA APPROVED	NUMERIC ASSESSMENT OF THIS
CHLORON	NITROBENZENE,		00088-73-3					
	ORAL: DIET	18 MONTHS MOUSE	LIVER	TUMORS	82	2.5E-2	7.1E-7	010099
CHLORON	NITROBENZENE,		00100-00-5		_			
	ORAL: DIET	18 MONTHS MOUSE	CARDIOVASCULAR SYSTEM	TUMORS	82	1.8E-2	5.1E-7	010100
CHLOROT	THALONIL	0018	97-45-6					
	ORAL: DIET	27-32 MONTHS	KIDNEY	TUMOR	<b>B</b> 2	1.1E-2	3.1E-7	010384
			REVIEW, NUMBER SUBJECT EAST TABLE 1: SUBCHRONIO		OXICITY	(OTHER THAN CARCINOGENIC	TY).	
CHROMIC	JM(VI)	018540	)-29-9					
	INHALATION: OCCUPATIONAL	HUMAN	LUNG	TUMORS	IRIS	4.1E+	1	IRIS 005091
			SEE APPENDIX A-II, DOS EAST TABLE 1: SUBCHRONIO			(OTHER THAN CARCINOGENIC	ITY).	

CHRYSENE 000218-01-9

> 010185 IRIS

GENERAL COMMENT: FOR RCRA ACTIVITIES ONLY, CONTACT THE HEALTH ASSESSMENT SECTION (202) 260-4761 FOR RCRA APPROVED NUMERIC ASSESSMENT OF THIS COMPOUND, ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

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**CUNIT RISK [SLOPE FACTOR]** EXPERIMENT LENGTH INHALATION ORAL INHALATION REFERENCE ORAL CHEMICAL (ug/L)<sup>-1</sup> ROUTE SPECIES TARGET CANCER GROUP] (mg/kg/day)<sup>-1</sup>(mg/kg/day)<sup>-1</sup> (ug/cu m)-1 **COKE OVEN EMISSIONS** 008007-45-2 INHALATION: IRIS 2.2E+0 IRIS 005039 OCCUPATIONAL LUNG **TUMORS** INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON NEAST. GENERAL COMMENT: FORMERLY LISTED UNDER COAL TARS. CREOSOTE, COAL TAR 008001-58-9 005042 IRIS CRESOL, M- / (3-METHYLPHENOL) 000108-39-4 010187 IRIS GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). CRESOL, O- / (2-METHYLPHENOL) 000095-48-7 010186 IRIS GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). CRESOL, P- / (4-METHYLPHENOL) 000106-44-5 010188 GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). CROTONALDEHYDE 000123-73-9 010190 5.4E-5 ORAL: DRINKING 113 WKS IRIS 1.9E+0 WATER RAT LIVER TUMOR ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

	Đ	CPERIMENT LENGTH			[EPA		INHALATION	(UNIT R	ISK) INHALATION	REFERENCE
CHEMICAL	ROUTE	SPECIES	TARGET	CANCER	GROUP]	(mg/kg/day) <sup>-1</sup> (mg/	kg/day) <sup>-1</sup>	(ug/L) <sup>-1</sup>	(ug/cu m)-1	
CYANAZI	NE	021725-46-2								
	ORAL: DIET	2 YEARS RAT	MANNARY GLAND	ADENOMA/ CARCINOMA, COMBINED	C	8.4E-1		2.4E-5		010944
	GENERAL COMMEN	T: ALSO SEE MEAST 1	ABLE 1: SUBCHRONIC	AND CHRONIC TO	OXICITY (	OTHER THAN CARCI	NOGENICITY).			
DDD		000072-54-8			IRIS	IRIS		IRIS		010291
					IKIS	IRIS		IRIS		010271
DDE		000072-55-9			IRIS	IRIS		IRIS		010292
DDT	ORAL: DIET	000050-29-3			IRIS	IRIS	3.4E-1	IRIS	IRIS	005044
	OWE. DIEI	MOUSE, RAT	LIVER	TUMORS	IRIS	1413	J.4E-1	IKIS	IRIS	003044
		OPE] COMMENT: BASED T: ALSO SEE HEAST 1						N FEAST.		
DECABRO	OMODIPHENYL	ETHER 001	163-19-5							
	GENERAL COMMEN	T: ALSO SEE HEAST 1	ABLE 1: SUBCHRONIC	AND CHRONIC TO	IRIS XICITY (	OTHER THAN CARCI	NOGENICITY).			010102
DIALLATI	F	002303-16-4								
DINEERI	ORAL	19 MONTHS MOUSE	LIVER	TUMORS	82	6.1E-2		1.7E-6		010103
DIBENZO	(A,H)ANTHRA	CENE 0000	53-70-3							
	GENERAL COMMEN	T: FOR RCRA ACTIVII	IES ONLY, CONTACT T	HE HEALTH ASSI	IRIS ESSMENT S	SECTION (202) 260	-4761 FOR RCRA	APPROVED N	UMERIC ASSESSM	010191 ENT OF THIS

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						[SLOPE FACTOR]	[UNIT	RISKI	
CHEMICAL	ROUTE	CPERIMENT LENGTH SPECIES	TARGET	CANCER	(EPA GROUP)	ORAL INHALATION (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	ORAL (ug/L) <sup>-1</sup>	INHALATION (ug/cu m)-1	REFERENCE
DIBROMO	0-3-CHLOROPR	OPANE, 1,2 0	00096-12-8						
	ORAL: DIET		STOMACH KIDNEY LIVER	TUMORS TUMORS TUMORS	<b>8</b> 2	1_4E+0	4E-5		010484
	[EPA GROUP] CO	MMENT: UNDER REVII OPE] COMMENT: SEE	EV, CLASSIFICATION APPENDIX A-II: DOS	SUBJECT TO CHA E CONVERSIONS	NGE. ON NEAST.				
	INHALATION: INTERMITTENT	RAT, MOUSE	NASAL CAVITY	TUMORS	<b>82</b>	2.4E-3		6.9E-7	010519
DIBROMO	OCHLOROMET	HANE 00	0124-48-1						
	GENERAL COMMEN	T: FORMERLY LISTER	AS CHLORODIBRONOM	ETHANE. ALSO S	IRIS EE MEAST	IRIS TABLE 1: SUBCHRONIC AND CHRONI	IRIS IC TOXICITY	(OTHER THAN CAI	010891 CCINOGENICITY).
DIBROMO	OETHANE, 1,2-	000106	6-93-4						
	GENERAL COMMEN	T: FORMERLY LISTER	UNDER ETHYLENE DI	BROMIDE	IRIS	IRIS	IRIS		005818
	INHALATION: INTERMITTENT	88-103 WEEKS RAT	NASAL CAVITY	TUMORS	IRIS	7.6E-1		IRIS	005071
			APPENDIX A-II, DOS TABLE 1: SUBCHRONI			(OTHER THAN CARCINOGENICITY).			
DICHLOR	O-2-BUTENE, 1	I, <b>4</b> - 00076	4-41-0						
	INHALATION: INTERMITTENT	90 DAYS Rat	NASAL PASSAGES	TUMORS	<b>B2</b>	9.3E+0		2.6E-3	005053
	INHALATION [SL	OPE] COMMENT: SEE	APPENDIX A-II: DOS	E CONVERSIONS	ON HEAST.				

GENERAL COMMENT: FORMERLY LISTED UNDER DICHLOROBUTENES

[SLOPE FACTOR] CUNIT RISKT EXPERIMENT LENGTH (EPA ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL GROUP] (ma/kg/day)-1 (mg/kg/day)-1 ROUTE SPECIES TARGET CANCER (ug/L)<sup>-1</sup> (ug/cu m)-1 **DICHLOROBENZENE. 1.4-**000106-46-7 ORAL: GAVAGE 103 VEEKS **B**2 2.4E-2 6.8E-7 005050 MOUSE LIVER **TUMORS** ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). DICHLOROBENZIDINE, 3,3'-000091-94-1 IRIS IRIS IRIS 005815 DICHLOROETHANE, 1,1-000075-34-3 IRIS 005055 GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **DICHLOROETHANE, 1,2-**000107-06-2 ORAL: GAVAGE 78 WEEKS. 1915 IRIS 9.1E-2 IRIS IRIS 005058 RAT CIRCULATORY SARCONA SYSTEM INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY) DICHLOROETHYLENE, 1,1-000075-35-4 IRIS IRIS IRIS 005060 INHALATION 12 MONTHS IRIS 1.2E+0 IRIS 005059

**ADENOCARCINOMA** 

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

KIDNEY

MOUSE

CHEMICAL	ROUTE	ERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	(SLOPE FACTOR) ORAL INHALATION (mg/kg/dmy) <sup>-1</sup> (mg/kg/dmy) <sup>-1</sup>	(UNIT II ORAL (Ug/L) <sup>-1</sup>	ISK) INHALATION (ug/cu m)-1	REFERENCE
DICHLOR	OPROPANE, 1,2 ORAL: GAVAGE	- 000078	1-87-5 LIVER	TUMORS	82	6.8E-2	1.9E-6		005062
			W, NUMBER SUBJECT T ABLE 1: SUBCHRONIC		OXICITY	(OTHER THAN CARCINOGENICITY)			
DICHLOR	OPROPENE, 1,3 ORAL: GAVAGE	- / (TELONE II) 104 WEEKS MOUSE	000542-75-6 BLADDER RESPIRATORY SYSTEM FORESTOMACH	CARCINOMA ALVEOLAR/ BRONCHIOLAR ADENOMA PAPILLOMA/ CARCINOMA	IRIS	1.8E-1	5E-6		010946
	ORAL: GAVAGE	104 WEEKS RAT	LIVER FORESTOMACH	NEOPLASTIC NODULE/CARC PAPILLONA/ CARCINONA	INONA				
						OF COMBINED TUMORS LISTED. (OTHER THAN CARCINOGENICITY)			
	INHALATION: INTERMITTENT	2 YEARS MOUSE	LUNG	ADENONA	IRIS	1.3E-1		3.7E-5	010104
	INHALATION (SLO GENERAL COMMENT	PE] COMMENT: SEE A: ALSO SEE HEAST	APPENDIX A-II: DOSE FABLE 1: SUBCHRONIC	CONVERSIONS OF	N HEAST. OXICITY	(OTHER THAN CARCINOGENICITY).			
DIELDRIN	ORAL: DIET	000060-57-1 MOUSE	LIVER	CARCINONA	IRIS	IRIS 1.6E+1	IRIS	IRIS	005816
			APPENDIX A-II, DOSE BASED ON ROUTE-TO-RE		TION.				

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

ABLE 3: CARCINOGENICITY Merch 1994

CHENICAL	ROUTE	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP] (	[SLOPE FACTOR] ORAL INHALATION mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	[UNIT RISK]  ORAL INHALATION (Ug/L)-1 (ug/cu m)-1	REFERENCE
DIETHYLS	STILBESTROL ORAL: DIET	000056-!	53-1 Mammary Gland	CARCINONA	<b>A</b>	4.7E+3	1.3E-1	010485
DIMETHO	XYBENZIDINE, ORAL: DIET	3,3'- 00011! Lifetime Hamster	9-90-4 Forestomach	PAPILLONA	<b>B</b> 2	1.4E-2	€E-7	010106
DIMETHY	LANILINE HYDF ORAL: DIET	ROCHLORIDE, 2,4 18 Months Mouse	- 021436-96-4 LLING	TUMORS	C	5.8E-1	1.7€-5	010108
DIMETHY	LANILINE, 2,4- ORAL: DIET	000095-6 18 MONTHS MOUSE	8-1 LUMG	TUMORS	c	7.5E-1	2.1E-5	010107
DIMETHY	LBENZIDINE, 3, Oral: Gavage	3'- 000119- 30 days Rat	93-7 Manhary	TUMORS	B2	9.2E+0	2.6E-4	010109

DIMETHYLBENZ(A)ANTHRACENE, 7,12- 000057-97-6

010297

GENERAL COMMENT: FOR RCRA ACTIVITIES ONLY, CONTACT THE HEALTH ASSESSMENT SECTION (202) 260-4761 FOR RCRA AFPROVED NUMERIC ASSESSMENT OF THIS COMPOUND.

**DIMETHYLHYDRAZINE, 1,1-** 000057-14-7

GENERAL COMMENT: NO EPA GROUP CLASSIFICATION WAS PROVIDED IN THE REFERENCE DOCUMENT, THEREFORE THIS COMPOUND WAS REMOVED FROM TABLE 3.

**DIMETHYLHYDRAZINE, 1,2-** 000540-73-8

GENERAL COMMENT: CONTACT THE SIDEREIND HEALTH BICK TECHNICAL SIDERET CENTER, (547) 540 7700

GENERAL COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

DIMETHYLSULFATE 000077-78-1

IRIS 010112

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[SLOPE FACTOR] **CUNIT RISKI** EXPERIMENT LENGTH INHALATION INHALATION REFERENCE (EPA ORAL ORAL CHEMICAL ROUTE SPECIES TARGET CANCER GROUP] (mg/kg/day)<sup>-1</sup>(mg/kg/day)<sup>-1</sup> (ug/L)<sup>-1</sup> (ug/cu m)-1

**DINITROTOLUENE, 2,4-**000121-14-2

> IRIS IRIS 005066 GENERAL COMMENT: LISTED AS "DINITROTOLUENE MIXTURE, 2,4-/2,6-" ON IRIS. ALSO SEE NEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**DINITROTOLUENE, 2,6-**000606-20-2

> IRIS IRIS 005068 IRIS GENERAL COMMENT: LISTED AS "DINITROTOLUENE MIXTURE, 2,4-/2,6-" ON IRIS. ALSO SEE NEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

DIOXANE, 1.4-000123-91-1

IRIS IRIS IRIS 010298

**DIPHENYLHYDRAZINE, 1,2-**000122-66-7

> ORAL: DIET 2 YEARS IRIS IRIS 8.0E-1 IRIS IRIS 005070

LIVER RAT **TUMORS** 

INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-11, DOSE CONVERSIONS ON HEAST.

**DIRECT BLACK 38** 001937-37-7

> ORAL: DIET 93 DAYS 8.6E+0 2.4E-4 010113

RAT LIVER TUMORS

ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

**DIRECT BLUE 6** 002602-46-2

> ORAL: DIET 91 DAYS 010114 8.1E+0 2.3E-4

> > RAT LIVER TUMORS

ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

CHEMICAL	ROUTE EX	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL INHALATIO (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	(UNIT RISK) N ORAL INHALATIO (Ug/L) <sup>-1</sup> (Ug/cu m)	
DIRECT E	ROWN 95 ORAL: DIET	016071-8 91 days Rat	6-6 LIVER	TUMORS	A	9.3E+0	2.6E-4	010115
	ORAL (SLOPE) C	OPPIENT: UNDER REVI	EV, NUMBER SUBJECT	TO CHANGE.				
DIRECT S	KY BLUE 6B	002610-0	5-1		82			010116
	(EPA GROUP) COM	MENT: BASED ON THE	FACT THAT 3,3-DINE	THOXYBENZIDIN	E, A KNOW	N EPA GROUP B2 CARCINOGEN	, IS A METABOLITE OF DIRECT	SKY BLUE 68.
EPICHLO	ROHYDRIN	000106-8	9-8		IRIS	IRIS	IRIS	010198
	INHALATION: INTERMITTENT	30 DAYS, OBSERVED LIFETIME RAT	NASAL CAVITY	TUMORS	IRIS	4.2 <u>E</u> -3	IRIS	010117
	INHALATION [SLO GENERAL COMMENT	PE] COMMENT: SEE A : ALSO SEE HEAST T	PPENDIX A-II, DOSE ( ABLE 1: SUBCHRONIC /	CONVERSIONS OF	HEAST.	OTHER THAN CARCINOGENICITY	n.	
ETHYL A	CRYLATE ORAL: GAVAGE	000140-88 104 WEEKS RAT	3-5 Forestomach	TUMORS	B2	4.8E-2	1.4E-6	010118
ETHYLEN	E OXIDE ORAL: GAVAGE	000075-21 LIFETIME, TWICE W	-	TUMORS	<b>B1</b>	1.02E+0	2. <b>9</b> E-5	010421
	ORAL [SLOPE] CO	MMENT: UNDER REVIE	W, NUMBER SUBJECT TO	D CHANGE.				
	INHALATION: INTERMITTENT	2 YEARS RAT	BLOOD BRAIN	GLIONA GLIONA	81	3.5E-1	1E-4	010422

INHALATION (UNIT RISK) COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

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						[SLOP	E FACTOR]	[UNIT R	(SK)	
CHENICAL	ROUTE	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	ORAL	INHALATION )' <sup>1</sup> (mg/kg/day)' <sup>1</sup>	ORAL (ug/L) <sup>-1</sup>	INHALATION (ug/cu m)-1	REFERENCE
ETHYLEN	E THIOUREA	000096-	45-7							
	ORAL: GAVAGE	2 YEARS MOUSE	LIVER	ADENONA/ CARCINONA,	B2 , COMBINED	1.1E-1		3.4E-6		010947
	GENERAL COMMENT	: ALSO SEE HEAST	TABLE 1: SUBCHRONIC	AND CHRONIC	TOXICITY	COTHER THAN	CARCINOGENICITY)			
FOLPET		000133-07-3								
	GENERAL COMMENT	: ALSO SEE HEAST	TABLE 1: SUBCHRONIC	AND CHRONIC	IRIS TOXICITY	IRIS (OTHER THAN	CARCINOGENICITY).	IRIS		010120
FORMAL	DEHYDE INHALATION	000050-0 24 MONTHS RAT	0-0 Nasal Cavity	TUNORS	IRIS		4.5E-2		IRIS	010121
	INHALATION (SLOF GENERAL COMMENT:	PE] COMMENT: SEE /	APPENDIX A-II: DOSE TABLE 1: SUBCHRONIC	CONVERSIONS AND CHRONIC	ON HEAST. TOXICITY	OTHER THAN	CARCINOGENICITY).			
FURAZOL	IDONE	000067-45	i <b>-8</b>							
	ORAL: DIET	45 WEEKS RAT	HAPPIARY	TUMORS	82	3.8E+0		1E-4		005106
	GENERAL COMMENT:	FORMERLY LISTED	UNDER NITROFURANS							
FURIUM	ORAL: DIET	000531-82-8 28 WEEKS NOUSE	BLOOD	LEUKENIA	B2	5.0E+1		1.4E-3		005108
	GENERAL COMMENT:	FORMERLY LISTED	UNDER NITROFURANS							
GLYCIDAI	LDEHYDE	000765-3	4-4							
	GENERAL COMMENT:	: ALSO SEE HEAST 1	ABLE 1: SUBCHRONIC	AND CHRONIC	IRIS TOXICITY	OTHER THAN	CARCINOGENICITY).			010122

**ISLOPE FACTOR1 CUNIT RISKI** EXPERIMENT LENGTH **TEPA** ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL ROUTE SPECIES TARGET CANCER (ug/L)<sup>-1</sup> SROUP] (mg/kg/dey)<sup>-1</sup>(mg/kg/dey)<sup>-1</sup> (ug/cu m)-1 **HEPTACHLOR** 000076-44-8 ORAL: DIET IRIS IRIS 4.5E+0 IRIS IRIS 005820 MOLISE LIVER CARCINONA INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-11, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE MEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **HEPTACHLOR EPOXIDE** 001024-57-3 ORAL: DIET **18-24 MONTHS** IRIS IRIS 9.1E+0 IRIS IRIS 010424 MOUSE LIVER CARCINONA INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON MEAST. GENERAL COMMENT: ALSO SEE NEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **HEXACHLOROBENZENE** 000118-74-1 ORAL: DIET IRIS IRIS 1.6E+0 IRIS IRIS 010365 RAT LIVER TIMORS INHALATION (SLOPE) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE MEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **HEXACHLOROBUTADIENE** 000087-68-3 **22-24 MONTHS** IRIS IRIS 7.8E-2 IRIS IRIS 005088 ORAL: DIET KIDNEY **TUMORS** INHALATION (SLOPE) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). HEXACHLOROCYCLOHEXANE, ALPHA- 000319-84-6 24 WEEKS 6.3E+0 IRIS 010123 ORAL: DIET IRIS IRIS IRIS MOLISE LIVER TURORS

INHALATION (SLOPE) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

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CHEMICAL	ROUTE EXT	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE   ORAL (mg/kg/day) <sup>-1</sup> (	INHALATION	[UNIT R ORAL (ug/L) <sup>-1</sup>	ISK] INHALATION (ug/cu m)-1	REFERENCE
HEXACH	LOROCYCLOHE	XANE, BETA-	000319-85-7							
	ORAL: DIET	110 WEEKS Mouse	LIVER	TUMORS	IRIS	IRIS	1.8E+0	IRIS	IRIS	010124
	INHALATION (SLO	PE] COMMENT: BASE	ON ROUTE TO ROUTE	EXTRAPOLATION	. SEE AP	PENDIX A-II, D	OSE CONVERSIONS	ON HEAST.		
UEVACU		VANE DELTA	000010 00 0							•
REXACT	LORUCTCLUME	XANE, DELTA-	000319-86-8		IRIS					010125
	GENERAL COMMENT	: ALSO SEE HEAST 1	ABLE 1: SUBCHRONIC	AND CHRONIC TO	OXICITY	(OTHER THAN CA	ACTINOGENICITY).			
HEXACH	LOROCYCLOHE	XANE, EPSILON-	006108-10-7							
		•			IRIS					010126
	GENERAL COMMENT	: ALSO SEE HEAST	TABLE 1: SUBCHRONIC	AND CHRONIC TO	OXICITY	(OTHER THAN C	ARCINOGENICITY).			
HEXACH	LOROCYCLOHE	XANE, GAMMA-	000058-89-9							
	ORAL: DIET	110 WEEKS MOUSE	LIVER	TUMORS	82-C	1.3E+0		3.7E-5		005098
	OBAL TOLONEL CO		== - <del></del>							
	GENERAL COMMENT	: ALSO SEE HEAST	EV, NUMBER SUBJECT 1 TABLE 1: SUBCHRONIC	AND CHRONIC TO	OXICITY	(OTHER THAN C	ARCINOGENICITY).			
LIEVAGU										
HEXACH	LURUCYCLOME ORAL: DIET	XANE-TECHNICA 6-20 MONTHS	AL 000608-73-1		IRIS	IRIS	1.8E+0	IRIS	IRIS	010127
	GIOLE. DIEI	MOUSE	LIVER	TUMORS	1619	IKIS	1.02*0	1619	IKIƏ	010127
	INHALATION (SLO	PE] COMMENT: BASE	ON ROUTE TO ROUTE	EXTRAPOLATION	. SEE AP	PENDIX A-II, E	OOSE CONVERSIONS	ON KEAST.		
HEXACH	LOROETHANE	00006	7-72-1							
	ORAL: GAVAGE	78 WEEKS MOUSE	LIVER	CARCINONA	IRIS	IRIS	1.4E-2	IRIS	IRIS	005090
			ON ROUTE TO ROUTE					ON HEAST.		

						[SLOPE	FACTOR]	CUNIT I	riskj	
CHEMICAL	ROUTE <u>Đ</u>	CPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	ORAL (mg/kg/day) <sup>-1</sup> (	INHALATION (mg/kg/day) <sup>-1</sup>	ORAL (ug/L) <sup>-1</sup>	INHALATION (ug/cu m)-1	REFERENCE
HYDRAZI	NE	000302-01-	2							
					IRIS	IRIS		IRIS		010129
	INHALATION: INTERMITTENT	1 YEAR RAT	NASAL CAVITY	TUMORS	IRIS		1.7E+1		IRIS	010128
	INHALATION [SL	OPE] COMMENT: SEE	APPENDIX A-II, DOSE	CONVERSIONS O	W WEAST.					
HYDRAZI	NE SULFATE	010034	-93-2							
					IRIS	IRIS		IRIS		010131
	INHALATION: INTERMITTENT	1 YEAR RAT	MASAL CAVITY	TUMORS	IRIS		1.7E+1		IRIS	010130
		OPE] COMMENT: SEE T: LISTED UNDER "N		CONVERSIONS O	W NEAST.					
INDENO[	1,2,3-CD]PYRE	NE 00019	3-39-5							
					IRIS					010192
ISOPHOR	ONE	000078-59	-1					IRIS		005094
	GENERAL COMMEN	T: ALSO SEE HEAST	TABLE 1: SUBCHRONIC	C AND CHRONIC 1	IRIS TOXICITY	IRIS (OTHER THAN CA	RCINOGENICITY).	IKI2		003094
LEAD		007439-92-1								005096
	GENERAL COMMEN	T: ALSO SEE HEAST	TABLE 1: SUBCHRONIC	C AND CHRONIC 1	IRIS	(OTHER THAN CA	RCINOGENICITY).			<del>003090</del>
LINURON	l	000330-55-2								040787
	GENERAL COMMEN	T: ALSO SEE HEAST	TABLE 1: SUBCHRONIC	C AND CHRONIC 1	IRIS FOXICITY	(OTHER THAN C	ARCINOGENICITY).			010383

**March 1994** 

CHEMICAL	ROUTE	<u>DOPERIMENT LENGTH</u> SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL INHAL (mg/kg/day) <sup>-1</sup> (mg/kg/day		
метнох	(Y-5-NITROAN ORAL: DIET	ILINE, 2- 000 104 WEEKS RAT	)099-59-2 skin	CARCINONA	B2	4.6E-2	1.3E-6	010132
METHYL	HYDRAZINE GENERAL CONNE		0-34-4 CLASSIFICATION W	AS PROVIDED IN THE	REFERENC	E DOCUMENT, THEREFORE	THIS COMPOUND WAS REMOVED FROM	N TABLE 3.
METHYL	-5-NITROANIL ORAL: DIET	INE, 2- 0000 98 WEEKS HOUSE	)99-55-8 LIVER	CARCINONA	С	3.3E-2	9.4E-7	010140
METHYL	ANILINE HYDF	ROCHLORIDE, 2- 93 WEEKS RAT	000636-21-! skin	5 FIBROMA	82	1.8E-1	3.1E-6	010134
METHYL	ANILINE, 2- ORAL: DIET	000095 93 Weeks rat	-53-4 skin	FIBROMA	B2	2.4E-1	6.9E-6	010133
	GENERAL COPPLE	NT: THE 1984 HEEP	CALLED THIS COM	POUND O-TOLUIDINE,	THE 1967	NEEP CALLED IT 2-METH	YLANILINE.	
METHYL	CHOLANTHR/	•	00056-49-5	TAPT THE HEALTH ACC	ECCMENT	SECTION (202) 240-4744	FOR RCRA APPROVED NUMERIC AS	010299 SESSMENT OF THIS

METHYLENE CHLORIDE / (DICHLOROMETHANE) 000075-09-2

COMPOUND.

IRIS IRIS IRIS 005100 IRIS 005904

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

**CUNIT RISK** 

March 1994

**ESLOPE FACTOR1** 

**EXPERIMENT LENGTH IEPA** ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL ROUTE SPECIES TARGET CANCER GROUP] (mg/kg/day)<sup>-1</sup> (mg/kg/day)<sup>-1</sup> (ua/L)-1 (ug/cu m)-1

METHYLENE-BIS(BENZENEAMINE), 4,4- / (4,4'-METHYLENEDIANILINE) 000101-77-9

GENERAL COMMENT: NO EPA GROUP CLASSIFICATION WAS PROVIDED IN THE REFERENCE DOCUMENT, THEREFORE THIS COMPOUND WAS REMOVED FROM TABLE 3.

METHYLENE-BIS(2-CHLOROANILINE), 4,4'- 000101-14-4

ORAL: DIET 2 YEARS **B2** 1.3E-1 1.3E-1 3.7E-6 3.7E-5 010425

RAT LUNG TUMORS

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST. BASED ON ROUTE TO ROUTE EXTRAPOLATION.

METHYLENE-BIS(N,N'-DIMETHYL)ANILINE, 4,4'- 000101-61-1

IRIS IRIS IRIS 010137

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

METOLACHLOR 051218-45-2

IRIS 010951

MIREX 002385-85-5

> ORAL: DIET 2 YEARS **B2** 010952

RAT LIVER **ADENONA** LIVER CARCINOMA

[EPA GROUP] COMMENT: UNDER REVIEW, CLASSIFICATION SUBJECT TO CHANGE.

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)

**NIAGARA BLUE 4B** 002429-74-5

> **B2** 010141

[EPA GROUP] COMMENT: BASED ON THE FACT THAT 3,3-DIMETHOXYBENZIDINE, A KNOWN EPA GROUP B2 CARCINOGEN, IS A METABOLITE OF NIAGARA BLUE 48.

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CHEMICAL	ROUTE EXP	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACT ORAL (mg/kg/day) <sup>-1</sup> (mg/	INHALATION	(UNIT RI ORAL (Ug/L) <sup>-1</sup>	SK) INHALATION (ug/cu m)-1	REFERENCE
NICKEL R	EFINERY DUST									
	INHALATION: OCCUPATIONAL	HUMAN	RESPIRATORY SYSTEM	TUMORS	IRIS		8.4E-1		IRIS	005103
			APPENDIX A-II, DOSE AS NICKEL. ALSO SEI			ONIC AND CHRONIC	TOXICITY (UND	ER NICKEL).		
NICKEL S	UBSULFIDE	012035-7	2-2							
	INHALATION: OCCUPATIONAL	HUMAN	RESPIRATORY SYSTEM	TUMORS	IRIS		1.7E+0		IRIS	005768
			APPENDIX A-II, DOSE AS NICKEL. ALSO SE			ONIC AND CHRONIC	TOXICITY (UND	ER NICKEL).		
NITROFU	RAZONE	000059-8	7-0							
	ORAL: DIET	46 WEEKS RAT	HAMMARY	TUMORS	B2	1.5E+0		4.3E-5		005110
NITROPR	OPANE, 2-	000079-4	6-9							
	INHALATION: INTERMITTENT	22 MONTHS RAT	LIVER	TUNORS	82		9.4E+0		2.7E-3	010142
	INHALATION [SLO	PE] COMMENT: SEE	APPENDIX A-II: DOSE	CONVERSIONS O	N HEAST.					
NITROSO	-DI-N-BUTYLAN	AINE N	24-16-3							
MITAUSU	ORAL: DRINKING	LIFETIME	24-10-3		IRIS	IRIS	5.4E+0	IRIS	IRIS	010143
	WATER	MOUSE	BLADDER ESOPHAGUS	TUMORS TUMORS						

INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

CHEMICAL	ROUTE	XPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP] (	[SLOPE FACTOR]  ORAL INHALATION  (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	(UMIT R ORAL (Ug/L) <sup>-1</sup>	ISK] INHALATION (ug/cu m)-1	REFERENCE
NITROSO	-DI-N-PROPYL	AMINE, N- 000	0621-64-7		IRIS	IRIS	IRIS		010147
NITROSO	-N-ETHYLURE/ ORAL: DRINKING WATER	<del>-</del>	59-73-9 Intestine	CASTROINTEST	B2 INAL	1.4E+2			010426

[EPA GROUP] COMMENT: UNDER REVIEW, CLASSIFICATION SUBJECT TO CHANGE. GENERAL COMMENT: FOR RCRA ACTIVITIES ONLY, CONTACT THE HEALTH ASSESSMENT SECTION (202) 382-4761 FOR RCRA-APPROVED NUMERIC ASSESSMENT FOR THIS COMPOUND, UNDER REVIEW.

NITROSO-N-METHYLUREA, N-000684-93-5

> ORAL: GAVAGE 308 DAYS 82 010427 GUINEA PIG PANCREAS **ADENOCARCINONA**

[EPA GROUP] COMMENT: UNDER REVIEW, CLASSIFICATION SUBJECT TO CHANGE.

GENERAL COMMENT: THE CRAVE WORK GROUP (04/01/92) STATES THERE IS NO ACCEPTABLE QUANTITATION FOR NITROSO-N-METHYLUREA, N-.

NITROSODIETHANOLAMINE, N-001116-54-7

IRIS IRIS IRIS 010144

NITROSODIETHYLAMINE, N-000055-18-5

> ORAL: DRINKING 6 OR 12 MONTHS 1.5E+2 IRIS IRIS IRIS IRIS 010145

WATER RAT LIVER TUMORS

INHALATION (SLOPE) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

NITROSODIMETHYLAMINE, N-000062-75-9

> ORAL: DRINKING IRIS IRIS 5.1E+1 IRIS IRIS 010146

WATER LIVER RAT TUMORS

INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

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						(SLOPE FAC		(LWIT R	_	
CHEMICAL	ROUTE	SPECIES	TARGET	CANCER	(EPA GROUP)	ORAL (mg/kg/day) <sup>-1</sup> (mg	INHALATION /kg/day) <sup>-1</sup>	OKAL (ug/L) <sup>-1</sup>	(ug/cu m)-1	REFERENCE
NITROSO	DIPHENYLAM	IINE, N- 000	086-30-6		IRIS	IRIS		IRIS		005112
NITROSO	METHYLETHY	/LAMINE, N- C	)10595-95-6		IRIS	IRIS		IRIS		010148
NITROSO	METHYLVINY INHALATION	'LAMINE, N O	04549-40-0  UPPER RESPIRATORY TRACT	CARCINOMAS	<b>8</b> 2					010149
	ORAL: DRIMKIN WATER	<b>6</b>	UPPER DIGESTIVE TRACT	CARCINOMAS						
NITROSO	PYRROLIDINE	., N- 0009	30-55-2							
	ORAL: DIET	LIFETIME RAT	LIVER	TUMORS	IRIS	IRIS	2.1E+0	IRIS	IRIS	010300
	INHALATION [S	LOPE] COMMENT: BAS	SED ON ROUTE TO ROUTE !	EXTRAPOLATION	. SEE APP	ENDIX A-II, DOS	E CONVERSIONS	ON HEAST.		
PARATHI		000056-38	3-2 t table 1: Subchronic /	AND CHRONIE T	IRIS	OTHER THAN CARE	INOCENICITY			005116
	GENERAL CUTTE	NI: ALSO SEE HEAS	I TABLE 1: SUBCHKUMIC	AND CHRONIC I	OKILIII (	OTHER THAN CARE	INOGENICITY.			
PENTABR	OMO-6-CHLC ORAL: DIET	OROCYCLOHEXA 2 YEARS RAT	NE, 1,2,3,4,5- 00	0087-84-3 TUMORS	С	2.3E-2		6.6E-7		010150
	ORAL [SLOPE]	COMMENT: BASED ON	RESULTS WITH THE ALPH	A ISOMER.						

						[SLOPE FACTOR]	DWIT R		
CHEMICAL	ROUTE	ERIMENT LENGTH SPECIES	TARGET	CANCER	(EPA GROUP)	ORAL INHALATION (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	ORAL (ug/L) <sup>-1</sup>	INHALATION (ug/cu m)-1	REFERENCE
PENTACH	ILORONITROBE	NZENE 000	082-68-8						
	ORAL	72 WEEKS Mouse	LIVER	TUMORS	C	2.6E-1	7.4E-6		010151
			W, NUMBER SUBJECT 1 ABLE 1: SUBCHRONIC		OXICITY (	OTHER THAN CARCINOGENICITY).			
PENTACH	ILOROPHENOL	00008	7-86-5						
	GENERAL COMMENT:	: ALSO SEE HEAST 1	ABLE 1: SUBCHRONIC	AND CHRONIC TO	IRIS OXICITY (	IRIS OTHER THAN CARCINOGENICITY).	IRIS		010381
PHENYLE	NEDIAMINE, O-	000095	-54-5						
	ORAL: DIET	548 DAYS RAT	LIVER	TUMORS	<b>8</b> 2	4.7E-2	1.3E-6		010152
			VER TUMORS IN RATS TABLE 1: SUBCHRONIC			NEDIAMINE DIHYDROCHLORIDE.			
PHENYLP	HENOL, 2-	000090-4	3-7						
	ORAL: DIET	637 DAYS RAT	URINARY BLADDER	TUMORS	C	1.94E-3	5.5E-8		010153
POLYBRO	MINATED BIPH	ENYLS							
	ORAL: GAVAGE	25 VEEKS RAT	LIVER	CARCINONA	82	8.9E+0	2.5E-4		010154
			LIVER	NEOPLASTIC					
	GENERAL COMMENT	: ALSO SEE HEAST 1	TABLE 1: SUBCHRONIC	AND CHRONIC T	OXICITY (	(OTHER THAN CARCINOGENICITY)			
POLYCHI	ORINATED BIP	HENYLS 001	336-36-3		IRIS	IRIS	IRIS		005118

					[SLOPE FACTOR]	QUNIT RI	(UNIT RISK)		
CHEMICAL	ROUTE	PERIMENT LENGTH SPECIES	TARGET	CANCER	(EPA Group)	ORAL INHALATION (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	ORAL (ug/L) <sup>-1</sup>	INHALATION (ug/cu m)-1	REFERENCE
PROPYLE	NE OXIDE	000075-	56-9						
					IRIS	IRIS	IRIS		010156
	INHALATION: INTERMITTENT	2 YEARS MOUSE	NASAL CAVITY	TUMORS	IRIS	1.3E-2		IRIS	010155
	INHALATION (SLO GENERAL COMMENT	PE] COMMENT: SEE : ALSO SEE HEAST	APPENDIX A-II, DOS TABLE 1: SUBCHRONIO	E CONVERSIONS OF CONTRACT TO AND CHRONIC T	N HEAST. OXICITY (	OTHER THAN CARCINOGENICITY)			
QUINOLII	NE	000091-22-5							
G00	ORAL: DIET	20-40 MEEKS RAT	LIVER	TUMORS	c	1.2 <del>E+</del> 1	3.5E-4		010158
RDX / (C	YCLONITE)	000121-8	2-4						
	GENERAL COMMENT	: ALSO SEE HEAST	TABLE 1: SUBCHRONI	C AND CHRONIC T	IRIS OXICITY (	IRIS OTHER THAN CARCINOGENICITY)	IRIS •		010157
SELENIUI	M SULFIDE	007446-3	34-6						040404
	ORAL [SLOPE] CO	MMENT: STUDY RESL	ULTS WERE CONSIDERE	D INCONCLUSIVE	IRIS FOR QUANT	ITATIVE RISK ASSESSMENT.			010194
SIMAZIN	E	000122-34-9							
	ORAL: DIET	2 YEARS RAT	HAMMARY	CARCINONA	C	1.25-1	3.4E-6		010195
			IEW, MUMBER SUBJECT TABLE 1: SUBCHRONI		OXICITY (	OTHER THAN CARCINOGENICITY)	•		
SODIUM	DIETHYLDITHIC	CARBAMATE	000148-18-5						
	ORAL: DIET	77 WEEKS Mouse	LIVER	TUMORS	C	2.7E-1	7.7E-6		005126
	GENERAL COMMENT	: ALSO SEE HEAST	TABLE 1: SUBCHRONI	C AND CHRONIC T	OXICITY (	OTHER THAN CARCINOGENICITY)	) <b>.</b>		

[SLOPE FACTOR] [UNIT RISK]

EXPERIMENT LENGTH

EXPERIMENT LENGTH

CHEMICAL ROUTE

SPECIES

TARGET

CANCER

GROUP] (mg/kg/day)<sup>-1</sup> (ug/l)<sup>-1</sup> (ug/cu m)-1

STYRENE 000100-42-5

010480

GENERAL COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

TCDD, 2,3,7,8- 001746-01-6

ORAL: DIET 720 DAYS B2 1.5E+5 1.5E+5 4.5E+0 3.3E-5 005128 (PG/CU N)-1

RAT RESPIRATORY TUMORS

SYSTEM LIVER TUMORS

ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST.

INHALATION [UNIT RISK] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE. BASED ON ROUTE TO ROUTE EXTRAPOLATION. AN ABSORPTION FACTOR OF 75% IS USED TO CALCULATE THE UNIT RISK FROM THE SLOPE FACTOR.

TETRACHLOROETHANE, 1,1,1,2- 000630-20-6

ORAL: GAVAGE 103 WEEKS IRIS IRIS 2.6E-2 IRIS IRIS 010302

LIVER TUNOR

INHALATION (SLOPE) COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

TETRACHLOROETHANE, 1,1,2,2- 000079-34-5

ORAL: GAVAGE 75 WEEKS IRIS IRIS 2.0E-1 IRIS IRIS 005130

MOUSE LIVER CARCINOMA

INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.

TETRACHLOROETHYLENE 000127-18-4

010482

GENERAL COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

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CHEMICAL	ROUTE EX	PERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACT Oral (mg/kg/day) <sup>-1</sup> (mg/	INHALATION	(UNIT R CRAL (UG/L)-1	ISK) INHALATION (ug/cu m)-1	REFERENCE
TETRACH	HLOROTOLUENI ORAL: GAVAGE	E, PARA, ALPHA, 17.5 VEEKS MOUSE	, ALPHA, ALPHA- LUNG	005216 ADENOCARCIN	B2	2.0E+1		5.7E-4		005028
TETRACH	HLOROVINPHOS ORAL: DIET	S / (STIROFOS) 560 days Mouse	000961-11-5 LIVER	TUMORS	c	2.4E-2		6. <b>9</b> E-7		010159
	[EPA GROUP] COM GENERAL COMMENT	MENT: UNDER REVIEW : ALSO SEE HEAST 1	, CLASSIFICATION SU TABLE 1: SUBCHRONIC	BJECT TO CHAN AND CHRONIC TO	Œ. OXICITY (	OTHER THAN CARCI	NOGENICITY).			
TOLUENE	E-2,4-DIAMINE ORAL: DIET	000095- 84-103 WEEKS RAT	80-7 Hammary	TUMORS	<b>B2</b>	3.2E+0		9.1E-5		010160
TOLUIDIN	NE, P- ORAL: DIET	000106-49-0 18 MONTHS MOUSE	LIVER	TUMORS	С	1.9E-1		5.4E-6		010162
ТОХАРНІ	ORAL: DIET	008001-35- 18 MONTHS MOUSE	LIVER	TUMORS	IRIS	IRIS	1.1E+0	IRIS	IRIS	005134
			APPENDIX A-II, DOSE BASED ON ROUTE TO RO							
TRICHLO	ROANILINE HYD ORAL: DIET	DROCHLORIDE, 2 18 MONTHS MOUSE	2,4,6- C	)33663-50-2 Tumors	2 C	2.9E-2		8.2E-7		005142
TRICHLO	ROANILINE, 2,4 ORAL: DIET	,6- 000634 18 MONTHS MOUSE	-93-5 vascular system	TUMORS	С	3.4E-2		1E-6		010487

1.1E-6

[SLOPE FACTOR] **(UNIT RISK)** EXPERIMENT LENGTH (EPA ORAL INHALATION ORAL INHALATION REFERENCE CHEMICAL ROUTE SPECIES TARGET CANCER GROUP] (mg/kg/day) (mg/kg/day) 1 (ug/L)<sup>-1</sup> (ug/cu m)-1 TRICHLOROETHANE, 1,1,2-000079-00-5 5.7E-2 005144 ORAL: GAVAGE 78 WEEKS IRIS IRIS IRIS IRIS NOUSE LIVER CARCINONA INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). TRICHLOROETHYLENE 000079-01-6 010483 GENERAL COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300. TRICHLOROPHENOL, 2,4,6-000088-06-2 1E-2 IRIS IRIS 010428 ORAL: DIET 107 WEEKS IRIS IRIS MOUSE BLOOD LEUKEMIA INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). TRICHLOROPROPANE, 1,2,3-000096-18-4 ORAL: GAVAGE **B2** 7E+0 Æ-4 010849 RAT MULTIPLE SITES TUMORS, BENIGN/MALIGNANT, [EPA GROUP] COMMENT: UNDER REVIEW, CLASSIFICATION SUBJECT TO CHANGE. GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY). **TRIFLURALIN** 001582-09-8 IRIS IRIS 010163 GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).

**B2** 

3.7E-2

**TUMORS** 

000512-56-1

**UTERUS** 

103 WEEKS MOUSE

TRIMETHYL PHOSPHATE

ORAL: GAVAGE

010164

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CHEMICAL	ROUTE	XPERIMENT LENGTH SPECIES	TARGET	CANCER	(EPA GROUP)	[SLOPE FACTOR] ORAL INHALATION (mg/kg/day) <sup>-1</sup> (mg/kg/day) <sup>-1</sup>	[UNIT R ORAL (ug/L) <sup>-1</sup>	ISK] INHALATION (ug/cu m)-1	REFERENCE
TRINITRO	TOLUENE, 2,4			RONIC AND CHRONIC TO	IRIS KICITY	IRIS (OTHER THAN CARCINOGENICITY).	IRIS		010476
VINYL CH	ORAL: DIET	000075-01 1001 days Rat	LUNG LIVER	TUMORS TUMORS	A	1.9E+0	5.4E-5		010368
	ORAL [SLOPE] ( INHALATION: INTERMITTENT	COMMENT: UNDER REVII  1 YEAR RAT	EV, NUMBER SUB Liver	JECT TO CHANGE. TUNORS	A	3.0E-1		8.4E-5	010367

INHALATION (SLOPE) COMMENT: SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST.

INHALATION [UNIT RISK] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.

GENERAL COMMENT: THE MOST RECENTLY REVIEWED QUANTITATIVE TOXICITY VALUES LISTED HERE APPEAR IN EPA DOCUMENTS PUBLISHED IN 1984 AND 1985. THE AGENCY IS AWARE THAT THESE VALUES DO NOT INCORPORATE CONSIDERABLE INFORMATION THAT IS NOW AVAILABLE. THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT'S POSITION IS THAT THESE TOXICITY VALUES DO NOT REFLECT STATE-OF-THE-ART SCIENCE FOR VINYL CHLORIDE. EPA NOW HAS INDIVIDUAL ANIMAL DATA, NOT AVAILABLE WHEN THE ORAL UNIT RISK WAS CALCULATED, THAT MAY INFLUENCE THIS VALUE. ADDITIONAL INFORMATION THAT MAY BE FACTORED INTO A REVISED QUANTITATIVE TOXICLES DATA ON INCREASED SENSITIVITY OBSERVED IN YOUNG ANIMALS AND DATA ON METABOLISM/PHARMACOKINETICS. A UNIT RISK FOR AIR THAT CONSIDERS INFORMATION ON YOUNG AGE EXPOSURE INCREASES THE RISK (I.E., LOWERS THE RISK SPECIFIC DOSE) BY AT LEAST 3-FOLD. THE CONSIDERATION OF METABOLISM PHARMACOKINETICS WILL FURTHER INCREASE THE RISK. ONE UNPUBLISHED PHYSIOLOGICALLY-BASED PHARMACOKINETIC MODEL PREDICTION RESULTS IN A 100-FOLD INCREASED RISK.

**ACEPHATE** 

030560-19-1

010086 CHEVRON CHEMICAL COMPANY. 1982. MRID NO 00105197. AVAILABLE FROM EPA. WRITE TO FOI, EPA, MASHINGTON, DC 20460.

US EPA. 1984. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR ACEPHATE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE, WASHINGTON, DC.

US EPA. 1988. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

**ACROLEIN** 

000107-02-8

005001 US EPA. 1987. HEALTH EFFECTS ASSESSMENT FOR ACROLEIN. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON DC.

US EPA. 1992. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

**ACRYLAMIDE** 

000079-06-1

010087 JOHNSON K, S GORZINSKI, KM BODNER, ET AL. 1986. CHRONIC TOXICITY AND ONCOGENICITY STUDY ON ACRYLAMIDE INCORPORATED IN THE DRINKING WATER OF FISCHER 344 RATS. DOW CHEMICAL, USA, MIDLAND, MI.

US EPA. 1985. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR ACRYLAMIDE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON, DC.

US EPA. 1988. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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# **ACRYLONITRILE**

### 000107-13-1

005004 BIO/DYNAMICS, INC. 1980. A 24-MONTH ORAL TOXICITY/CARCINOGENICITY STUDY OF ACRYLONITRILE ADMINISTERED IN DRINKING WATER TO FISCHER 344 RATS. FINAL REPORT, VOL 1-4. PREPARED BY BIO/DYNAMICS, INC., DIVISION OF BIOLOGY AND SAFETY EVALUATION, EAST MILLSTONE, NJ. UNDER PROJECT NO 77-1744 (BDN-77-27) FOR MONSANTO COMPANY, ST LOUIS, MO.

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000107-06-2

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# **DIMETHYLANILINE HYDROCHLORIDE, 2,4-**

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# **DIMETHYLANILINE, 2,4-**

000095-68-1

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# DIMETHYLBENZ[A]ANTHRACENE, 7,12-

000057-97-6

010297 FOR RCRA ACTIVITIES ONLY, CONTACT THE HEALTH ASSESSMENT SECTION (202) 260-4761 FOR RCRA APPROVED NUMERIC ASSESSMENT OF THIS COMPOUND.

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# DIMETHYLSULFATE

000077-78-1

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March 1994

# **DIRECT BLACK 38**

### 001937-37-7

010113 NCI (NATIONAL CANCER INSTITUTE). 1978. 13-WEEK SUBCHRONIC TOXICITY STUDIES ON DIRECT BLUE 6, DIRECT BLACK 38 AND DIRECT BROWN 95 DYES. NCI CARCINOGENESIS TECH REP SER NO. 108. P 127.

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## **DIRECT BLUE 6**

## 002602-46-2

010114 NCI (NATIONAL CANCER INSTITUTE). 1978. 13-WEEK SUBCHRONIC TOXICITY STUDIES OF DIRECT BLUE 6, DIRECT BLACK 38, AND DIRECT BROWN 95 DYES. NCI CARCINOGENESIS. TECH REP SER NO. 108. P 127.

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## 002385-85-5

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### **NIAGARA BLUE 4B**

### 002429-74-5

010141 US EPA. 1987. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR NIAGARA BLUE 4B. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON, DC.

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### **DIRECT SKY BLUE 6B**

### 002610-05-1

010116 US EPA. 1987. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR DIRECT SKY BLUE 68. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON. DC.

# **EPICHLOROHYDRIN**

## 000106-89-8

010117 LASKIN S, AR SELLAKUMAR, N KUSCHNER, ET AL. 1980. INHALATION CARCINOGENICITY OF EPICHLOROHYDRIN IN NON-INBRED SPRAGUE-DAWLEY RATS. J NATL CANCER INST. 65(4): 751-757.

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US EPA. 1986. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

### ETHYL ACRYLATE

#### 000140-88-5

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## ETHYLENE OXIDE

### 000075-21-8

010421 DUNKELBERG H. 1982. CARCINOGENICITY OF ETHYLENE OXIDE AND 1,2-PROPYLENE OXIDE UPON INTRA-GASTRIC ADMINISTRATION TO RATS. BR J CANCER. 46: 924-933.

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# **ETHYLENE THIOUREA**

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#### HYDRAZINE

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#### March 1994

# **HYDRAZINE SULFATE**

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010130 MACEMEN JD, EN VERNOT, CC HAUN, ER KINKEAD AND A HALL, 111. 1981. CHRONIC INHALATION TOXICITY OF HYDRAZINE: ONCOGENIC EFFECTS. AIR FORCE AEROSPACE MEDICAL RESEARCH LABORATORY, WRIGHT-PATTERSON AIR FORCE BASE, OH.

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000193-39-5

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LINURON

000330-55-2

010383 REFER TO IRIS.

US EPA. 1989. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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000099-59-2

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000099-55-8

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US EPA. 1987. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR 2-METHYL-5-NITROANILINE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE. WASHINGTON, DC.

METHYLANILINE HYDROCHLORIDE, 2-

000636-21-5

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**METHYLANILINE, 2-**

000095-53-4

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METHYLCHOLANTHRACENE, 3-

000056-49-5

010299 FOR RCRA ACTIVITIES ONLY, CONTACT THE HEALTH ASSESSMENT SECTION (202) 260-4761 FOR RCRA APPROVED NUMERIC ASSESSMENT OF THIS COMPOUND.

# METHYLENE CHLORIDE / (DICHLOROMETHANE)

000075-09-2

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US EPA. 1989. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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US EPA. 1989. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

# METHYLENE-BIS(2-CHLOROANILINE), 4,4'-

000101-14-4

010425 KOMMINENI C, DH GROTH, IJ FROCKT, RW VOELKER AND RP STANOVICK. 1979. DETERMINATION OF THE TUMORIGENIC POTENTIAL OF METHYLENE-BIS-ORTHO-CHLOROANILINE. J ENVIRON PATH TOXICOL. 2(5): 149-171.

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March 1994

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## **STYRENE**

## 000100-42-5

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TETRACHLOROETHYLENE

000127-18-4

010482 CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.

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## 000095-80-7

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US EPA. 1987. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	r Unit Intake or Exposure
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi
Actinium (89)	Ac-225 <sup>†</sup>	014265-85-1	1.000E+01	D	Y	1.0E-03	1.7E-11	2.4E-09	7.6E-09
	Ac-227 <sup>†</sup>	014952-40-0	2.180E+01	Υ	Υ	1.0E-03	2.8E-10	8.0E-08	2.6E-11
	Ac-227+D	014952-40-0(+D)	2.180E+01	Υ	Y	1.0E-03	3.5E-10	8.8E-08	8.5E-07
	Ac-228 <sup>†</sup>	014331-83-0	6.130E+00	н	Y	1.0E-03	5.0E-13	2.6E-11	2.9E-06
Americium (95)	Am-241 <sup>†</sup>	014596-10-2	4.320E+02	Υ	W	1.0E-03	2.4E-10	3.2E-08	4.9E-09
	Am-242	013981-54-9	1.600E+01	н	w	1.0E-03	3.6E-13	1.2E-11	5.8E-09
	Am-242m	013981-54-9(m)	1.520E+02	Υ	W	1.0E-03	2.3E-10	2.8E-08	1.2E-10
	Am-243 <sup>†</sup>	014993-75-0	7.380E+03	Υ	w	1.0E-03	2.4E-10	3.2E-08	2.4E-08
	Am-243+D	014993-75-0(+D)	7.380E + 03	Υ	W	1.0E-03	2.4E-10	3.2E-08	2.5E-07
Antimony (51)	Sb-122	014374-79-9	2.700E+00	D	W	1.0E-01	2.0E-12	3.4E-12	1.4E-06
	Sb-124	014683-10-4	6.020E+01	D	W	1.0E-01	2.9E-12	2.2E-11	6.5E-06
	Sb-125	014234-35-6	2.770E+00	Υ	w	1.0E-01	8.4E-13	1.1E-11	1.2E-06
	Sb-126	015756-32-8	1.240E+01	D	w	1.0E-01	2.8E-12	8.8E-12	9.1E-06
	Sb-126m	015756-32-8(m)	1.900E+01	М	w	1.0E-01	7.2E-14	2.6E-14	5.1E-06
	Sb-127	013968-50-8	3.850E+00	D	w	1.0E-01	2.0E-12	4.1E-12	2.1E-06
	Sb-129	014331-88-5	4.400E+00	Н	w	1.0E-01	5.9E-13	5.1E-13	4.9E-06
Argon (18)	Ar-41	014163-25-8	1.830E+00	Н	*	1.0E+00		5.8E-16	4.4E-06
Astatine (85)	At-217 <sup>†</sup>	017239-90-6	3.230E-02	s	D	1.0E + 00	4.5E-18	5.6E-17	7.7E-10

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil		
Barium (56)	Ba-131	014914-75-1	1.180E+01	D	D	1.0E-01	4.8E-13	3.6E-13	1.2E-06		
	Ba-133	013981-41-4	1.050E+01	Υ	D	1.0E-01	1.2E-12	3.6E-12	8.4E-07		
	Ba-133m	013981-41-4(m)	3.890E+01	Н	D	1.0E-01	6.2E-13	3.5E-13	9.5E-08		
	Ba-137m†	013981-97-0(m)	2.550E+00	M	D	1.0E-01	2.4E-15	6.0E-16	2.0E-06		
	Ba-139	014378-25-7	8.310E+01	М	D	1.0E-01	2.1E-13	1.5E-13	7.7E-08		
	Ba-140	014798-08-4	1.280E+01	D	D	1.0E-01	2.7E-12	2.0E-12	5.4E-07		
Beryllium (4)	Be-7	013966-02-4	5.340E+01	D	Y	5.0E-03	3.0E-14	2.7E-13	1.5E-07		
Bismuth (83)	Bi-206	015776-19-9	6.240E+00	D	w	5.0E-02	2.2E-12	4.3E-12	1.1E-05		
	Bi-207	013982-38-2	3.340E+01	Υ	W	5.0E-02	1.4E-12	1.8E-11	4.9E-06		
	Bi-210 <sup>†</sup>	014331-79-4	5.010E+00	D	w	5.0E-02	1.6E-12	8.0E-11	0.0E+00		
	Bi-211 <sup>†</sup>	015229-37-5	2.130E+00	М	W	5.0E-02	1.2E-14	1.9E-13	1.3E-07		
	Bi-212 <sup>†</sup>	014913-49-6	6.055E+01	М	w	5.0E-02	3.1E-13	6.6E-12	5.9E-07		
	Bi-213 <sup>†</sup>	015776-20-2	4.570E+01	М	W	5.0E-02	2.3E-13	3.0E-13	4.1E-07		
	Bi-214 <sup>†</sup>	014733-03-0	1.990E+01	М	w	5.0E-02	1.3E-13	2.1E-12	5.3E-06		
Bromine (35)	Br-82	014686-69-2	3.530E+01	Н	D	1.0E + 00	1.1E-12	8.7E-13	8.9E-06		
Cadmium (20)	Cd-109	014109-32-1	4.640E+02	D	Υ	5.0E-02	7.9E-12	6.5E-11	7.3E-10		
	Cd-115	014336-68-6	5.350E+01	н	Y	5.0E-02	1.7E-12	2.6E-12	6.3E-07		
	Cd-115m	014336-68-6(m)	4.460E+01	D	Υ	5.0E-02	5.2E-12	3.9E-11	7.5E-08		

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
lement Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class	GI Absorption Factor (f₁) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi		
Calcium (20)	Ca-45	013966-05-7	1.630E+02	D	w	3.0E-01	9.9E-13	5.1E-12	5.8E-18		
	Ca-47	001439-99-2	4.540E+00	D	w	3.0E-01	2.0E-12	4.6E-12	3.6E-06		
Carbon (6)	C-11	014333-33-6	2.050E+01	М	D	1.0E+00	4.9E-14	2.1E-14	3.2E-06		
	C-14	014762-75-5	5.730E+03	Υ	*	1.0E+00	9.0E-13	6.4E-15	0.0E+00		
	C-15	015929-23-4	2.450E+00	s	D	1.0E + 00	8.1E-16	2.1E-16			
Cerium (58)	Ce-141	013967-74-3	3.250E+01	D	Υ	3.0E-04	8.3E-13	8.4E-12	1.3E-07		
	Ce-143	014119-19-8	3.300E+01	Н	Y	3.0E-04	1.3E-12	2.2E-12	6.7E-07		
	Ce-144	014762-78-8	2.840E+02	D	Υ	3.0E-04	6.1E-12	3.4E-10	2.5E-08		
Cesium (55)	Cs-131	014914-76-2	9.690E+00	D	D	1.0E + 00	1.4E-13	1.0E-13	2.8E-09		
	Cs-134	013967-70-9	2.060E+00	Υ	D	1.0E + 00	4.1E-11	2.8E-11	5.2E-06		
	Cs-134m	013967-70-9(m)	2.900E+00	Н	D	1.0E + 00	4.1E-14	3.9E-14	2.0E-08		
	Cs-135	015726-30-4	2.300E+06	Υ	D	1.0E+00	4.0E-12	2.7E-12	0.0E + 00		
	Cs-136	014234-29-8	1.320E+01	D	D	1.0E + 00	6.7E-12	4.6E-12	7.2E-06		
	Cs-137 <sup>†</sup>	010045-97-3	3.020E+01	Υ	D	1.0E + 00	2.8E-11	1.9E-11	0.0E + 00		
	Cs-137+D	010045-97-3(+D)	3.020E+01	Υ	D	1.0E + 00	2.8E-11	1.9E-11	2.0E-06		
	Cs-138	015758-29-9	3.220E+01	M	D	1.0E + 00	1.9E-13	9.6E-14	8.3E-06		
Chlorine (17)	CI-36	013981-43-6	3.010E+05	Υ	D	1.0E + 00	1.8E-12	1.4E-12	0.0E+00		
	CI-38	014158-34-0	3.720E+01	М	D	1.0E + 00	2.3E-13	1.3E-13	5.7E-06		

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

					IODD	GI	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil	
Chromium (24)	Cr-51	014392-02-0	2.770E+01	D	Y	1.0E-01	4.3E-14	3.0E-13	9.2E-08	
Cobalt (27)	Co-57	013981-50-5	2.710E+02	D	Υ	3.0E-01	5.8E-13	8.2E-12	1.9E-07	
	Co-58	01381-38-9	7.080E + 01	D	Y	3.0E-01	1.6E-12	9.8E-12	3.3E-06	
	Co-58m	01381-38-9(m)	9.150E+00	н	Y	3.0E-01	3.2E-14	7.3E-14	4.8E-11	
	Co-60	010198-40-0	5.270E+00	Υ	Y	3.0E-01	1.5E-11	1.5E-10	8.6E-06	
Copper (29)	Cu-64	013981-25-4	1.270E+01	н	Y	5.0E-01	1.7E-13	2.0E-13	6.0E-07	
Curium (96)	Cm-242	015510-73-3	1.630E+02	D	W	1.0E-03	1.3E-11	3.9E-09	3.4E-11	
	Cm-243	015757-87-6	2.850E+01	Y	W	1.0E-03	1.9E-10	2.6E-08	1.6E-07	
	Cm-244	013981-15-2	1.810E+01	Υ	W	1.0E-03	1.6E-10	2.2E-08	3.0E-11	
	Cm-245	015621-76-8	8.500E+03	Υ	W	1.0E-03	2.4E-10	3.2E-08	5.3E-08	
	Cm-246	015757-90-1	4.750E+03	Υ	w	1.0E-03	2.4E-10	3.2E-08	2.7E-11	
	Cm-247	015758-32-4	1.560E+07	Υ	W	1.0E-03	2.2E-10	3.0E-08	9.2E-07	
	Cm-248	015758-33-5	3.390E+05	Y	W	1.0E-03	9.1E-10	1.2E-07	2.2E-11	
Dysprosium (66)	Dy-165	013967-64-1	2.330E+00	Н	W	3.0E-04	1.5E-13	1.1E-13	5.7E-08	
	Dy-166	015840-01-4	8.160E+01	Н	W	3.0E-04	1.9E-12	5.0E-12	2.7E-08	
Erbium (63)	Er-169	015840-13-8	9.400E+00	D	W	3.0E-04	4.4E-13	1.5E-12	8.4E-12	
	Er-171	014391-45-8	7.520E+00	Н	W	3.0E-04	4.6E-13	3.9E-13	9.4E-07	
Europium (63)	Eu-152	014683-23-9	1.360E+01	Υ	W	1.0E-03	2.1E-12	1.1E-10	3.6E-06	

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	r Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f₁) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi
	Eu-154	015585-10-1	8.800E+00	Y	W	1.0E-03	3.0E-12	1.4E-10	4.1E-06
	Eu-155	014391-16-3	4.960E+00	Υ	W	1.0E-03	4.5E-13	1.8E-11	5.9E-08
	Eu-156	014280-35-4	1.520E+01	D	w	1.0E-03	2.5E-12	1.1E-11	4.8E-06
Fluorine (9)	F-18	013981-56-1	1.100E+02	М	D	1.0E+00	9.8E-14	7.1E-14	3.1E-06
Francium (87)	Fr-221 <sup>†</sup>	015756-41-9	4.800E + 00	М	D	1.0E + 00	5.9E-14	9.2E-13	6.2E-08
	Fr-223 <sup>†</sup>	015756-98-6	2.180E+00	М	D	1.0E + 00	1.7E-13	4.2E-13	4.1E-08
Gadolinium (64)	Gd-153	014276-65-4	2.420E+02	D	W	3.0E-04	3.1E-13	5.8E-12	7.3E-08
	Gd-159	014041-42-0	1.860E+01	Н	W	3.0E-04	5.9E-13	6.3E-13	8.9E-08
Gallium (31)	Ga-67	014119-09-6	3.260E+00	D	W	1.0E-03	2.1E-13	3.6E-13	3.3E-07
	Ga-72	013982-22-4	1.410E+01	Н	W	1.0E-03	1.3E-12	1.2E-12	9.8E-06
Germanium (32)	Ge-71	014374-81-3	1.180E+01	D	W	1.0E + 00	6.8E-15	1.3E-13	2.3E-11
Gold (79)	Au-196	014914-16-0	6.180E+00	D	Υ	1.0E-01	3.9E-13	8.7E-13	1.3E-06
	Au-198	010043-49-0	2.700E+00	D	Υ	1.0E-01	1.2E-12	2.1E-12	1.2E-06
Holmium (67)	Ho-166	013967-65-2	2.680E+01	Н	w	3.0E-04	1.7E-12	2.2E-12	6.5E-08
Hydrogen (1)	H-3	010028-17-8	1.230E+01	Υ	*	1.0E + 00	5.4E-14	7.8E-14	0.0E + 00
Indium (49)	In-113m	014885-78-0(m)	1.660E+00	Н	W	2.0E-02	4.9E-14	2.9E-14	7.0E-07
	In-114	013981-55-0	7.190E+01	S	W	2.0E-02	5.5E-15	1.5E-15	1.0E-07
	In-114m	013981-55-0(m)	4.950E+01	D	W	2.0E-02	5.4E-12	4.2E-11	1.8E-07

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil		
	In-115	014191-71-0	4.600E + 15	Υ	w	2.0E-02	3.2E-11	2.2E 10	0.0E + 00		
	In-115m	014191-71-0(m)	4.360E+00	н	w	2.0E-02	1.2E-13	9.4E-14	3.9E-07		
lodine (53)	I-122	018287-75-7	3.620E + 00	М	D	1.0E+00	2.5E-14	7.9E-15	3.0E-06		
	I-123	015715-08-9	1.310E+01	Н	D	1.0E + 00	1.0E-12	5.7E-13	2.4E-07		
	I-125	014158-31-7	6.010E+01	D	D	1.0E + 00	7.9E-12	5.3E-12	2.9E-09		
	I-126	014158-32-8	1.290E+01	D	D	1.0E + 00	1.5E-11	9.8E-12	1.3E-06		
	I-129	015046-84-1	1.570E+07	Υ	D	1.0E + 00	1.9E-10	1.2E-10	4.1E-09		
	I-130	014914-02-4	1.240E+01	Н	D	1.0E+00	9.1E-12	5.1E 12	7.0E-06		
	I-131	010043-66-0	8.040E+00	D	D	1.0E + 00	3.6E-11	2.4E-11	1.5E-06		
	I-132	014683-16-0	2.300E+00	н	D	1.0E + 00	1.0E-12	5.8E·13	7.7E-06		
	I-133	014834-67-4	2.080E+01	Н	D	1.0E+00	2.1E-11	1.2E-11	2.0E-06		
	I-134	014914-27-3	5.260E+01	М	D	1.0E+00	2.8E-13	1.6E-13	9.0E-06		
	I-135	014834-68-5	6.610E+00	Н	D	1.0E + 00	4.2E-12	2.4E-12	5.5E-06		
	Ir-190	014981-91-0	1.180E+01	D	Υ	1.0E-02	1.4E-12	4.8E-12	4.2E-06		
Iridium (77)	lr-192	014694-69-0	7.400E+01	D	Υ	1.0E-02	1.7E-12	2.7E-11	2.4E-06		
	Ir-194	014158-35-1	1.920E+01	н	Y	1.0E-02	1.6E-12	1.8E-12	2.8E-07		
Iron (26)	Fe-55	014681-59-5	2.700E+00	Υ	w	1.0E-01	2.7E-13	8.4E-13	0.0E + 00		
	Fe-59	014596-12-4	4.460E + 01	D	W	1.0E-01	2.8E-12	9.7E-12	4.1E-06		

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

					CDD		Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	er Unit Intake or Exposure
Element (Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	t	CRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi
Krypton (36)	Kr-83m	013965-98-5(m)	1.830E+00	н	*	1.0E + 00		6.2E-17	3.4E-11
	Kr-85	013983-27-2	1.070E+01	Υ	*	1.0E+00		4.7E-17	7.0E-09
	Kr-85m	013983-27-2(m)	4.480E+00	Н	*	1.0E+00		4.9E-16	3.4E-07
	Kr-87	014809-68-8	7.630E+01	М	*	1.0E + 00		2.2E-15	2.8E-06
	Kr-88	014995-61-0	2.840E+00	Н	*	1.0E + 00		4.7E-15	7.3E-06
	Kr-89	016316-03-3	3.160E+00	М	*	1.0E + 00		2.6E-15	6.5E-06
	Kr-90	015741-13-6	3.230E+01	s	*	1.0E+00		2.4E-15	4.2E-06
Lanthanum (57)	La-140	013981-28-7	4.020E+01	Н	W	1.0E-03	2.3E-12	3.0E-12	8.0E-06
Lead (82)	Pb-203	014687-25-3	5.200E+01	Н	D	2.0E-01	3.2E-13	2.6E-13	5.9E-07
	Pb-209 <sup>†</sup>	014119-30-3	3.250E+00	Н	D	2.0E-01	8.5E-14	7.0E-14	0.0E + 00
	Pb-210 <sup>†</sup>	014255-04-0	2.230E+01	Υ	D	2.0E-01	5.1E-10	1.3E-09	1.3E-10
	Pb-210+D	014255-04-0(+D)	2.230E+01	Υ	D	2.0E-01	6.6E-10	4.0E-39	1.6E-10
	Pb-211 <sup>†</sup>	015816-77-0	3.610E+01	М	D	2.0E-01	1.8E-13	2.8E-12	1.6E-07
	Pb-212 <sup>†</sup>	015092-94-1	1.060E+01	Н	D	2.0E-01	5.5E-12	4.3E-11	2.8E-07
	Pb-214 <sup>†</sup>	015067-28-4	2.680E+01	М	D	2.0E-01	1.7E-13	2.9E-12	6.4E-07
Lutetium (71)	Lu-177	014265-75-9	6.710E+00	D	Υ	3.0E-04	6.2E-13	1.9E-12	6.7E-08
Manganese (25)	Mn-52	014092-99-0	5.590E+00	D	W	1.0E-01	2.2E-12	3.7E-12	1.2E-05
	Mn-54	013966-31-9	3.130E+02	D	W	1.0E-01	1.1E-12	5.3E-12	2.9E-06

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	r Unit Intake or Exposure
Element Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi
	Mn-56	014681-52-8	2.580E+00 H	1	W	1.0E-01	4.0E-13	2.8E-13	6.1E-06
Mercury (80)	Hg-197	013981-51-6	6.410E+01 H	н	W	2.0E-02	2.7E-13	4.5E-13	5.5E-08
	Hg-203	013982-78-0	4.660E+01 E	D	W	2.0E-02	6.6E-13	4.8E-12	5.7E-07
Molybdenum (42)	Mo-99	014119-15-4	6.600E+01 H	Н	Υ	8.0E-01	1.5E-12	2.6E-12	4.9E-07
Neodymium (60)	Nd-147	014269-74-0	1.100E+01 [	D	Υ	3.0E-04	1.3E-12	5.5E-12	3.0E-07
	Nd-149	015749-81-2	1.730E+00 H	Н	Υ	3.0E-04	1.9E-13	1.6E-13	9.8E-07
Neptunium (93)	Np-236	015700-36-4	1.150E+05	Υ	W	1.0E-03	2.3E-13	3.9E-12	8.9E-08
	Np-237 <sup>†</sup>	013994-20-2	2.140E+06	Υ	W	1.0E-03	2.2E-10	2.9E-08	7.8E-09
	Np-237 + D	013994-20-2(+D)	2.140E+06	Υ	W	1.0E-03	2.2E-10	2.9E-08	4.3E-07
	Np-238	015766-25-3	2.120E+00	D	W	1.0E-03	1.1E-12	3.3E-12	1.7E-06
	Np-2391	013968-59-7	2.360E+00	D	W	1.0E-03	9.4E-13	1.5E-12	2.3E-07
	Np-240	015690-84-3	6.500E+01	М	W	1.0E-03	1.3E-13	6.5E-14	3.2E-06
	Np-240m	015690-84-3(m)	7.400E+00	М	W	1.0E-03	2.9E-14	9.0E-15	9.3E-07
Nickel (28)	Ni-59	014336-70-0	7.500E+04	Υ	W	5.0E-02	9.1E-14	7.0E-13	0.0E + 00
	Ni-63	013981-37-8	1.000E+02	Υ	W	5.0E-02	2.4E-13	1.8E-12	0.0E + 00
	Ni-65	014833-49-9	2.520E+00	Н	W	5.0E-02	2.6E-13	1.9E-13	1.9E-06
Niobium (41)	Nb-93m	007440-03-1(m)	1.460E+01	Υ	Υ	1.0E-02	1.5E-13	1.9E-11	5.3E-11
	Nb-94	014681-63-1	2.030E+04	Υ	Υ	1.0E-02	2.1E-12	2.1E-10	5.4E-06

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	r Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil
	Nb-95	013967-76-5	3.510E+01	D	Y	1.0E-02	6.5E-13	5.1E-12	2.6E-06
	Nb-95m	013967-76-5(m)	8.660E+01	Н	Y	1.0E-02	6.6E-13	1.6E-12	8.1E-08
	Nb-97	018496-04-3	7.210E+01	М	Υ	1.0E-02	1.2E-13	6.9E-14	2.2E-06
	Nb-97m	018496-04-3(m)	6.000E+01	s	Y	1.0E-02	2.4E-15	1 1E-15	2.5E-06
Osmium (76)	Os-185	015766-50-4	9.360E+01	D	Υ	1.0E-02	6.3E-13	8.9E-12	2.2E-06
	Os-191	014119-24-5	1.540E+01	D	Υ	1.0E-02	6.7E-13	3.6E-12	8.5E-08
	Os-191m	014119-24-5(m)	1.300E+01	Н	Υ	1.0E-02	1.2E-13	2.1E-13	3.4E-09
	Os-193	016057-77-5	3.000E+01	Н	Υ	1.0E-02	9.6E-13	1.2E-12	1.7E-07
Palladium (46)	Pd-100	015690-69-4	3.640E+00	D	Y	5.0E-03	1.0E-12	2.3E-12	
	Pd-101	015749-54-9	8.480E+00	Н	Υ	5.0E-03	1.2E-13	1.1E-13	
	Pd-103	014967-68-1	1.700E+01	D	Υ	5.0E-03	2.2E-13	1.4E-12	7.3E-10
	Pd-107	017637-99-9	6.500E+06	Υ	Y	5.0E-03	4.4E-14	6.4E-12	0.0E + 00
	Pd-109	014981-64-7	1.350E+01	Н	Υ	5.0E-03	7.9E-13	8.1E-13	2.2E-09
Phosphorus (15)	P-32	014596-37-3	1.430E+01	D	D	8.0E-01	3.5E-12	3.0E-12	0.0E + 00
	P-33	015749-66-3	2.540E+01	D	D	8.0E-01	5.6E-13	4.6E-13	0.0E + 00
Platinum (78)	Pt-191	015706-36-2	2.710E+00	D	D	1.0E-02	3.8E-13	3.0E-13	6.3E-07
	Pt-193	015735-70-3	5.000E+01	Υ	D	1.0E-02	3.5E-14	8.2E-14	0.0E + 00
	Pt-193m	015735-70-3(m)	4.330E+00	D	D	1.0E-02	5.3E-13	4.0E-13	7.6E-09

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

				9.	000		Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	L	CRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi	
	Pt-197	015735-74-7	1.830E+01	Н	D	1.0E-02	4.9E-13	3.2E-13	3.1E-08	
	Pt-197m	015735-74-7(m)	9.440E+01	М	D	1.0E-02	1.2E-13	8.7E-14	1.6E-07	
Plutonium (94)	Pu-236	015411-92-4	2.850E+00	Υ	Υ	1.0E-03	5.0E-11	2.4E-08	3.4E-11	
	Pu-238	013981-16-3	8.780E+01	Υ	Υ	1.0E-03	2.2E-10	3.9E-08	2.8E-11	
	Pu-239	015117-48-3	2.410E + 04	Υ	Υ	1.0E-03	2.3E-10	3.8E-08	1.7E-11	
	Pu-240	014119-33-6	6.570E+03	Υ	Υ	1.0E-03	2.3E-10	3.8E-08	2.7E-11	
	Pu-241	014119-32-5	1.440E+01	Υ	Υ	1.0E-03	3.6E-12	2.3E-10	0.0E + 00	
	Pu-242	013982-10-0	3.760E+05	Υ	Υ	1.0E-03	2.2E-10	3.6E-08	2.3E-11	
	Pu-243	015706-37-3	4.960E+00	Н	Υ	1.0E-03	1.1E-13	1.0E-13	1.8E-08	
	Pu-244	014119-34-7	8.260E+07	Υ	Υ	1.0E-03	2.2E-10	3.6E-08	1.9E-11	
Polonium (84)	Po-210 <sup>†</sup>	013981-52-7	1.380E+02	D	W	1.0E-01	1.5E-10	2.6E-09	2.9E-11	
	Po-212 <sup>†</sup>	015389-34-1	2.980E-07	s	W	1.0E-01	2.2E-23	6.1E-22	0.0E + 00	
	Po-213 <sup>†</sup>	015756-57-7	4.200E-06	S	W	1.0E-01	3.2E-22	8.0E-21	1.0E-10	
	Po-214 <sup>†</sup>	015735-67-8	1.640E-04	S	W	1.0E-01	1.0E-20	2.8E-19	2.8E-10	
	Po-215 <sup>†</sup>	015706-52-2	1.780E-03	S	W	1.0E-01	2.8E-19	5.7E-18	4.6E-10	
	Po-216 <sup>†</sup>	015756-58-8	1.460E-01	S	W	1.0E-01	3.0E-17	4.8E-16	5.0E-11	
	Po-218 <sup>†</sup>	015422-24-9	3.050E+00	М	W	1.0E-01	2.8E-14	5.8E-13	0.0E + 00	
Potassium (19)	K-40	013966-00-2	1.280E+09	Υ	Đ	1.0E + 00	1.1E-11	7.6E-12	5.4E-07	

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

						Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	r Unit intake or Exposure
lement Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi
	K-42	014378-21-3	1.240E+01 F	I D	1.0E+00	8.9E-13	1.2E-12	9.6E-07
Praseodymium (59)	Pr-142	014191-64-1	1.910E+01 H	I Y	3.0E-04	1.6E-12	1.8E-12	2.1E-07
	Pr-143	014981-79-4	1.360E+01 D	Υ Υ	3.0E-04	1.4E-12	7.0E-12	3.0E-14
	Pr-144	014119-05-2	1.730E+01 N	1 Y	3.0E-04	9.5E-14	3.6E-14	1.2E-07
	Pr-144m	014119-05-2(m)	7.200E+00 N	1 Y	3.0E-04	3.7E-14	1.6E-14	2.1E-09
Promethium (61)	Pm-147	014380-75-7	2.620E+00 Y	Y	3.0E-04	3.1E-13	3.0E-11	6.0E-12
	Pm-148	014683-19-3	5.370E+00 E	Υ Υ	3.0E-04	3.1E-12	7.9E-12	1.9E-06
	Pm-148m	014683-19-3(m)	4.130E+01 E	Y	3.0E-04	2.5E-12	4.8E-11	6.5E-06
	Pm-149	015765-31-8	5.310E+01 H	I Y	3.0E-04	1.2E-12	1.9E-12	3.3E-08
Protactinium (91)	Pa-231 <sup>†</sup>	014331-85-2	3.730E+04 \	′ Y	1.0E-03	9.2E-11	3.6E-08	2.6E-08
	Pa-233†	013981-14-1	2.700E+01 [	Y	1.0E-03	1.0E-12	8.6E-12	4.2E-07
	Pa-234	015100-28-4	6.700E+00 H	l Y	1.0E-03	6.8E-13	5.4E-13	5.9E-06
	Pa-234m <sup>†</sup>	015100-28-4(m)	1.170E+00 M	И Y	1.0E-03	5.8E-15	1.6E-15	3.6E-08
Radium (88)	Ra-223†	015623-45-7	1.140E+01 [	w w	2.0E-01	6.4E-11	3.1E-09	2.3E-07
	Ra-224 <sup>†</sup>	013233-32-4	3.620E+00 [	) W	2.0E-01	3.8E-11	1.2E-09	2.3E-08
	Ra-225 <sup>↑</sup>	013981-53-8	1.480E+01 [	) W	2.0E-01	5.1E-11	1.5E-09	1.9E-09
	Ra-226 <sup>†</sup>	013982-63-3	1.600E+03 \	v w	2.0E-01	1.2E-10	3.0E-09	1.2E-08
	Ra-226 + D	013982-63-3(+D)	1.600E+03	v w	2.0E-01	1.2E-10	3.0E-09	6.0E-06

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

					1000		Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi		
	Ra-228†	015262-20-1	5.750E+00	Υ	w	2.0E-01	1.0E-10	6.6E-10	0.0E + 00		
	Ra-228 + D	015262-20-1(+D)	5.750E+00	Υ	w	2.0E-01	1.0E-10	6.9E-10	2.9E-06		
Radon (86)	Rn-219 <sup>†</sup>	014835-02-0	3.960E+00	s	*	1.0E+00		4.6E-14	1.6E-07		
	Rn-220 <sup>†</sup>	022481-48-7	5.560E+01	s	*	1.0E+00		1.2E-13	1.7E-09		
	Rn-222 <sup>†</sup>	014859-67-7	3.820E+00	D	*	1.0E+00	1.4E-12	7.3E-13	1.2E-09		
	Rn-222 + D	014859-67-7(+D)	3.820E+00	D	*	1.0E + 00	1.7E-12	7.7E-12	5.9E-06		
Rhodium (45)	Rh-103m	007440-16-6(m)	5.610E+01	М	Υ	5.0E-02	6.9E-15	3.9E-15	7.9E-11		
	Rh-105	014913-89-4	3.540E+01	Н	Y	5.0E-02	4.3E-13	5.9E-13	2.2E-07		
	Rh-105m	014913-89-4(m)	4.500E+01	S	Υ	5.0E-02	6.2E-16	3.4E-16	2.2E-08		
	Rh-106	014234-34-5	2.990E+01	S	Υ	5.0E-02	4.4E-15	1.2E-15	6.7E-07		
Rubidium (37)	Rb-82	014391-63-0	1.250E+00	М	D	1.0E+00	1.2E-14	3.5E-15	3.5E-06		
	Rb-86	014932-53-7	1.870E+01	D	D	1.0E+00	5.9E-12	4.5E-12	3.3E-07		
	Rb-87	013982-13-3	4.730E + 10	Υ	D	1.0E+00	3.4E-12	2.4E-12	0.0E+00		
	Rb-88	014928-36-0	1.780E+01	М	D	1.0E + 00	1.7E-13	7.9E-14	2.4E-06		
	Rb-89	014191-65-2	1.540E+01	М	D	1.0E + 00	9.3E-14	3.9E-14	7.4E-06		
Ruthenium (44)	Ru-97	015758-35-7	2.900E+00	D	Υ	5.0E-02	1.7E-13	2.6E-13	4.2E-07		
	Ru-103	013968-53-1	3.940E+01	D	Υ	5.0E-02	9.0E-13	8.4E-12	1.5E-06		
	Ru-105	014331-95-4	4.440E+00	н	Υ	5.0E-02	3.7E-13	3.3E-13	2.6E-06		

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

					CRP	0.	Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ı	Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi		
	Ru-106	013967-48-1	3.680E+02	D	Y	5.0E-02	9.5E-12	4.4E-10	0.0E+00		
Samarium (62)	Sm-147	014392-33-7	1.060E + 11	Υ	w	3.0E-04	1.6E-11	7.2E-09	0.0E+00		
	Sm-151	015715-94-3	9.000E + 01	Υ	w	3.0E-04	1.1E-13	8.7E-12	4.0E-13		
	Sm-153	015766-00-4	4.670E + 01	Н	w	3.0E-04	8.7E-13	1.3E-12	4.6E-08		
Scandium (21)	Sc-46	013967-63-0	8.380E+01	D	Υ	1.0E-04	1.6E-12	2.7E-11	6.9E-06		
	Sc-47	014391-96-9	3.420E+00	D	Υ	1.0E-04	6.4E-13	1.2E-12	2.3E-07		
	Sc-48	014391-86-7	4.370E+01	н	Υ	1.0E-04	1.9E-12	2.3E-12	1.2E-05		
Selenium (34)	Se-75	014265-71-5	1.200E + 02	D	w	8.0E-01	5.8E-12	6.0E-12	8.1E-07		
Silicon (14)	Si-31	014276-49-4	1.570E+02	М	w	1.0E-02	2.2E-13	1.7E-13	3.0E-09		
Silver (47)	Ag-105	014928-14-4	4.130E+01	D	Υ	5.0E-02	7.3E-13	4.0E-12			
	Ag-108	014391-65-2	2.370E+00	М	Υ	5.0E-02	8.5E-15	2.4E-15	5.1E-08		
	Ag-108m	014391-65-2(m)	1.270E+02	Υ	Υ	5.0E-02	3.5E-12	1.5E-10	5.0E-06		
	Ag-109m	014378-38-2(m)	3.960E+01	S	Υ	5.0E-02	3.3E-16	8.9E-17	1.3E-09		
	Ag-110	014391-76-5	2.460E+01	S	Υ	5.0E-02	3.0E-15	8.1E-16	1.0E-07		
	Ag-110m	014391-76-5(m)	2.500E+02	D	Υ	5.0E-02	4.7E-12	6.9E-11	9.3E-06		
	Ag-111	157690-04-0	7.460E+00	D	Υ	5.0E-02	1.6E-12	4.8E-12	7.7E-08		
Sodium (11)	Na-22	013966-32-0	2.600E + 00	Υ	D	1.0E+00	6.8E-12	4.8E-12	7.2E-06		
	Na-24	013982-04-2	1.500E+01	Н	D	1.0E+00	1.0E-12	9.6E-13	1.6E-05		

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil		
Strontium (38)	Sr-82	014809-50-8	2.500E+01	D	D	3.0E-01	7.5E-12	6.9E-12	1.3E-10		
	Sr-85	013967-73-2	6.480E+01	D	D	3.0E-01	7.7E-13	1.0E-12	1.4E-06		
	Sr-85m	013967-73-2(m)	6.770E+01	М	D	3.0E-01	1.2E-14	6.1E-15	4.8E-07		
	Sr-89	014158-27-1	5.060E+01	D	D	3.0E-01	3.0E-12	2.9E-12	4.7E-10		
	Sr-90 <sup>†</sup>	010098-97-2	2.860E+01	Υ	D	3.0E-01	3.3E-11	5.6E-11	0.0E + 00		
	Sr-90+D	010098-97-2(+D)	2.860E+01	Υ	D	3.0E-01	3.6E-11	6.2E-11	0.0E + 00		
	Sr-91	014331-91-0	9.500E+00	Н	D	3.0E-01	8.5E-13	6.9E-13	2.4E-06		
	Sr-92	014928-29-1	2.710E+00	н	D	3.0E-01	5.7E-13	4.5E-13	4.6E-06		
Sulfur (16)	S-35	015117-53-0	8.740E+01	D	D	8.0E-01	2.2E-13	1.9E-13	0.0E + 00		
Tantalum (73)	Ta-182	013982-00-8	1.150E+02	D	Υ	1.0E-03	1.7E-12	4.3E-11	4.1E-06		
Technetium (43)	Tc-95	014809-56-4	2.000E+01	Н	w	8.0E-01	5.6E-14	2.3E-14	2.4E-06		
	Tc-95m	014809-56-4(m)	6.100E+01	D	W	8.0E-01	1.8E-12	4.0E-12	1.9E-06		
	Tc-96	014808-44-7	4.280E+00	D	W	8.0E-01	1.8E-12	2.1E-12	8.3E-06		
	Tc-96m	014808-44-7(m)	5.150E+01	М	W	8.0E-01	2.3E-14	2.1E-14	7.0E-08		
	Tc-97	015759-35-0	2.600E+06	Υ	W	8.0E-01	1.5E-13	9.8E-13	3.5E-10		
	Tc-97m	015759-35-0(m)	8.900E+01	D	w	8.0E-01	1.1E-12	5.0E-12	3.7E-10		
	Tc-99	014133-76-7	2.130E+05	Υ	w	8.0E-01	1.3E-12	8.3E-12	6.0E-13		
	Tc-99m	014133-76-7(m)	6.020E + 00	н	w	8.0E-01	5.0E-14	2.7E-14	2.3E-07		

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

				1000		Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	er Unit Intake or Exposure
Element (Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi
Tellurium (52)	Te-125m	014390-73-9(m)	5.800E+01 E	w	2.0E-01	8.5E-13	5.4E-12	2.6E-09
	Te-127	013981-49-2	9.350E+00 H	w	2.0E-01	2.3E-13	2.3E-13	1.4E-08
	Te-127m	013981-49-2(m)	1.090E+02 D	w	2.0E-01	2.2E-12	1.6E-11	8.4E-10
	Te-129	014269-71-7	6.960E+01 N	1 W	2.0E-01	1.1E-13	6.8E-14	1.3E-07
	Te-129m	014269-71-7(m)	3.360E+01 E	w	2.0E-01	3.2E-12	2.0E-11	6.3E-08
	Te-131	014683-12-6	2.500E+01 N	ı w	2.0E-01	2.8E-13	1.4E-13	1.2E-06
	Te-131m	014683-12-6(m)	3.000E+01 F	W	2.0E-01	3.8E-12	5.4E-12	4.7E-06
	Te-132	014234-28-7	7.820E+01 F	W	2.0E-01	3.0E-12	5.3E-12	4.0E-07
Terbium (65)	Tb-158	015759-55-4	1.500E+02 Y	w	3.0E-04	1.2E-12	9.4E-11	
	Tb-160	013981-29-8	7.230E+01 D	W	3.0E-04	1.8E-12	1.9E-11	3.6E-06
Thallium (81)	TI-202	015720-57-7	1.220E+01 D	D	1.0E+00	8.4E-13	6.0E-13	1.3E-06
	TI-204	013968-51-9	3.780E+00 Y	D	1.0E + 00	1.7E-12	1.3E-12	8.7E-10
	TI-207†	014133-67-6	4.770E+00 N	i D	1.0E + 00	1.3E-14	4.5E-15	7.5E-09
	TI-208 <sup>†</sup>	014913-50-9	3.050E+00 N	I D	1.0E + 00	1.8E-14	5.0E-15	1.3E-05
	TI-209 <sup>†</sup>	015690-73-0	2.200E+00 N	I D	1.0E+00	1.4E-14	4.3E-15	6.9E-06
Thorium (90)	Th-227 <sup>†</sup>	015623-47-9	1.870E+01 D	Υ	2.0E-04	4.5E-12	4.9E-09	1.6E-07
	Th-2281	014274-82-9	1.910E+00 Y	Υ	2.0E-04	1.1E-11	7.7E-08	5.5E-10
	Th-228+D	014274-82-9(+D)	1.910E+00 Y	Υ	2.0E-04	5.5E-11	7.8E-08	5.6E-06

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

						Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil
	Th-229 <sup>†</sup>	015594-54-4	7.340E+03	Y Y	2.0E-04	2.1E- <u>-</u> 11	7.3E-08	5.8E-08
	Th-229 + D	015594-54-4(+D)	7.340E+03	Y Y	2.0E-04	8.9E-11	7.7E-08	6.8E-07
	Th-230 <sup>†</sup>	014269-63-7	7.700E+04	Y Y	2.0E-04	1.3E-11	2.9E-08	5.4E-11
	Th-231 <sup>†</sup>	014932-40-2	2.550E+01	- Y	2.0E-04	4.0E-13	4.9E-13	2.3E-09
	Th-232 <sup>†</sup>	007440-29-1	1.410E+10	Y Y	2.0E-04	1.2E-11	2.8E-08	2.6E-11
	Th-234 <sup>†</sup>	015065-10-8	2.410E+01	) Y	2.0E-04	4.0E-12	3.2E-11	3.5E-09
Thulium (69)	Tm-170	013981-30-1	1.290E+02	o w	3.0E-04	1.5E-12	2.1E-11	3.8E-09
	Tm-171	014333-45-0	1.920E+00	y W	3.0E-04	1.2E-13	3.1E-12	3.3E-10
Tin (50)	Sn-113	013966-06-8	1.150E+02	o w	2.0E-02	8.7E-13	9.4E-12	3.3E-09
	Sn-121	014683-06-8	2.710E+01	н w	2.0E-02	2.7E-13	3.2E-13	
	Sn-121m	014683-06-8(m)	5.550E+01	y w	2.0E-02	5.4E-13	9.3E-12	
	Sn-125	014683-08-0	9.640E+00	o w	2.0E-02	3.6E-12	1.2E-11	1.1E-06
	Sn-126	015832-50-5	1.000E+05	r w	2.0E-02	5.6E-12	7.7E-11	3.3E-08
Tungsten (74)	W-181	015749-46-9	1.210E+02	О С	3.0E-01	8.3E-14	6.4E-14	2.2E-08
	W-185	014932-41-3	7.510E+01	D D	3.0E-01	4.7E-13	3.0E-13	4.7E-11
	W-187	014983-48-3	2.380E+01	н р	3.0E-01	6.3E-13	3.9E-13	1.5E-06
Uranium (92)	U-232	014158-29-3	7.200E + 01	Y	5.0E-02	3.7E-11	6.0E-08	4.6E-11
	U-233	013968-55-3	1.590E+05	Υ	5.0E-02	1.6E-11	2.7E-08	4.2E-11

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	r Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f₁) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soi	
	U-234†	013966-29-5	2.450E+05	Y	Y	5.0E-02	1.6E-11	2.6E-08	3.0E-11
	U-235†	015117-96-1	7.040E+08	Υ	Y	5.0E-02	1.6E-11	2.5E-08	2.4E-07
	U-235 + D	015117-96-1(+D)	7.040E+08	Υ	Υ	5.0E-02	1.6E-11	2.5E-08	2.4E-07
	U-236	013982-70-2	2.340E+07	Υ	Υ	5.0E-02	1.5E-11	2.5E-08	2.4E-11
	U-237	014269-75-1	6.750E+00	D	Υ	5.0E-02	8.9E-13	2.6E-12	1.3E-07
	U-238 <sup>†</sup>	007440-61-1	4.470E+09	Υ	Υ	5.0E-02	1.6E-11	2.4E-08	2.1E-11
	U-238+D	007440-61-1(+D)	4.470E+09	Y	Y	5.0E-02	2.0E-11	2.4E-08	5.1E-08
	U-240	015687-53-3	1.410E+01	Н	Υ	5.0E-02	1.2E-12	1.2E-12	1.5E-10
Vanadium (23)	V-48	014331-97-6	1.600E+01	D	W	1.0E-02	2.2E-12	7.6E-12	9.9E-06
Xenon (54)	Xe-122	015151-09-4	2.010E+01	Н	*	1.0E + 00		3.3E-15	8.4E-08
	Xe-123	015700-10-4	2.140E+00	Н	*	1.0E+00		7.4E-16	1.7E-06
	Xe-125	013994-18-8	1.680E+01	Н	*	1.0E+00		4.2E-16	4.7E-07
	Xe-127	013994-19-9	3.640E+01	D	*	1.0E + 00		4.0E-16	5.0E-07
	Xe-129m	013965- <del>9</del> 9-6(m)	8.890E+00	D	*	1.0E + 00		6.0E-16	1.3E-08
	Xe-131m	014683-11-5(m)	1.180E+01	D	*	1.0E + 00		4.3E-16	4.5E-09
	Xe-133	014932-42-4	5.250E+00	D	*	1.0E + 00		4.3E-16	2.3E-08
	Xe-133m	014932-42-4(m)	2.190E+00	D	*	1.0E + 00		5.4E-16	3.3E-08
	Xe-135	014995-62-1	9.110E+00	Н	*	1.0E+00		8.0E-16	6.2E-07

Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Picocuries<sup>b</sup>)

						Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil		
	Xe-135m	014995-62-1(m)	1.540E+01 N	1 *	1.0E+00		2.1E-16	1.3E-06		
	Xe-137	014835-21-3	3.830E+00 N	1 *	1.0E+00		1.8E-15	6.0E-07		
	Xe-138	015751-81-2	1.410E+01 N	1 *	1.0E + 00		2.8E-15	4.0E-06		
Yttrium (39)	Y-90 <sup>†</sup>	010098-91-6	6.410E+01 H	Υ	1.0E-04	3.2E-12	5.5E-12	0.0E + 00		
rttilum (39)	Y-91	014234-24-3	5.850E+01 D	Y	1.0E-04	2.8E-12	4.3E-11	1.2E-08		
	Y-91m	014234-24-3(m)	4.970E+01 M	1 Y	1.0E-04	2.2E-14	3.5E-14	1.7E-06		
	Y-92	015751-59-4	3.540E+00 H	ı Y	1.0E-04	7.1E-13	5.3E-13	8.6E-07		
	Y-93	014981-70-5	1.010E+01 H	I Y	1.0E-04	1.4E-12	1.4E-12	3.0E-07		
Zinc (30)	Zn-65	013982-39-3	2.440E+02 [	Y	5.0E-01	8.5E-12	1.6E-11	2.0E-06		
	Zn-69	013982-23-5	5.560E+01 N	Λ Y	5.0E-01	5.6E-14	3.2E-14	1.8E-11		
	Zn-69m	013982-23-5(m)	1.380E+01 H	ı Y	5.0E-01	4.3E-13	5.7E-13	1.3E-06		
Zirconium (40)	Zr-93	015751-77-6	1.530E+06 \	v w	2.0E-03	1.7E-13	6.5E-12	0.0E + 00		
	Zr-95	013967-71-0	6.400E+01 (	w w	2.0E-03	9.9E-13	1.0E-11	2.5E-06		
	Zr-97	014928-30-4	1.690E+01 I	ı w	2.0E-03	2.4E-12	2.5E-12	6.1E-07		

# Table 4A: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> (In Units of Picocuries<sup>b</sup>)

**MARCH 1994** 

						Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)	

# **ENDNOTES:**

<sup>&</sup>lt;sup>a</sup> EPA classifies all radionuclides as Group A (known human) carcinogens. Radionuclide slope factors are calculated by EPA's Office of Radiation and Indoor Air (ORIA) to assist HEAST users with risk-related evaluations and decision-making at various stages of the remediation process. Ingestion and inhalation slope factors are best estimates (i.e., median or 50th percentile values) of the age-averaged, lifetime excess cancer incidence (fatal and nonfatal cancer) risk per unit of activity inhaled or ingested, expressed as risk/picocurie (pCi) in Table 4A or as risk/becquerel (Bq) in Table 4B. External exposure slope factors are best estimates of the lifetime excess cancer incidence risk for each year of exposure to external radiation from photon-emitting radionuclides distributed uniformly in a thick layer of soil, and are expressed as risk/yr per pCi/gram of soil (Table 4A) or as risk/yr per Bq/gram of soil (Table 4B). For a discussion on the derivation of radionuclide slope factors and guidance on their use, refer to the User's Guide section on radionuclide carcinogenicity.

b A curie (Ci), the customary unit of activity, is equal to 3.7 x  $10^{10}$  nuclear transformations per second. 1 picocurie (pCi) =  $10^{-12}$  Ci.

For each radionuclide listed, slope factors correspond to the risks per unit intake or exposure for that radionuclide only, except when marked with a "+D" to indicate that the risks from radioactive decay chain products are also included. Radionuclides designated with a "†" are members of a decay chain. Refer to Exhibit 1 in the User's Guide section on radionuclide carcinogenicity for guidance on determining slope factors for partial or complete radioactive decay chains.

d Chemical Abstract Service Reference Number (CASRN). For risk calculations involving decay chains, a CASRN should be reported for the parent radionuclide and each chain member.

e Radioactive half-life: S = Second, M = Minute, D = Day, Y = Year. For those radionuclides with decay products (+D), half-lives are listed for the parent radionuclide.

f Lung clearance classification recommended by the International Commission on Radiological Protection (ICRP): Y = Year, W = Week, D = Day, \* = Gas.

g Gastrointestinal (GI) absorption factors are the fractional amounts of each radionuclide absorbed across the GI tract into the bloodstream. Lung clearance classifications and GI absorption factors are provided for reference only. Do not use these factors to adjust (i.e., multiply or divide) inhalation or ingestion slope factors. See the User's Guide for instructions.

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soil	
Actinium (89)	Ac-225†	014265-85-1	1.000E+01	D	Y	1.0E-03	4.6E-10	6.5E-08	2.1E-07	
	Ac-227 <sup>†</sup>	014952-40-0	2.180E+01	Υ	Υ	1.0E-03	7.6E-09	2.2E-06	7.0E-10	
	Ac-227 + D	014952-40-0(+D)	2.180E+01	Υ	Υ	1.0E-03	9.5E-09	2.4E-06	2.3E-05	
	Ac-228	014331-83-0	6.130E+00	Н	Υ	1.0E-03	1.4E-11	7.0E-10	7.8E-05	
Americium (95)	Am-241 <sup>†</sup>	014596-10-2	4.320E+02	Υ	w	1.0E-03	6.5E-09	8.6E-07	1.3E-07	
	Am-242	013981-54-9	1.600E+01	Н	w	1.0E-03	9.7E-12	3.2E-10	1.6E-07	
	Am-242m	013981-54-9(m)	1.520E+02	Υ	w	1.0E-03	6.2E-09	7.6E-07	3.2E-09	
	Am-243 <sup>1</sup>	014993-75-0	7.380E+03	Υ	w	1.0E-03	6.5E-09	8.6E-07	6.5E-07	
	Am-243 + D	014993-75-0(+D)	7.380E+03	Υ	w	1.0E-03	6.5E-09	8.6E-07	6.8E-06	
Antimony (51)	Sb-122	014374-79-9	2.700E+00	D	w	1.0E-01	5.4E-11	9.2E-11	3.8E-05	
	Sb-124	014683-10-4	6.020E+01	D	W	1.0E-01	7.8E-11	5.9E-10	1.8E-04	
	Sb-125	014234-35-6	2.770E+00	Υ	W	1.0E-01	2.3E-11	3.0E-10	3.2E-05	
	Sb-126	015756-32-8	1.240E+01	D	W	1.0E-01	7.6E-11	2.4E-10	2.5E-04	
	Sb-126m	015756-32-8(m)	1.900E+01	М	w	1.0E-01	1.9E-12	7.0E-13	1.4E-04	
	Sb-127	013968-50-8	3.850E+00	D	W	1.0E-01	5.4E-11	1.1E-10	5.7E-05	
	Sb-129	014331-88-5	4.400E+00	Н	w	1.0E-01	1.6E-11	1.4E-11	1.3E-04	
Argon (18)	Ar-41	014163-25-8	1.830E+00	Н	*	1.0E+00		1.6E-14	1.2E-04	
Astatine (85)	At-217 <sup>†</sup>	017239-90-6	3.230E-02	s	D	1.0E+00	1.2E-16	1.5E-15	2.1E-08	

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi	
Barium (56)	Ba-131	014914-75-1	1.180E+01	D	D	1.0E-01	1.3E-11	9.7E-12	3.2E-05
	Ba-133	013981-41-4	1.050E+01	Υ	D	1.0E-01	3.2E-11	9.7E-11	2.3E-05
	Ba-133m	013981-41-4(m)	3.890E+01	Н	D	1.0E-01	1.7E-11	9.5E-12	2.6E-06
	Ba-137m <sup>†</sup>	013981-97-0(m)	2.550E+00	М	D	1.0E-01	6.5E-14	1.6E-14	5.4E-05
	Ba-139	014378-25-7	8.310E+01	М	D	1.0E-01	5.7E-12	4.1E-12	2.1E-06
	Ba-140	014798-08-4	1.280E+01	D	D	1.0E-01	7.3E-11	5.4E-11	1.5E-05
Beryllium (4)	Be-7	013966-02-4	5.340E+01	D	Υ	5.0E-03	8.1E-13	7.3E-12	4.1E-06
Bismuth (83)	Bı-206	015776-19-9	6.240E+00	D	w	5.0E-02	5.9E-11	1.2E-10	3.0E-04
	Bi-207	013982-38-2	3.340E+01	Υ	w	5.0E-02	3.8E-11	4.9E-10	1.3E-04
	Bi-210 <sup>1</sup>	014331-79-4	5.010E+00	D	W	5.0E-02	4.3E-11	2.2E-09	0.0E + 00
	Bi-211 <sup>†</sup>	015229-37-5	2.130E+00	М	W	5.0E-02	3.3E-13	5.2E-12	3.6E-06
	Bi-212 <sup>†</sup>	014913-49-6	6.055E+01	М	W	5.0E-02	8.4E-12	1.8E-10	1.6E-05
	Bi-213†	015776-20-2	4.570E+01	М	W	5.0E-02	6.2E-12	8.1E-12	1.1E-05
	Bı-214 <sup>†</sup>	014733-03-0	1.990E+01	М	W	5.0E-02	3.5E-12	5.7E-11	1.4E-04
Bromine (35)	Br-82	014686-69-2	3.530E+01	Н	D	1.0E+00	3.0E-11	2.4E-11	2.4E-04
Cadmium (20)	Cd-109	014109-32-1	4.640E+02	D	Y	5.0E-02	2.1E-10	1.8E-09	2.0E-08
	Cd-115	014336-68-6	5.350E+01	н	Y	5.0E-02	4.6E-11	7.0E-11	1.7E-05
	Cd-115m	014336-68-6(m)	4.460E+01	D	Y	5.0E-02	1.4E-10	1.1E-09	2.0E-06

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Po	er Unit Intake or Exposure
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
Calcium (20)	Ca-45	013966-05-7	1.630E+02	D	w	3.0E-01	2.7E-11	1.4E-10	1.6E-16
	Ca-47	001439-99-2	4.540E+00	D	W	3.0E-01	5.4E-11	1.2E-10	9.7E-05
Carbon (6)	C-11	014333-33-6	2.050E+01	М	D	1.0E+00	1.3E-12	5.7E-13	8.6E-05
	C-14	014762-75-5	5.730E+03	Υ	*	1.0E+00	2.4E-11	1.7E-13	0.0E + 00
	C-15	015929-23-4	2.450E+00	S	D	1.0E + 00	2.2E-14	5.7E-15	
Cerium (58)	Ce-141	013967-74-3	3.250E+01	Đ	Υ	3.0E-04	2.2E-11	2.3E-10	3.5E-06
	Ce-143	014119-19-8	3.300E+01	Н	Υ	3.0E-04	3.5E-11	5.9E-11	1.8E-05
	Ce-144	014762-78-8	2.840E+02	D	Υ	3.0E-04	1.6E-10	9.2E 09	6.8E-07
Cesium (55)	Cs-131	014914-76-2	9.690E+00	D	D	1.0E + 00	3.8E-12	2.7E-12	7.6E-08
	Cs-134	013967-70-9	2.060E+00	Υ	D	1.0E + 00	1.1E-09	7.6E-10	1.4E-04
	Cs-134m	013967-70-9(m)	2.900E+00	Н	D	1.0E + 00	1.1E-12	1.1E-12	5.4E-07
	Cs-135	015726-30-4	2.300E+06	Y	D	1.0E + 00	1.1E-10	7.3E-11	0.0E + 00
	Cs-136	014234-29-8	1.320E+01	D	D	1.0E + 00	1.8E-10	1.2E-10	1.9E-04
	Cs-137 <sup>†</sup>	010045-97-3	3.020E+01	Υ	D	1.0E+00	7.6E-10	5.1E-10	0.0E + 00
	Cs-137+D	010045-97-3(+D)	3.020E+01	Υ	D	1.0E+00	7.6E-10	5.1E-10	5.4E-05
	Cs-138	015758-29-9	3.220E+01	М	D	1.0E+00	5.1E-12	2.6E-12	2.2E-04
Chlorine (17)	CI-36	013981-43-6	3.010E+05	Υ	D	1.0E+00	4.9E-11	3.8E-11	0.0E + 00
	CI-38	014158-34-0	3.720E + 01	М	D	1.0E+00	6.2E-12	3.5E-12	1.5E-04

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
Element (Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	L	CRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soil		
Chromium (24)	Cr-51	014392-02-0	2.770E+01	D	Y	1.0E-01	1.2E-12	8.1E-12	2.5E-06		
Cobalt (27)	Co-57	013981-50-5	2.710E+02	D	Υ	3.0E-01	1.6E-11	2.2F-10	5.1E-06		
	Co-58	01381-38-9	7.080E+01	D	Υ	3.0E-01	4.3E-11	2.6E-10	8.9E-05		
	Co-58m	01381-38-9(m)	9.150E+00	Н	Υ	3.0E-01	8.6E-13	2.0E-12	1.3E-09		
	Co-60	010198-40-0	5.270E+00	Υ	Υ	3.0E-01	4.1E-10	4.1E-09	2.3E-04		
Copper (29)	Cu-64	013981-25-4	1.270E+01	н	Υ	5.0E-01	4.6E-12	5.4E-12	1.6E-05		
Curium (96)	Cm-242	015510-73-3	1.630E+02	D	W	1.0E-03	3.5E-10	1.1E-07	9.2E-10		
	Cm-243	015757-87-6	2.850E+01	Υ	W	1.0E-03	5.1E-09	7.0E-07	4.3E-06		
	Cm-244	013981-15-2	1.810E+01	Υ	W	1.0E-03	4.3E-09	5.9E-07	8.1E-10		
	Cm-245	015621-76-8	8.500E+03	Υ	W	1.0E-03	6.5E-09	8.6E-07	1.4E-06		
	Cm-246	015757-90-1	4.750E+03	Υ	W	1.0E-03	6.5E-09	8.6E-07	7.3E-10		
	Cm-247	015758-32-4	1.560E+07	Υ	W	1.0E-03	5.9E-09	8.1E-07	2.5E-05		
	Cm-248	015758-33-5	3.390E+05	Y	W	1.0E-03	2.5E-08	3.2E-06	5:9E-10		
Dysprosium (66)	Dy-165	013967-64-1	2.330E+00	Н	W	3.0E-04	4.1E-12	3.0E-12	1.5E-06		
	Dy-166	015840-01-4	8.160E+01	Н	W	3.0E-04	5.1E-11	1.4E-10	7.3E-07		
Erbium (63)	Er-169	015840-13-8	9.400E+00	D	W	3.0E-04	1.2E-11	4.1E-11	2.3E-10		
	Er-171	014391-45-8	7.520E+00	Н	W	3.0E-04	1.2E-11	1.1E-11	2.5E-05		
Europium (63)	Eu-152	014683-23-9	1.360E+01	Υ	W	1.0E-03	5.7E-11	3.0E-09	9.7E-05		

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> (In Units of Becquerels<sup>b</sup>)

**MARCH 1994** 

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pa	er Unit Intake or Exposure
Element Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	Eu-154	015585-10-1	8.800E + 00	Y	w	1.0E-03	8.1E-11	3.8E-09	1.1E-04
	Eu-155	014391-16-3	4.960E+00	Υ	w	1.0E-03	1.2E-11	4.9E-10	1.6E-06
	Eu-156	014280-35-4	1.520E+01	D	W	1.0E-03	6.8E-11	3.0E-10	1.3E-04
Florin (9)	F-18	013981-56-1	1.100E + 02	M	D	1.0E + 00	2.6E-12	1.95-12	8.4E-05
Francium (87)	Fr-2211	015756-41-9	4.800E+00	М	D	1.0E + 00	1.6E-12	2.5E-11	1.7E-06
	Fr-223 <sup>†</sup>	015756-98-6	2.180E + 00	М	D	1.0E+00	4.6E-12	1.15-11	1.1E-06
Gadolinium (64)	Gd-153	014276-65-4	2.420E + 02	D	W	3.0E-04	8.4E-12	1.6E-10	2.0E-06
	Gd-159	014041-42-0	1.860E + 01	н	w	3.0E-04	1.6E-11	1.7E-11	2.4E-06
Gallium (31)	Ga-67	014119-09-6	3.260E + 00	D	w	1.0E-03	5.7E-12	9.7E-12	8.9E-06
	Ga-72	013982-22-4	1.410E + 01	Н	W	1.0E-03	3.5E-11	3.2E-11	2.6E-04
Germanium (32)	Ge-71	014374-81-3	1.180E+01	D	W	1.0E + 00	1.8E-13	3.5E-12	6.2E-10
Gold (79)	Au-196	014914-16-0	6.180E+00	D	Y	1.0E-01	1.1E-11	2.4E-11	3.5E-05
	Au-198	010043-49-0	2.700E + 00	D	Υ	1.0E-01	3.2E-11	5.75-11	3.2E-05
Holmium (67)	Ho-166	013967-65-2	2.680E + 01	Н	W	3.0E-04	4.6E-11	5.9E-11	1.8E-06
Hydrogen (1)	H-3	010028-17-8	1.230E+01	Υ	*	1.0E + 00	1.5E-12	2.1F-12	0.0E+00
Indium (49)	In-113m	014885-78-0(m)	1.660E + 00	н	w	2.0E-02	1.3E-12	7.8E-13	1.9E-05
	In-114	013981-55-0	7.190E + 01	s	w	2.0E-02	1.5E-13	4.1E-14	2.7E-06
	In-114m	013981-55-0(m)	4.950E+01	D	W	2.0E-02	1.5E-10	1.1E-09	4.9E-06

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
							Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi	
	ln-115	014191-71-0	4.600E + 15	Y	w	2.0E-02	8.6E-10	5.9E-09	0.0E+00	
	In-115m	014191-71-0(m)	4.360E+00	Н	W	2.0E-02	3.2E-12	2.5E-12	1.1E-05	
lodine (53)	I-122	018287-75-7	3.620E+00	М	D	1.0E + 00	6.8E-13	2.1E-13	8.1E-05	
	I-123	015715-08-9	1.310E+01	н	D	1.0E + 00	2.7E-11	1.5E-11	6.5E-06	
	I-125	014158-31-7	6.010E+01	D	D	1.0E+00	2.1E-10	1.4E-10	7.8E-08	
	I-126	014158-32-8	1.290E+01	D	D	1.0E + 00	4.1E-10	2.6E-10	3.5E-05	
	I-129	015046-84-1	1.570E + 07	Υ	D	1.0E+00	5.1E-09	3.2E-09	1.1E-07	
	I-130	014914-02-4	1.240E+01	Н	D	1.0E+00	2.5E-10	1.4E-10	1.9E-04	
	I-131	010043-66-0	8.040E+00	D	D	1.0E+00	9.7E-10	6.5E-10	4.1E-05	
	I-132	014683-16-0	2.300E+00	Н	D	1.0E + 00	2.7E-11	1.6E-11	2.1E-04	
	I-133	014834-67-4	2.080E+01	Н	D	1.0E+00	5.7E-10	3.2E-10	5.4E-05	
	I-134	014914-27-3	5.260E+01	М	D	1.0E+00	7.6E-12	4.3E-12	2.4E-04	
	I-135	014834-68-5	6.610E+00	н	D	1.0E + 00	1.1E-10	6.5£-11	1.5E-04	
	lr-190	014981-91-0	1.180E+01	D	Υ	1.0E-02	3.8E-11	1.3E-10	1.1E-04	
Iridium (77)	lr-192	014694-69-0	7.400E+01	D	Y	1.0E-02	4.6E-11	7.3E-10	6.5E-05	
	lr-194	014158-35-1	1.920E+01	н	Υ	1.0E-02	4.3E-11	4.9E-11	7.6E-06	
Iron (26)	Fe-55	014681-59-5	2.700E+00	Υ	W	1.0E-01	7.3E-12	2.3F-11	0.0E + 00	
	Fe-59	014596-12-4	4.460E+01	D	w	1.0E-01	7.6E-11	2.6E-10	1.1E-04	

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

Element (Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	Gi Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
							Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi	
Krypton (36)	Kr-83m	013965-98-5(m)	1.830E+00	н	*	1.0E + 00		1.7E-15	9.2E-10	
	Kr-85	013983-27-2	1.070E + 01	Υ	*	1.0E+00		1.3E-15	1.9E-07	
	Kr-85m	013983-27-2(m)	4.480E + 00	Н	*	1.0E+00		1.3E-14	9.2E-06	
	Kr-87	014809-68-8	7.630E + 01	М	*	1.0E + 00		5.9E-14	7.6E-05	
	Kr-88	014995-61-0	2.840E+00	Н	*	1.0E + 00		1.3E-13	2.0E-04	
	Kr-89	016316-03-3	3.160E + 00	М	*	1.0E + 00		7.0E-14	1.8E-04	
	Kr-90	015741-13-6	3.230E + 01	s	*	1.0E + 00		6.5E-14	1.1E-04	
Lanthanum (57)	La-140	013981-28-7	4.020E + 01	Н	w	1.0E-03	6.2E-11	8.1E-11	2.2E-04	
Lead (82)	Pb-203	014687-25-3	5.200E + 01	Н	D	2.0E-01	8.6E-12	7.0E-12	1.6E-05	
	Pb-209 <sup>†</sup>	014119-30-3	3.250E + 00	Н	D	2.0E-01	2.3E-12	1.9E-12	0.0E+00	
	Pb-210 <sup>†</sup>	014255-04-0	2.230E+01	Υ	D	2.0E-01	1.4E-08	3.5E-08	3.5E-09	
	Pb-210+D	014255-04-0(+D)	2.230E + 01	Υ	D	2.0E-01	1.8E-08	1.1E-07	4.3E-09	
	Pb-211 <sup>†</sup>	015816-77-0	3.610E+01	М	D	2.0E-01	4.9E-12	7.6E-11	4.3E-06	
	Pb-212 <sup>†</sup>	015092-94-1	1.060E+01	н	D	2.0E-01	1.5E-10	1.2E-09	7.6E-06	
	Pb-214 <sup>†</sup>	015067-28-4	2.680E + 01	М	D	2.0E-01	4.6E-12	7.8E-11	1.7E-05	
Lutetium (71)	Lu-177	014265-75-9	6.710E +00	D	Υ	3.0E-04	1.7E-11	5.1E-11	1.8E-06	
Manganese (25)	Mn-52	014092-99-0	5.590E + 00	D	w	1.0E-01	5.9E-11	1.0E-10	3.2E-04	
	Mn-54	013966-31-9	3.130E+02	D	w	1.0E-01	3.0E-11	1.4E-10	7.8E-05	

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

Element (Atomic <b>N</b> umber)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>			GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
					ICRP Lung Class <sup>f</sup>		Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soil	
	Mn-56	014681-52-8	2.580E+00	Н	W	1.0E-01	1.1E-11	7.6E-12	1.6E-04	
Mercury (80)	Hg-197	013981-51-6	6.410E+01	н	w	2.0E-02	7.3E-12	1.2E-11	1.5E-06	
	Hg-203	013982-78-0	4.660E+01	D	W	2.0E-02	1.8E-11	1.3E-10	1.5E-05	
Molybdenum (42)	Mo-99	014119-15-4	6.600E+01	н	Υ	8.0E-01	4.1E-11	7.0E-11	1.3E-05	
Neodymium (60)	Nd-147	014269-74-0	1.100E+01	D	Υ	3.0E-04	3.5E-11	1.5E-10	8.1E-06	
	Nd-149	015749-81-2	1.730E+00	н	Υ	3.0E-04	5.1E-12	4.3E-12	2.6E-05	
Neptunium (93)	Np-236	015700-36-4	1.150E+05	Υ	w	1.0E-03	6.2E-12	1.1E-10	2.4E-06	
	Np-237 <sup>†</sup>	013994-20-2	2.140E+06	Υ	W	1.0E-03	5.9E-09	7.8E-07	2.1E-07	
	Np-237+D	013994-20-2(+D)	2.140E+06	Υ	W	1.0E-03	5.9E-09	7.8E-07	1.2E-05	
	Np-238	015766-25-3	2.120E+00	D	W	1.0E-03	3.0E-11	8.9E-11	4.6E-05	
	Np-239 <sup>†</sup>	013968-59-7	2.360E+00	D	, W	1.0E-03	2.5E-11	4.1E-11	6.2E-06	
	Np-240	015690-84-3	6.500E+01	М	W	1.0E-03	3.5E-12	1.8E-12	8.6E-05	
	Np-240m	015690-84-3(m)	7.400E+00	М	W	1.0E-03	7.8E-13	2.4E-13	2.5E-05	
Nickel (28)	Ni-59	014336-70-0	7.500E+04	Υ	W	5.0E-02	2.5E-12	1.9E-11	0.0E + 00	
	Ni-63	013981-37-8	1.000E+02	Υ	W	5.0E-02	6.5E-12	4.9E-11	0.0E + 00	
	Nı-65	014833-49-9	2.520E+00	н	W	5.0E-02	7.0E-12	5.1E-12	5.1E-05	
Niobium (41)	Nb-93m	007440-03-1(m)	1.460E+01	Υ	Υ	1.0E-02	4.1E-12	5.1E-10	1.4E-09	
	Nb-94	014681-63-1	2.030E+04	Υ	Y	1.0E-02	5.7E-11	5.7E-09	1.5E-04	

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> (In Units of Becquerels<sup>b</sup>)

**MARCH 1994** 

Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>®</sup>			GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure			
					ICRP Lung Class <sup>f</sup>		Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi	
	Nb-95	013967-76-5	3.510E + 01	D	Υ	1.0E-02	1.8E-11	1.4E-10	7.0E-05	
	Nb-95m	013967-76-5(m)	8.660E+01	Н	Υ	1.0E-02	1.8E-11	4.3E-11	2.2E-06	
	Nb-97	018496-04-3	7.210E + 01	М	Υ	1.0E-02	3.2E-12	1.9E-12	5.9E-05	
	Nb-97m	018496-04-3(m)	6.000E + 01	s	Υ	1.0E-02	6.5E-14	3.0E-14	6.8E-05	
Osmium (76)	Os-185	015766-50-4	9.360E+01	D	Υ	1.0E-02	1.7E-11	2.4E-10	5.9E-05	
	Os-191	014119-24-5	1.540E+01	D	Υ	1.0E-02	1.8E-11	9.7E-11	2.3E-06	
	Os-191m	014119-24-5(m)	1.300E + 01	Н	Υ	1.0E-02	3.2E-12	5.7E-12	9.2E-08	
	Os-193	016057-77-5	3.000E + 01	Н	Y	1.0E-02	2.6E-11	3.2E-11	4.6E-06	
Palladium (46)	Pd-100	015690-69-4	3.640E + 00	D	Y	5.0E-03	2.7E-11	6.2E-11		
	Pd-101	015749-54-9	8.480E + 00	Н	Υ	5.0E-03	3.2E-12	3.0E-12		
	Pd-103	014967-68-1	1.700E + 01	D	Y	5.0E-03	5.9E-12	3.8E-11	2.0E-08	
	Pd-107	017637-99-9	6.500E+06	Υ	Υ	5.0E-03	1.2E-12	1.7E-10	0.0E+00	
	Pd-109	014981-64-7	1.350E + 01	Н	Υ	5.0E-03	2.1E-11	2.2E-11	5.9E-08	
Phosphorus (15)	P-32	014596-37-3	1.430E+01	D	D	8.0E-01	9.5E-11	8.1E-11	0.0E+00	
	P-33	015749-66-3	2.540E+01	D	D	8.0E-01	1.5E-11	1.2E-11	0.0E+00	
Platinum (78)	Pt-191	015706-36-2	2.710E + 00	D	D	1.0E-02	1.0E-11	8.1E-12	1.7E-05	
	Pt-193	015735-70-3	5.000E + 01	Υ	D	1.0E-02	9.5E-13	2.2E-12	0.0E+00	
	Pt-193m	015735-70-3(m)	4.330E+00	D	D	1.0E-02	1.4E-11	1.1E-11	2.1E-07	

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

						Lifetime Exc	Slope Factor ess Total Cancer Risk Po	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive	ICRP Lung Class	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	Pt-197	015735-74-7	1.830E+01 F	D	1.0E-02	1.3E-11	8.6E-12	8.4E-07
	Pt-197m	015735-74-7(m)	9.440E+01 N	1 D	1.0E-02	3.2E-12	2.4E-12	4.3E-06
Plutonium (94)	Pu-236	015411-92-4	2.850E+00 Y	Y	1.0E-03	1.4E-09	6.5E-07	9.2E-10
	Pu-238	013981-16-3	8.780E+01 Y	Y	1.0E-03	5.9E-09	1.1E-06	7.6E-10
	Pu-239	015117-48-3	2.410E+04 Y	Y	1.0E-03	6.2E-09	1.0E-06	4.6E-10
	Pu-240	014119-33-6	6.570E+03 Y	Y	1.0E-03	6.2E-09	1.0E-06	7.3E-10
	Pu-241	014119-32-5	1.440E+01 Y	Y	1.0E-03	9.7E-11	6.2E-09	0.0E + 00
	Pu-242	013982-10-0	3.760E+05 Y	Y	1.0E-03	5.9E-09	9.7E-07	6.2E-10
	Pu-243	015706-37-3	4.960E+00 H	ı Y	1.0E-03	3.0E-12	2.7E-12	4.9E-07
	Pu-244	014119-34-7	8.260E+07 \	Y	1.0E-03	5.9E-09	9.7E-07	5.1E-10
Polonium (84)	Po-210 <sup>†</sup>	013981-52-7	1.380E+02 [	w w	1.0E-01	4.1E-09	7.0E-08	7.8E-10
	Po-212 <sup>†</sup>	015389-34-1	2.980E-07 S	s w	1.0E-01	5.9E-22	1.6E-20	0.0E + 00
	Po-213 <sup>†</sup>	015756-57-7	4.200E-06	s w	1.0E-01	8.6E-21	2.2E-19	2.7E-09
	Po-214 <sup>†</sup>	015735-67-8	1.640E-04 S	s w	1.0E-01	2.7E-19	7.6E-18	7.6E-09
	Po-215 <sup>†</sup>	015706-52-2	1.780E-03	s w	1.0E-01	7.6E-18	1.5E-16	1.2E-08
	Po-216 <sup>†</sup>	015756-58-8	1.460E-01	s w	1.0E-01	8.1E-16	1.3E-14	1.4E-09
	Po-218 <sup>†</sup>	015422-24-9	3.050E+00 M	и w	1.0E-01	7.6E-13	1.6E-11	0.0E+00
Potassium (19)	K-40	013966-00-2	1.280E+09	/ D	1.0E + 00	3.0E-10	2.1E-10	1.5E-05

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	K-42	014378-21-3	1.240E+01	н	D	1.0E + 00	2.4E-11	3.2E-11	2.6E-05
Praseodymium (59)	Pr-142	014191-64-1	1.910E+01	н	Y	3.0E-04	4.3E-11	4.9E-11	5.7E-06
	Pr-143	014981-79-4	1.360E+01	D	Y	3.0E-04	3.8E-11	1.9E-10	8.1E-13
	Pr-144	014119-05-2	1.730E+01	М	Υ	3.0E-04	2.6E-12	9.7E-13	3.2E-06
	Pr-144m	014119-05-2(m)	7.200E + 00	М	Υ	3.0E-04	1.0E-12	4.3E-13	5.7E-08
Promethium (61)	Pm-147	014380-75-7	2.620E + 00	Υ	Y	3.0E-04	8.4E-12	8.1E-10	1.6E-10
	Pm-148	014683-19-3	5.370E + 00	D	Υ	3.0E-04	8.4E-11	2.1E-10	5.1E-05
	Pm-148m	014683-19-3(m)	4.130E+01	D	Y	3.0E-04	6.8E-11	1.3E-09	1.8E-04
	Pm-149	015765-31-8	5.310E+01	Н	Υ	3.0E-04	3.2E-11	5.1E 11	8.9E-07
Protactinium (91)	Pa-231 <sup>†</sup>	014331-85-2	3.730E + 04	Υ	Υ	1.0E-03	2.5E-09	9.7E-07	7.0E-07
	Pa-2331	013981-14-1	2.700E + 01	D	Y	1.0E-03	2.7E-11	2.3E-10	1.1E-05
	Pa-234	015100-28-4	6.700E + 00	Н	Υ	1.0E-03	1.8E-11	1.5E-11	1.6E-04
	Pa-234m <sup>†</sup>	015100-28-4(m)	1.170E+00	М	Υ	1.0E-03	1.6E-13	4.3E-14	9.7E-07
Radium (88)	Ra-223 <sup>†</sup>	015623-45-7	1.140E+01	D	W	2.0E-01	1.7E-09	8.4E-08	6.2E-06
	Ra-224 <sup>†</sup>	013233-32-4	3.620E+00	D	W	2.0E-01	1.0E-09	3.2E-08	6.2E-07
	Ra-225 <sup>†</sup>	013981-53-8	1.480E+01	D	W	2.0E-01	1.4E-09	4.1E-08	5.1E-08
	Ra-226 <sup>†</sup>	013982-63-3	1.600E + 03	Υ	W	2.0E-01	3.2E-09	8.1E-08	3.2E-07
	Ra-226 + D	013982-63-3(+D)	1.600E+03	Υ	W	2.0E-01	3.2E-09	8.1E-08	1.6E-04

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ss Total Cancer Risk Pe	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f₁) <sup>g</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	Ra-228†	015262-20-1	5.750E+00	Υ	w	2.0E-01	2.7E-09	1.8E-08	0.0E + 00
	Ra-228+D	015262-20-1(+D)	5.750E+00	Υ	w	2.0E-01	2.7E-09	1.9E-08	7.8E-05
Radon (86)	Rn-219 <sup>†</sup>	014835-02-0	3.960E+00	s	*	1.0E + 00		1.2F-12	4.3E-06
	Rn-220 <sup>†</sup>	022481-48-7	5.560E+01	s	*	1.0E+00		3.2E-12	4.6E-08
	Rn-222 <sup>†</sup>	014859-67-7	3.820E+00	D	*	1.0E+00	3.8E-11	2.0F-11	3.2E-08
	Rn-222+D	014859-67-7(+D)	3.820E+00	D	*	1.0E + 00	4.6E-11	2.1E-10	1.6E-04
Rhodium (45)	Rh-103m	007440-16-6(m)	5.610E+01	М	Y	5.0E-02	1.9E-13	1.1F-13	2.1E-09
	Rh-105	014913-89-4	3.540E+01	н	Y	5.0E-02	1.2E-11	1.6E-11	5.9E-06
	Rh-105m	014913-89-4(m)	4.500E+01	s	Y	5.0E-02	1.7E-14	9.2F-15	5.9E-07
	Rh-106	014234-34-5	2.990E+01	s	Y	5.0E-02	1.2E-13	3.2E-14	1.8E-05
Rubidium (37)	Rb-82	014391-63-0	1.250E+00	М	D	1.0E+00	3.2E-13	9.5E-14	9.5E-05
	Rb-86	014932-53-7	1.870E+01	D	D	1.0E + 00	1.6E-10	1.2E-10	8.9E-06
	Rb-87	013982-13-3	4.730E + 10	Υ	D	1.0E + 00	9.2E-11	6.5E-11	0.0E + 00
	Rb-88	014928-36-0	1.780E+01	М	D	1.0E+00	4.6E-12	2.1F-12	6.5E-05
	Rb-89	014191-65-2	1.540E+01	М	D	1.0E+00	2.5E-12	1.1E-12	2.0E-04
Ruthenium (44)	Ru-97	015758-35-7	2.900E+00	D	Υ	5.0E-02	4.6E-12	7.0E-12	1.1E-05
	Ru-103	013968-53-1	3.940E+01	D	Υ	5.0E-02	2.4E-11	2.3E-10	4.1E-05
	Ru-105	014331-95-4	4.440E+00	Н	Υ	5.0E-02	1.0E-11	8.9E 12	7.0E-05

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Po	er Unit Intake or Exposure
Element Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	Ru-106	013967-48-1	3.680E + 02	D	Y	5.0E-02	2.6E-10	1.2E 08	0.0E + 00
Samarıum (62)	Sm-147	014392-33-7	1.060E + 11	Υ	w	3.0E-04	4.3E-10	1.9E-07	0.0E + 00
	Sm-151	015715-94-3	9.000E + 01	Y	w	3.0E-04	3.0E-12	2.4E-10	1.1E-11
	Sm-153	015766-00-4	4.670E+01	н	w	3.0E-04	2.4E-11	3.5E-11	1.2E-06
Scandium (21)	Sc-46	013967-63-0	8.380E+01	D	Y	1.0E-04	4.3E-11	7.3E-10	1.9E-04
	Sc-47	014391-96-9	3.420E+00	D	Y	1.0E-04	1.7E-11	3.2E-11	6.2E-06
	Sc-48	014391-86-7	4.370E+01	Н	Y	1.0E-04	5.1E-11	6.2E-11	3.2E-04
Selenium (34)	Se-75	014265-71-5	1.200E + 02	D	W	8.0E-01	1.6E-10	1.6E-10	2.2E-05
Silicon (14)	Sı-31	014276-49-4	1.570E + 02	М	w	1.0E-02	5.9E-12	4.6E-12	8.1E-08
Silver (47)	Ag-105	014928-14-4	4.130E+01	D	Y	5.0E-02	2.0E-11	1.1E-10	
	Ag-108	014391-65-2	2.370E + 00	М	Y	5.0E-02	2.3E-13	6.5E-14	1.4E-06
	Ag-108m	014391-65-2(m)	1.270E + 02	Υ	Y	5.0E-02	9.5E-11	4.1E-09	1.4E-04
	Ag-109m	014378-38-2(m)	3.960E + 01	s	Y	5.0E-02	8.9E-15	2.4E-15	3.5E-08
	Ag-110	014391-76-5	2.460E + 01	s	Y	5.0E-02	8.1E-14	2.2E-14	2.7E-06
	Ag-110m	014391-76-5(m)	2.500E + 02	D	Y	5.0E-02	1.3E-10	1.9E-09	2.5E-04
	Ag-111	157690-04-0	7.460E + 00	D	Y	5.0E-02	4.3E-11	1.3E 10	2.1E-06
Sodium (11)	Na-22	013966-32-0	2.600E + 00	Υ	D	1.0E+00	1.8E-10	1.3E-10	1.9E-04
	Na-24	013982-04-2	1.500E+01	Н	D	1.0E+00	2.7E-11	2.6E-11	4.3E-04

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element (Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi	
Strontium (38)	Sr-82	014809-50-8	2.500E+01	D	D	3.0E-01	2.0E-10	1.9E-10	3.5E-09
	Sr-85	013967-73-2	6.480E+01	D	D	3.0E-01	2.1E-11	2.7E-11	3.8E-05
	Sr-85m	013967-73-2(m)	6.770E+01	М	D	3.0E-01	3.2E-13	1.6E-13	1.3E-05
	Sr-89	014158-27-1	5.060E+01	D	D	3.0E-01	8.1E-11	7.8E-11	1.3E-08
	Sr-90†	010098-97-2	2.860E+01	Υ	D	3.0E-01	8.9E-10	1.5E-09	0.0E + 00
	Sr-90+D	010098-97-2(+D)	2.860E+01	Υ	D	3.0E-01	9.7E-10	1.7E-09	0.0E + 00
	Sr-91	014331-91-0	9.500E+00	н	D	3.0E-01	2.3E-11	1.9E-11	6.5E-05
	Sr-92	014928-29-1	2.710E+00	Н	D	3.0E-01	1.5E-11	1.2E-11	1.2E-04
Sulfur (16)	S-35	015117-53-0	8.740E+01	D	D	8.0E-01	5.9E-12	5.1E-12	0.0E + 00
Tantalum (73)	Ta-182	013982-00-8	1.150E+02	D	Υ	1.0E-03	4.6E-11	1.2E-09	1.1E-04
Technetium (43)	Tc-95	014809-56-4	2.000E+01	Н	W	8.0E-01	1.5E-12	6.2E-13	6.5E-05
	Tc-95m	014809-56-4(m)	6.100E+01	D	W	8.0E-01	4.9E-11	1.1E-10	5.1E-05
	Tc-96	014808-44-7	4.280E+00	D	W	8.0E-01	4.9E-11	5.7E-11	2.2E-04
	Tc-96m	014808-44-7(m)	5.150E+01	М	W	8.0E-01	6.2E-13	5.7E-13	1.9E-06
	Tc-97	015759-35-0	2.600E+06	Υ	W	8.0E-01	4.1E-12	2.6E-11	9.5E-09
	Tc-97m	015759-35-0(m)	8.900E+01	D	W	8.0E-01	3.0E-11	1.4E-10	1.0E-08
	Tc-99	014133-76-7	2.130E+05	Υ	W	8.0E-01	3.5E-11	2.2E-10	1.6E-11
	Tc-99m	014133-76-7(m)	6.020E+00	Н	W	8.0E-01	1.4E-12	7.3E-13	6.2E-06

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive L		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soil
Tellurium (52)	Te-125m	014390-73-9(m)	5.800E+01	D	W	2.0E-01	2.3E-11	1.5E-10	7.0E-08
	Te-127	013981-49-2	9.350E+00	Н	w	2.0E-01	6.2E-12	6.2E-12	3.8E-07
	Te-127m	013981-49-2(m)	1.090E + 02	D	w	2.0E-01	5.9E-11	4.3E-10	2.3E-08
	Te-129	014269-71-7	6.960E+01	М	w	2.0E-01	3.0E-12	1.8E-12	3.5E-06
	Te-129m	014269-71-7(m)	3.360E+01	D	W	2.0E-01	8.6E-11	5.4E-10	1.7E-06
	Te-131	014683-12-6	2.500E+01	М	W	2.0E-01	7.6E-12	3.8E-12	3.2E-05
	Te-131m	014683-12-6(m)	3.000E+01	Н	W	2.0E-01	1.0E-10	1.5E-10	1.3E-04
	Te-132	014234-28-7	7.820E+01	н	W	2.0E-01	8.1E-11	1.4E-10	1.1E-05
Terbium (65)	Tb-158	015759-55-4	1.500E+02	Υ	W	3.0E-04	3.2E-11	2.5E-09	
	Tb-160	013981-29-8	7.230E+01	D	W	3.0E-04	4.9E-11	5.1E-10	9.7E-05
Thallium (81)	TI-202	015720-57-7	1.220E+01	D	D	1.0E+00	2.3E-11	1.6E-11	3.5E-05
	TI-204	013968-51-9	3.780E+00	Υ	D	1.0E+00	4.6E-11	3.5E-11	2.4E-08
	TI-207 <sup>†</sup>	014133-67-6	4.770E+00	М	D	1.0E+00	3.5E-13	1.2E-13	2.0E-07
	TI-208 <sup>†</sup>	014913-50-9	3.050E+00	М	D	1.0E+00	4.9E-13	1.4E-13	3.5E-04
	TI-209†	015690-73-0	2.200E+00	М	D	1.0E+00	3.8E-13	1.2E-13	1.9E-04
Thorium (90)	Th-227 <sup>†</sup>	015623-47-9	1.870E+01	D	Υ	2.0E-04	1.2E-10	1.3E-07	4.3E-06
	Th-228†	014274-82-9	1.910E+00	Υ	Υ	2.0E-04	3.0E-10	2.1E-06	1.5E-08
	Th-228 + D	014274-82-9(+D)	1.910E+00	Υ	Υ	2.0E-04	1.5E-09	2.1E-06	1.5E-04

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	Th-229 <sup>†</sup>	015594-54-4	7.340E+03	Υ	Y	2.0E-04	5.7E-10	2.0E-06	
	Th-229 + D	015594-54-4(+D)	7.340E+03	Υ	Y	2.0E-04	2.4E-09	2.1E-06	1.8E-05
	Th-230†	014269-63-7	7.700E + 04	Υ	Υ	2.0E-04	3.5E-10	7.8 <del>E</del> -07	1.5E-09
	Th-231 <sup>†</sup>	014932-40-2	2.550E+01	Н	Υ	2.0E-04	1.1E-11	1.3E-11	6.2E-08
	Th-232 <sup>†</sup>	007440-29-1	1.410E+10	Υ	Υ	2.0E-04	3.2E-10	7.6E-07	7.0E-10
	Th-2341	015065-10-8	2.410E+01	D	Υ	2.0E-04	1.1E-10	8.6E-10	9.5E-08
Thulium (69)	Tm-170	013981-30-1	1.290E+02	D	W	3.0E-04	4.1E-11	5.7E-10	1.0E-07
	Tm-171	014333-45-0	1.920E+00	Υ	w	3.0E-04	3.2E-12	8.4E-11	8.9E-09
Tin (50)	Sn-113	013966-06-8	1.150E+02	D	W	2.0E-02	2.4E-11	2.5E-10	8.9E-08
	Sn-121	014683-06-8	2.710E+01	Н	W	2.0E-02	7.3E-12	8.6F-12	
	Sn-121m	014683-06-8(m)	5.550E+01	Υ	W	2.0E-02	1.5E-11	2.5E-10	
	Sn-125	014683-08-0	9.640E+00	D	W	2.0E-02	9.7E-11	3.2E-10	3.0E-05
	Sn-126	015832-50-5	1.000E + 05	Υ	W	2.0E-02	1.5E-10	2.1E-09	8.9E-07
Tungsten (74)	W-181	015749-46-9	1.210E+02	D	D	3.0E-01	2.2E-12	1.7E-12	5.9E-07
	W-185	014932-41-3	7.510E+01	D	D	3.0E-01	1.3E-11	8.1E-12	1.3E-09
	W-187	014983-48-3	2.380E+01	Н	D	3.0E-01	1.7E-11	1.1E-11	4.1E-05
Uranium (92)	U-232	014158-29-3	7.200E+01	Υ	Υ	5.0E-02	1.0E-09	1.6E-06	1.2E-09
	U-233	013968-55-3	1.590E + 05	Υ	Y	5.0E-02	4.3E-10	7.3E-07	1.1E-09

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soi
	U-234†	013966-29-5	2.450E+05	Υ	Υ	5.0E-02	4.3E-10	7.0E-07	8.1E-10
	U-235 <sup>†</sup>	015117-96-1	7.040E + 08	Υ	Y	5.0E-02	4.3E-10	6.8E-07	6.5E-06
	U-235 + D	015117-96-1(+D)	7.040E + 08	Υ	Υ	5.0E-02	4.3E-10	6.8E-07	6.5E-06
	U-236	013982-70-2	2.340E+07	Υ	Υ	5.0E-02	4.1E-10	6.8E-07	6.5E-10
	U-237	014269-75-1	6.750E + 00	D	Y	5.0E-02	2.4E-11	7.0E-11	3.5E-06
	U-238 <sup>†</sup>	007440-61-1	4.470E+09	Υ	Y	5.0E-02	4.3E-10	6.5E-07	5.7E-10
	U-238 + D	007440-61-1(+D)	4.470E + 09	Υ	Y	5.0E-02	5.4E-10	6.5E-07	1.4E-06
	U-240	015687-53-3	1.410E+01	н	Y	5.0E-02	3.2E-11	3.2F-11	4.1E-09
Vanadium (23)	V-48	014331-97-6	1.600E + 01	D	W	1.0E-02	5.9E-11	2.1E-10	2.7E-04
Xenon (54)	Xe-122	015151-09-4	2.010E+01	Н	*	1.0E+00		8.9E-14	2.3E-06
	Xe-123	015700-10-4	2.140E+00	Н	*	1.0E+00		2.0E-14	4.6E-05
	Xe-125	013994-18-8	1.680E+01	Н	*	1.0E+00		1.1E-14	1.3E-05
	Xe-127	013994-19-9	3.640E+01	D	*	1.0E+00		1.1E-14	1.4E-05
	Xe-129m	013965-99-6(m)	8.890E+00	D	*	1.0E+00		1.6E-14	3.5E-07
	Xe-131m	014683-11-5(m)	1.180E+01	D	*	1.0E+00		1.2E-14	1.2E-07
	Xe-133	014932-42-4	5.250E+00	D	*	1.0E+00		1.2E-14	6.2E-07
	Xe-133m	014932-42-4(m)	2.190E+00	D	*	1.0E + 00		1.5E-14	8.9E-07
	Xe-135	014995-62-1	9.110E+00	Н	*	1.0E + 00		2.2E-14	1.7E-05

Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> MARCH 1994 (In Units of Becquerels<sup>b</sup>)

							Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	er Unit Intake or Exposure
Element (Atomic Number)	Isotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>		ICRP Lung Class <sup>f</sup>	Gl Absorption Factor (f <sub>1</sub> ) <sup>9</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soil
	Xe-135m	014995-62-1(m)	1.540E+01	М	*	1.0E + 00		5.7E-15	3.5E-05
	Xe-137	014835-21-3	3.830E+00	М	*	1.0E + 00		4.9E-14	1.6E-05
	Xe-138	015751-81-2	1.410E+01	М	*	1.0E+00		7.6E-14	1.1E-04
Yttrium (39)	Y-90 <sup>†</sup>	010098-91-6	6.410E+01	Н	Υ	1.0E-04	8.6E-11	1.5E-10	0.0E + 00
	Y-91	014234-24-3	5.850E+01	D	Υ	1.0E-04	7.6E-11	1.3E-09	3.2E-07
	Y-91m	014234-24-3(m)	4.970E+01	М	Υ	1.0E-04	5.9E-13	9.5E-13	4.6E-05
	Y-92	015751-59-4	3.540E+00	Н	Y	1.0E-04	1.9E-11	1.6E-11	2.3E-05
	Y-93	014981-70-5	1.010E+01	Н	Y	1.0E-04	3.8E-11	3.8E-11	8.1E-06
Zinc (30)	Zn-65	013982-39-3	2.440E+02	D	Y	5.0E-01	2.3E-10	4.3E-10	5.4E-05
	Zn-69	013982-23-5	5.560E+01	M	Y	5.0E-01	1.5E-12	8.6E-13	4 9E-10
	Zn-69m	013982-23-5(m)	1.380E+01	Н	Υ	5.0E-01	1.2E-11	1.5E-11	3.5E-05
Zırconium (40)	Zr-93	015751-77-6	1.530E+06	Υ	W	2.0E-03	4.6E-12	1.8E-10	0.0E + 00
	Zr-95	013967-71-0	6.400E+01	D	W	2.0E-03	2.7E-11	2.7E-10	6.8E-05
	Zr-97	014928-30-4	1.690E+01	Н	W	2.0E-03	6.5E-11	6.8E-11	1.6E-05

# Table 4B: Radionuclide Carcinogenicity -- Slope Factors<sup>a</sup> (In Units of Becquerels<sup>b</sup>)

						Lifetime Exce	Slope Factor ess Total Cancer Risk Pe	r Unit Intake or Exposure
Element (Atomic Number)	lsotope <sup>c</sup>	CASRN <sup>d</sup>	Radioactive Half-life <sup>e</sup>	ICRP Lung Class <sup>f</sup>	GI Absorption Factor (f <sub>1</sub> ) <sup>g</sup>	Ingestion (Risk/Bq)	Inhalation (Risk/Bq)	External Exposure (Risk/yr per Bq/g soil)

#### **ENDNOTES:**

**MARCH 1994** 

<sup>&</sup>lt;sup>a</sup> EPA classifies all radionuclides as Group A (known human) carcinogens. Radionuclide slope factors are calculated by EPA's Office of Radiation and Indoor Air (ORIA) to assist HEAST users with risk-related evaluations and decision-making at various stages of the remediation process. Ingestion and inhalation slope factors are best estimates (i.e., median or 50th percentile values) of the age-averaged, lifetime excess cancer incidence (fatal and nonfatal cancer) risk per unit of activity inhaled or ingested, expressed as risk/picocurie (pCi) in Table 4A or as risk/becquerel (Bq) in Table 4B. External exposure slope factors are best estimates of the lifetime excess cancer incidence risk for each year of exposure to external radiation from photon-emitting radionuclides distributed uniformly in a thick layer of soil, and are expressed as risk/yr per pCi/gram of soil (Table 4A) or as risk/yr per Bq/gram of soil (Table 4B). For a discussion on the derivation of radionuclide slope factors and guidance on their use, refer to the User's Guide section on radionuclide carcinogenicity.

b A becquerel (Bq), the International System (SI) unit of activity, is equal to one nuclear transformations per second. 1 Bq ≈ 27 picocuries (pCi).

<sup>&</sup>lt;sup>c</sup> For each radionuclide listed, slope factors correspond to the risks per unit intake or exposure for that radionuclide only, except when marked with a "+D" to indicate that the risks from radioactive decay chain products are also included. Radionuclides designated with a "†" are members of a decay chain. Refer to Exhibit 1 in the User's Guide section on radionuclide carcinogenicity for guidance on determining slope factors for partial or complete radioactive decay chains.

d Chemical Abstract Service Reference Number (CASRN). For risk calculations involving decay chains, a CASRN should be reported for the parent radionuclide and each chain member.

e Radioactive half-life: S = Second, M = Minute, D = Day, Y = Year. For those radionuclides with decay products (+D), half-lives are listed for the parent radionuclide.

f Lung clearance classification recommended by the International Commission on Radiological Protection (ICRP): Y = Year, W = Week, D = Day, \* = Gas.

g Gastrointestinal (GI) absorption factors are the fractional amounts of each radionuclide absorbed across the GI tract into the bloodstream. Lung clearance classifications and GI absorption factors are provided for reference only. Do not use these factors to adjust (i.e., multiply or divide) inhalation or ingestion slope factors. See the User's Guide for instructions.

#### APPENDIX A: TECHNICAL INFORMATION

- I. DATA SOURCES AND SELECTION CRITERIA USED IN HEAST
- II. DOSE CONVERSIONS ON HEAST
- III. CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBERS CROSS REFERENCE
- IV. EFFECT LEVEL DEFINITIONS
- V. NATIONAL AMBIENT AIR QUALITY STANDARDS (NAAQS)



#### **APPENDIX A-I**

#### I. DATA SOURCES AND SELECTION CRITERIA USED IN HEAST

- A. Description of Sources and Documents Cited in HEAST
  - 1. The Integrated Risk Information System (IRIS)

IRIS is an on-line data base developed by the EPA for compilation of risk assessment and regulatory information on chemicals and physical agents. IRIS is the primary communications mechanism for distribution of health hazard assessment information derived by the various intra-Agency Work Groups. The primary intent of IRIS is to provide guidance to EPA personnel in making risk management decisions. An IRIS chemical file contains a Work Group verified summary of the available information on hazard and dose-response assessment for noncarcinogenic and/or carcinogenic effects for that chemical and is not an extensive toxicologic data base. Risk assessment values placed on IRIS are considered Agency consensus and take precedence over differing risk assessment values from other EPA sources. Each file includes referenced citations and EPA contacts for obtaining further information on any specific chemical or agent. The IRIS data base was made available to State and local governments, as well as to the public, in April 1988.

\* Questions concerning IRIS: Call RISK INFORMATION HOTLINE at (513) 569-7254

#### 2. EPA Work Groups:

Risk assessment values for chemicals currently being considered by EPA Work Groups, but not yet on IRIS, are included in HEAST. The EPA Reference Dose/Reference Concentration (RfD/RfC) Work Group and the Carcinogen Risk Assessment Verification Endeavor (CRAVE) Work Group validates Agency systemic toxicity and carcinogen risk assessments, respectively. These Work Groups are also responsible for resolving any conflicts regarding toxicity values developed by various Program Offices. Work Group members represent many different EPA offices and are scientists experienced in issues related to both the qualitative and quantitative risk assessment of carcinogenic and toxic agents. Values verified by these Work Groups have undergone extensive peer review and represent an Agency consensus. Verified risk assessment values are entered into the IRIS data base monthly.

\* Questions concerning RfD/RfC Work Group: Call Mike Dourson (ECAO-Cincinnati) at (513) 569-7533 or Annie Jarabek (ECAO-Research Triangle Park) at (919) 541-4847

- \* Questions concerning CRAVE Work Group: Call Jim Cogliano at (202) 260-3814 (OHEA-Washington, DC)
  - 3. Office of Research and Development (ORD/Office of Health and Environmental Assessment (OHEA) OSWER-OAQPS (Office of Solid Waste and Emergency Response-Office of Air Quality Planning and Standards) Documents:

A listing of most ORD/OHEA OSWER-OAQPS documents can be found in the Chemical Assessments and Related Activities (CARA) list (available through NTIS) or in the CERI (Center for Environmental Research Information) Office of Research and Development publications list. The CARA is produced by the Office of Health and Environmental Assessment (OHEA). All OSWER-OAQPS documents are subject to a minimum of internal EPA peer review or a maximum of EPA/Peer Review Workshop/Science Advisory Board and public comments prior to finalization.

\* Information on the availability of OSWER-OAQPS documents can be obtained from the following sources:

#### All Documents:

Technical Information Staff
Office of Health and Environmental Assessment (RD-689)
U.S. Environmental Protection Agency
401 M Street, S.W.
Washington, D.C. 20460
(202) 260-7345

#### **Published Documents:**

Center for Environmental Research Information (CERI)
Office of Research and Development
U.S. Environmental Protection Agency
26 W. Martin Luther King Drive
Cincinnati, OH 45268
(513) 569-7562

#### **Documents Available Through RCRA/Superfund:**

Hotline Number 1-800-424-9346

#### **Documents Available from NTIS:**

National Technical Information Service (NTIS) 5285 Port Royal Road Springfield, VA 22161 (703) 487-4650

Health Effects Assessments (HEAs): This document series was prepared by the Environmental Criteria and Assessment Office (ECAO-Cincinnati) for the Office of Emergency and Remedial Response (Superfund). HEAs are intended for use by the OERR in evaluating risk for its preliminary assessment process at uncontrolled sites, and for appraising clean-up alternatives in its remedial investigation/feasibility studies. HEAs are brief, quantitatively oriented, preliminary assessment of relevant health effects data. HEAs are subject to internal EPA review by staff within the Office of Health and Environmental Assessment. Final drafts of HEAs become part of the RCRA and Superfund dockets and are available through NTIS. This series has recently been incorporated into the following HEED series.

Health and Environmental Effects Documents (HEEDs): This document series is prepared by the Environmental Criteria and Assessment Office (ECAO-Cincinnati) for the Office of Solid Waste and Emergency Response (OSWER). HEEDs are intended to support listings under the Resource Conservation and Recovery Act (RCRA) as well as to provide health-related limits and goals for emergency and remedial actions under the Comprehensive Environmental Response, Compensation and Liability Act (CERCLA). Within a HEED, both published literature and information within Agency Program Offices are evaluated as they pertain to potential human health, aquatic life and environmental effects of hazardous waste constituents. Quantitative estimates, including reference doses for chronic and subchronic duration for both inhalation and oral exposures, carcinogenic potency factors, unit risk estimates for air and drinking water, and reportable quantities (RQs) based on chronic toxicity and carcinogenicity are determined when sufficient data are available. HEEDs are subject to internal EPA review by staff within the Office of Health and Environmental Assessment and the Office of Pesticides and Toxic Substances. Final drafts of HEEDs become part of the RCRA and Superfund public dockets and are available through NTIS.

Health and Environmental Effects Profiles (HEEPs): This document series was prepared by the Environmental Criteria and Assessment Office (ECAO-Cincinnati) for the Office of Solid Waste and Emergency Response (OSWER). HEEPs have been superseded by HEEDs since mid-FY87. HEEPs are intended to support listings of hazardous constituents of a wide range of waste streams under Section 3001 of the Resource Conservation and Recovery Act (RCRA), as well as to provide health-related limits for emergency actions under Section 010 of the Comprehensive Environmental Response, Compensation and Liability Act (CERCLA). HEEPs are summaries of the

literature concerning health hazards associated with environmental exposures to chemicals or compounds and are very similar to HEEDs as described above. HEEPs were subject to internal EPA review by staff within the Office of Health and Environmental Assessment. HEEPS are part of the RCRA and CERCLA public dockets. Final drafts are available through NTIS.

Air Quality Criteria Documents (AQCDs): This document series is prepared by the Environmental Criteria and Assessment Office (ECAO-Research Triangle Park) for the Office of Air and Radiation (OAR). AQCDs are intended to support National Ambient Air Quality Standards (NAAQS) set under Sections 108-110 of the Clean Air Act. These documents are evaluations of the available scientific literature on the potential health effects of air pollutants. AQCDs are subject to internal EPA review by staff within the Office of Health and Environmental Assessment and the Office of Air and Radiation. Further review is conducted by the Science Advisory Board/Clean Air Scientific Advisory Committee, and then, these documents are subject to peer review workshops and public comments. The AQCDs are mandated by the Clean Air Act and are revised at 5-year intervals. AQCDs become part of the OAR public docket and final drafts are available through NTIS.

Health Assessment Documents (HADs): This document series is prepared by the Environmental Criteria and Assessment Office (ECAO-Research Triangle Park and ECAO-Cincinnati) for the Office of Air and Radiation (OAR). HADs are intended for use by the Office of Air Quality Planning and Standards (OAQPS) to determine possible listing of hazardous air pollutants (HAP) under sections 111 and 112 of the Clean Air Act. These documents are evaluations of the available scientific literature on the potential health effects of air pollutants and serve as the scientific data base for establishing relationships between exposure concentrations and potential health risks. HADs are subject to internal EPA review by staff within the Office of Health and Environmental Assessment and the Office of Air and Radiation. Further review is conducted by the Science Advisory Board/Clean Air Scientific Advisory Committee, and then, these documents are subject to peer review workshops and public comments. HADs become part of the OAR public docket and final drafts are available through NTIS.

#### 4. Miscellaneous Documents:

<u>Drinking Water Criteria Documents (DWCDs):</u> The Environmental Criteria and Assessment Office (ECAO-Cin) prepares a portion of this document series for the Office of Water (OW). DWCDs are intended to assist the OW in deriving criteria standards for chemicals in drinking water, as required under Section 412(b)(3)(A) of the Safe Drinking Water Act, as amended in 1986. The DWCDs are comprehensive evaluations of potential health effects, including mechanisms of toxicity, with specific emphasis on data providing dose-response information. DWCDs contain Health

Advisories (Has) for 1-day, 10-day and longer-term exposures and drinking water equivalent levels for lifetime exposures. DWCDs are subject to internal EPA review by staff within the Office of Health and Environmental Assessment and the Office of Water. Selected documents are reviewed by the Science Advisory Board and are subject to peer review workshops and public comments. DWCDs become part of the Safe Drinking Water (SDW) public docket and final drafts are available through NTIS.

#### B. Selection Criteria and Sources of HEAST Values

Chemicals with derived noncarcinogenic and/or carcinogen risk assessment values that have had some level of peer review (i.e., those in peer reviewed EPA documents or under review by EPA Work Groups) are included in HEAST; this does not include many interim values (values not found in final EPA documents or not being considered by Work Groups) derived for various purposes within Superfund and other Program Offices. In updating the HEAST, the first source that is checked is the Integrated Risk Information System (IRIS) for revised or newly added risk assessment values. Secondly, the status of chemicals under discussion by the RfD/RfC and CRAVE Work Groups is reviewed. The Office of Health and Environmental Assessment's Chemical Assessments and Related Activities (CARA) list is also reviewed for new Office of Water, Office of Air Quality Planning and Standards, and Office of Solid Waste and Emergency Response risk assessment documents (HEEDs, HEEPs, HEAs, HADs, AQCDs, DWCDs).

The HEAST also contains chemicals commonly found at RCRA (Resource Conservation and Recovery Act) sites as identified by the Office of Solid Waste's Technical Assessment Branch. Questions about RCRA chemicals may be addressed by calling the Health Assessment Section (Office of Solid Waste) at (202) 260-4761. Finally, the Office of Radiation Programs provides data on radionuclides for Table 4A and 4B of the HEAST. Radionuclides included are those thought to be most commonly encountered at Superfund sites. Questions concerning radionuclides carcinogenicity should first be addressed by contacting the appropriate Regional Radiation Program Manager. A listing of these managers and several contacts in the Office of Radiation Programs can be found in Exhibit 2 of the User's Guide -Radionuclide Carcinogenicity.

#### APPENDIX A-II

#### II. DOSE CONVERSIONS ON HEAST

In January 1991, the decision was made to replace inhalation Reference Doses (RfDi) for noncancer toxicity and inhalation slope factors for carcinogenicity, previously available on the IRIS data base, with Reference Concentrations (RFC) and inhalation unit risks, respectively. RfCs and unit risks are expressed in terms of concentration in air (mg/m³), not in terms of "dose" (mg/kg-day) like the RfDs and the oral and inhalation slope factors. This presents a problem for the Superfund program, since the current Hazard Ranking System (HRS) and the Risk Assessment Guidance for superfund (RAGS): Human Health Evaluation manual, Parts A and B were developed using chronic daily intakes and health criteria expressed in units of mg/kg-day.

The decision to replace inhalation slope factors and RfDi values expressed in mg/kg-day with unit risk and RfC values expressed in mg/m<sub>3</sub> was based on two major factors: 1) the workgroups felt that it was technically more accurate to base toxicity values directly on measured air concentrations instead of making the metabolic pharmacokinetic and/or surface area adjustments required to estimate an "internal dose"; and 2) there are compounds that elicit route-of-entry effects (e.g., sensitizers and irritants) where the toxic effect is to the respiratory system or exchange boundary where a measure of "internal dose" might inappropriately imply effects to other organ systems or effects from other exposure routes.

Superfund recognizes the importance of these issues and is actively working with EPA's Office of Research and Development to evaluate the impact of these changes on its program regulations and guidance. In the short term, however, modification of program regulations and guidance is not a viable option. Therefore, the chairs of the RfD/RfC and CRAVE Work Groups were consulted regarding Superfund's need to make the conversion from a concentration in air to dose. There was agreement that, in many cases, converting the air concentration data to a dose (in mg/kg-day) may not add significant uncertainty to the Superfund risk assessment process, and therefore may be a reasonable use of the data given appropriate circumstances and Superfund program objectives. These Work Groups will continue to work with the Superfund program to identify specific instances where it is not appropriate to make the conversion from unit risk/RfC to inhalation slope factor/RfD due to the large uncertainty introduced by the assumptions used in the conversion.

Generally, the Superfund Health Risk Technical Support Center will be responsible for making all appropriate conversions and the values will be identified with appropriate highlights or footnotes in the Health Effects Assessment Summary Tables (HEAST). Therefore, <u>HEAST users are strongly advised against making such conversions themselves</u>. However, it is a critical responsibility of the risk assessor to

consult the original reports cited in the HEAST and to appropriately characterize or caveat the resulting risk estimates derived from these values so that managers are fully informed of their origin and related uncertainties.

#### APPENDIX A-III

## II. CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE

This section lists chemicals and their respective Chemical Abstracts Service Registry Number (CASRN) for cross referencing on the HEAST. Chemicals may be searched either alphabetically by chemical name or numerically by the CASRN.

#### CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY NAME)

	ACENADUTUENE	000083-32-9	BENZININE	000003 87 5	04000W TETRACIII 0010C	000057 37 5
	ACENAT II IICHE	000208-96-8	DENZOIC ACID	000092-87-5 <b>000065-85-</b> 0	CHI COAL	000056-23-5
	ACEDUATE	030560-19-1	BENZOIC ACID	000003-03-0	CHLORAL	000075-87-6
	ACENAPHTHENE ACENAPTHYLENE ACEPHATE ACETONE ACETONE CYANOHYDRIN ACETONITRILE ACETOPHENONE ACROLEIN ACRYLAMIDE ACRYLONITRILE ADIPONITRILE ADIPONITRILE ALACHLOR ALDICARB ALDRIN ALLIDOCHLOR ALLYL ALCOHOL ALLYL CHLORIDE ALUMINUM PHOSPHIDE AMETRYN AMINO-2-NAPHTOL, 1- AMINO-2-NAPHTOL HYDROCHLORIDE, 1-	000067-64-1	BENZIDINE BENZOIC ACID BENZOTRICHLORIDE BENZO[A] ANTHRACENE BENZO[B] FLUORANTHENE BENZO[K] FLUORANTHENE BENZYL ALCOHOL BENZYL CHLORIDE BERYLLIUM BIPHENYL, 1,1' BIS(2-CHLOROETHYL) ETHER BIS(2-CHLOROISOPROPYL) ETHER BIS(2-ETHYLHEXYL) PHTHALATE /	000098-07-7	CARBON TETRACHLORIDE CHLORAL CHLORANIL CHLORDANE CHLORINE CYANIDE	000118-75-2
	ACETONE CYANOUVODIN	000075-86-5	BENZU (A) AN I NKAUENE	000056-55-3	CHLORDANE	000057-74-9
	ACCIONE CIANUTIDAIN	000077-00-3	BENZU (A) PIKENE	000050-32-8	CHLORINE CYANIDE	000506-77-4
	ACETONIFICHE	000075-05-8	SENZU (B) FLUURAN I HENE	000205-99-2	CHLORO-1,3-BUTADIENE, 2- / (CHLORO)	
	ACROLETA	000098-86-2	BENZU (K) FLUORAN I HENE	000207-08-9		000126-99-8
	ACRYLAMICS	000107-02-8	BENZYL ALCOHOL	000100-51-6	CHLORO-2-METHYLANILINE, 4-	000095-69-2
	ACRYLAMIDE	000079-06-1	BENZTL CHLORIDE	000100-44-7	CHLORO-2-METHYLANILINE HYDROCHLORIE	•
	ACRYLIC ACID	000079-10-7	BERYLLIUM	007440-41-7	CHLORO-M-CRESOL, P- CHLOROACETALDEHYDE CHLOROACETIC ACID CHLOROANILINE, 2- CHLOROANILINE, 3- CHLOROANILINE, 4- CHLOROBENZENE CHLOROBENZILATE CHLOROBENZOIC ACID, P- CHLOROBENZOIT IFLUORIDE, 4- CHLOROBENZOTRIFLUORIDE, 4-	003165-93-3
	ACRYLONITRILE	000107-13-1	BIPHENYL, 1,1'	000092-52-4	CHLORO-M-CRESOL, P-	000059-50-7
	ADIPONITRILE	000111-69-3	BIS(2-CHLOROETHYL) ETHER	000111-44-4	CHLOROACETALDEHYDE	000107-20-0
	ALACHLOR	015972-60-8	BIS(2-CHLOROISOPROPYL) ETHER	039638-32-9	CHLOROACETIC ACID	000079-11-8
	ALDICARB	000116-06-3	BIS(2-ETHYLHEXYL) PHTHALATE /	(DEHP) 000117-81-7	CHLOROANILINE, 2-	000095-51-2
	ALDRIN	000309-00-2	BIS(CHLOROMETHYL) ETHER	000542-88-1	CHLOROANILINE, 3-	000108-42-9
	ALLIDOCHLOR	000093-71-0	BISPHENOL A	000080-05-7	CHLOROANILINE, 4-	000106-47-8
	ALLYL ALCOHOL	000107-18-6	BIS(2-ETHYLHEXYL) PHTHALATE / BIS(CHLOROMETHYL) ETHER BISPHENOL A BORON, ELEMENTAL BORON TRIFLUORIDE BROMINATED DIBENZO-P-DIOXINS BROMINATED DIBENZOFURANS BROMOACETONE BROMOCHLOROETHANES BROMODICHLOROMETHANE BROMOCCOMM	007440-42-8	CHLOROBENZENE	000108-90-7
	ALLYL CHLORIDE	000107-05-1	BORON TRIFLUORIDE	007637-07-2	CHLOROBENZILATE	000510-15-6
	ALUMINUM	007429-90-5	BROMINATED DIBENZO-P-DIOXINS	NO CASRN	CHLOROBENZOIC ACID, P-	000074-11-3
	ALUMINUM PHOSPHIDE	020859-73-8	BROMINATED DIBENZOFURANS	NO CASRN	CHLOROBENZOTRIFLUORIDE, 4-	000098-56-6
$\triangleright$	AMETRYN	000834-12-8	BROMOACETONE	000598-31-2	CHLOROBUTANE. 1-	000109-69-3
A-1	AMINO-2-NAPHTHOL. 1-	002834-92-6	BROMOCHLOROETHANES	NO CASRN	CHLOROBUTANE 2-	000078-86-4
$\vec{\Box}$	AMINO-2-NAPHTOL HYDROCHLORIDE, 1-	001198-27-2	BROMODICHI OROMETHANE	000075-27-4	CHLOROBENZOTRIFLUORIDE, 4- CHLOROBUTANE, 1- CHLOROCYCLOPENTADIENE CHLOROFORM	041851-50-7
0	ANTINOPHENOL M-	000591-27-5	RROMOETHENE / (VINYL RROWIDE)	000503-60-2	CHI UBUEUBM	000067-66-3
	AMINOPHENOL O-	000095-55-6	BROMOFORM	000075-25-2	CHLOROMETHANE / (METHYL CHLORIDE)	000074-87-3
	AMINOPHENOL D.	000123-30-8	BROMOMETHANE	000073-23-2	CHLOROMETHYL METHYL ETHER	000074-87-3
	AMINOPHENOL, P	000504-24-5	BROMOPHENYL PHENYL ETHER, 4-	000101-55-3	CHLOROMETHTE METHTE ETHER CHLOROMITROBENZENE, M-	000107-30-2
	AMMOUTA	007664-41-7	BROMODUCE	000101-33-3	CHI COCALTOCRENTENE C	000121-73-3
	ANTITUE	00/004-41-7	BROMOVYNII	002104-96-3	CHI COCKLITECTENE, U	000088-73-3
	ANTLINE	000062-53-3	SKUTUK TRIL	001689-84-5	CHLORUNI I KOBENZENE, P-	000100-00-5
	AMINO-2-MAPHOL HYDROCHLORIDE, 1- AMINOPHENOL, 0- AMINOPHENOL, P- AMINOPYRIDINE, 4- AMMONIA ANILINE ANTHRACENE ANTIMONY, METALLIC ANTIMONY PENTOXIDE	000120-12-7	BROMOXYNIL OCIANOATE	001689-99-2	CHLOROPHENOL, 2-	000095-57-8
	ANTIMONY, METALLIC	007440-36-0	BUSAN //	031512-74-0	CHLOROPHENOL, 3-	000108-43-0
	ANTIMONY PENTOXIDE	001314-60-9	BUSAN 90	002491-38-5	CHLOROPHENOL, 4-	000106-48-9
	ANTIMONY POTASSIUM TARTRATE	000304-61-0	BUTADIENE, 1,3-	000106-99-0	CHLOROPROPANE, 2-	000075-29-6
	ANTIMONY TETROXIDE	001332-81-6	BUTANOL, 1-	000071-36-3	CHLOROTHALONII.	001897-45-6
	ANTIMONY TRIOXIDE	001309-64-4	BUTYL BENZYL PHTHALATE, N-	000085-68-7	CHLOROTOLUENE, M-	000108-41-8
	ARAMITE	000140-57-8	BUTYLATE	002008-41-5	CHLOROTOLUENE, O-	000095-49-8
	AROCHLOR 1248	012672-29-6	BUTYLCHLORIDE, T-	000507-20-0	CHLOROTOLUENE, P-	000106-43-4
	ARSENIC, INORGANIC	007440-38-2	BUTYROLACTONE, GAMMA-	000096-48-0	CHLORPYRIFOS	002921-88-2
	ASBESTOS	001332-21-4	CACODYLIC ACID	000075-60-5	CHLORPYRIFOS METHYL	005598-13-0
	ATRAZINE	001912-24-9	CADMIUM	007440-43-9	CHLORTHIOPHOS	060238-56-4
	AZOBENZENE	000103-33-3	CALCIUM CYANIDE	000592-01-8	CHROMIUM(III)	016065-83-1
	BARIUM	007440-39-3	CAPROLACTAM	000105-60-2	CHROMIUM(VI)	018540-29-9
	BARIUM CYANIDE	000542-62-1	CAPTAFOL	002425-06-1	CHRYSENE	000218-01-9
	ANTIMONY POISSION TAKKATE ANTIMONY TETROXIDE ANTIMONY TRIOXIDE ARAMITE AROCHLOR 1248 ARSENIC, INORGANIC ASBESTOS ATRAZINE AZOBENZENE BARIUM BARIUM CYANIDE BENEFIN BENEAL CHLORIDE BENZAL CHLORIDE BENZAL DENVOE CYANONYDBIN	001861-40-1	CAPTAN	000133-06-2	CHLORONITROBENZENE, M- CHLORONITROBENZENE, O- CHLORONITROBENZENE, P- CHLOROPHEMOL, 2- CHLOROPHEMOL, 3- CHLOROPHEMOL, 4- CHLOROPHEMOL, 2- CHLOROTHALONII. CHLOROTOLUENE, M- CHLOROTOLUENE, P- CHLOROTOLUENE, P- CHLOROTOLUENE, P- CHLOROTOLUENE, P- CHLOROTOLUENE, P- CHLOROTOLUENE, CHLOROPYRIFOS CHLORYRIFOS CHCORPYRIFOS CHCORYRIFOS CHROMIUM(III) CHROMIUM(III) CHROMIUM(III) CHROMIUM(III) CHROMIUM(III) CHROMIUM(III) COPER COME EMISSIONS COPPER COPPER CYANIDE CREOSOTE, COAL TAR CRESOL, M- / (3-METHYLPHENOL) CRESOL, O- / (2-METHYLPHENOL)	008007-45-2
	RENZAL CHIORIDE	000098-87-3	CARRARYI	000063-25-2	COPPER	007440-50-8
	RENZALDEHYDE	000100-52-7	CARRAZOI F	000086-74-8	COPPER CYANIDE	000544-92-3
	BENZALDEHYDE BENZALDEHYDE CYANOHYDRIN	000532-28-5	CAPROFIIDAN	001563-66-2	CREOSOTE COAL TAR	008001-58-9
	BENZENE	000071-43-2	CADDON DIGHTETOE	000075-15-0	CRECOL M. / /Z-METHYLDHENOLY	000108-39-4
	BENZENETHIOL / (THIOPHENOL)	000071-43-2	CARRON MONOVINE	000630-05-0	CRESOL, 0- / (2-METHYLPHENOL)	000108-39-4
	DENZEREINIUL / (INTURNERUL)	000100-90-3	BROMOPHENYL PHENYL ETHER, 4- BROMOPHOS BROMOXYNIL BROMOXYNIL OCTANOATE BUSAN 77 BUSAN 90 BUTADIENE, 1,3- BUTADIENE, 1,3- BUTYLATE BUTYL BENZYL PHTHALATE, N- BUTYLATE BUTYLCHLORIDE, T- BUTYROLACTONE, GAMMA- CACODYLIC ACID CAMHUM CALCIUM CYANIDE CAPPOLACTAM CAPTAFOL CAPTAN CARBAZOLE CARBOFURAN CARBON DISULFIDE CARBON MONOXIDE	000030-03-0	CRESCE, U- / (E-MEINTERNOL)	000073-40-7

### CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY NAME) continued

						000/00 04 7
	CRESOL, P- / (4-METHYLPHENOL)	000106-44-5	DICHLOROPHENOXY ACETIC ACID, 2,4-	000094-75-7	DINITROTOLUENE, 2,3-	000602-01-7
	CROTOMALDEHYDE CLIMENE CYANAZINE CYANOGEN CYANOGEN CYANOGEN CYANOGEN CYCLOATE CYCLOHEXANOL CYCLOHEXYLAMINE CYCLOPENTADIENE DACTHAL DALAPON DDD DDE DDT DECABROMODIPHENYL ETHER	000123-73-9	DICHLOROPHENOXY) BUTYRIC ACID, 4-(2,		DINITROTOLUENE, 2,4	000121-14-2
	CUMENE	000098-82-8	(2,4-DB)	000094-82-6	DINITROTOLUENE, 2,5-	000619-15-8
	CYANAZINE	021725-46-2	DICHLOROPROPANE, 1,1-	000078-99-9	DINITROTOLUENE, 2,6-	000606-20-2
	CYANIDE	000057-12-5	DICHLOROPROPANE, 1,2-	000078-87-5	DINITROTOLUENE, 3,4-	000610-39-9
	CYANOGEN	000460-19-5	DICHLOROPROPANE, 1,3-	000142-28-9	DINOSEB	000088-85-7
	CYANOGEN RROWIDE	000506-68-3	DICHLOROPROPANE, 2,2-	000594-20-7	DIOXANE, 1,4-	000123-91-1
	CYCLOATE	001134-23-2	DICHLOROPROPENE, 1,3- / (TELONE II)		DIPHENYLAMINE, N.N-	000122-39-4
	CACI UNEANIOI	000108-93-0	DICHLORPROP	000120-36-5	DIPHENYLHYDRAZINE, 1,2-	000122-66-7
	CYCLOHEANINE	000108-91-8	DICYCLOPENTADIENE	000077-73-6	DIRECT BLACK 38	001937-37-7
	CICLOREATEMENE	000168-91-8	DIELDRIN	000060-57-1	DIRECT BLUE 6	002602-46-2
	CICLOPERIADIERE	000342-72-7	DIETHYL-P-NITROPHENYL PHOSPHATE	000311-45-5	NIDECT BOOLW OF	016071-86-6
	DACINAL	001861-32-1	· ·	000311-43-3	DIRECT BROWN 95 DIRECT LIGHTFAST BLUE	004399-55-7
	DALAPON	000075-99-0	DIETHYL PHTHALATE	000084-66-2	DIRECT FIGHTAST SLUE	002610-05-1
	DDD	000072-54-8	DIETHYLANILINE, N,N-	000091-66-7	DIRECT SKY BLUE OB	002010-03-1
	DDE	000072-55-9	DIETHYLENE GLYCOL MONOBUTYL ETHER	000112-34-5	DISULFOTON	000298-04-4
	DDT	000050-29-3	DIETHYLENE GLYCOL MONOETHYL ETHER	000111-90-0	ENDOSULFAN	000115-29-7
	DECABROMODIPHENYL ETHER	001163-19-5	DIETHYLFORMAMIDE	000617-84-5	ENDOTHALL	000145-73-3
	DI-N-OCTYL PHTHALATE	000117-84-0	DIETHYLHYDRAZINE, 1,2-	001615-80-1	ENDRIN	000072-20-8
	DIALLATE	002303-16-4	DIETHYLSTILBESTROL	000056-53-1	EPICHLOROHYDRIN	000106-89-8
	DIAZINON	000333-41-5	DIMETHOATE	000060-51-5	EPTC	000759-94-4
$\triangleright$	DIBENZOFURAN	000132-64-9	DIMETHOXYBENZIDINE, 3,3'-	000119-90-4	ETHOPROP	013194-48-4
Y	DIBENZO (A, H) ANTHRACENE	000053-70-3	DIMETHYLANILINE, 2,4-	000095-68-1	ETHOXYETHANOL, 2-	000110-80-5
<u></u>	DIBROMO-3-CHLOROPROPANE, 1,2	000096-12-8	DIMETHYLANILINE HYDROCHLORIDE, 2,4-	021436-96-4	DIRECT LIGHTFAST BLUE DIRECT SKY BLUE 6B DISULFOTON ENDOSULFAN ENDOTHALL ENDRIN EPICHLOROHYDRIN EPTC ETHOPROP ETHOXYETHANOL, 2- ETHOXYETHANOL ACETATE, 2-	000111-15-9
_	DIBROMOBENZENE, 1,4-	000106-37-6	DIMETHYLANILINE, N,N-	000121-69-7	ETHOXYETHANOL ACRYLATE, 2-	000106-74-1
	DIBROMOCHLOROMETHANE	000124-48-1	DIMETHYLBENZIDINE, 3,3'-	000119-93-7	ETHOXYETHANOL DODECANOATE, 2-	000106-13-8
	DIBROMOETHANE, 1,2-	000106-93-4	DIMETHYLBENZ [A] ANTHRACENE, 7,12-	000057-97-6	ETHOXYETHANOL PHOSPHATE, 2-	068554-00-7
	DIBUTYL PHTHALATE	000084-74-2	DIMETHYLFORMAMIDE, N,N-	000068-12-2	ETHOXYETHYL METHACRYLATE, 2-	002370-63-0
	DICAMBA	001918-00-9	DIMETHYLHYDRAZINE, 1,1-	000057-14-7	ETHYL ACETATE	000141-78-6
	DICHLORO-2-BUTENE, 1,4-	000764-41-0	DIMETHYLHYDRAZINE, 1,2-	000540-73-8	ETHYL ACRYLATE ETHYL BENZENE ETHYL CHLORIDE ETHYL ETHER ETHYL METHACRYLATE ETHYL-O-XYLENE, 4-	000140-88-5
		000095-50-1	DIMETHYLPHENOL, 2,3-	000526-75-0	FTHY! RENZENE	000100-41-4
	DICHLOROBENZENE, 1,2-			000105-67-9	ETHYL CHIODIDE	000075-00-3
	DICHLOROBENZENE, 1,3-	000541-73-1	DIMETHYLPHENOL, 2,4-	000105-87-4	CTUVI CTUCD	000060-29-7
	DICHLOROBENZENE, 1,4-	000106-46-7	DIMETHYLPHENOL, 2,5-		CIRIL CIRER	000097-63-2
	DICHLOROBENZIDINE, 3,3'-	000091-94-1	DIMETHYLPHENOL, 2,6-	000576-26-1	EIRIL MEIRACKILNIE	000934-80-5
	DICHLOROBUTENES	NO CASRN	DIMETHYLPHENOL, 3,4-	000095-65-8	EINTL-U-ATLEME, 4-	000103-69-5
	DICHLORODIFLUOROMETHANE	000075-71-8	DIMETHYLPHTHALATE	000131-11-3	ETHYLANILINE, N-	000109-78-4
	DICHLOROETHANE, 1,1-	000075-34-3	DIMETHYLSULFATE	000077-78-1	ETHYLENE CYANOHYDRIN	
	DICHLOROETHANE, 1,2-	000107-06-2	DIMETHYLTEREPHTHALATE	000120-61-6	ETHYLENE DIAMINE	000107-15-3
	DICHLOROETHYLENE, 1,1-	000075-35-4	DIMETHYLUREA, N,N-	000598-94-7	ETHYLENE GLYCOL	000107-21-1
	DICHLOROETHYLENE, 1,2- (MIXED	ISOMERS)	DINITRO-O-CRESOL, 4,6-	000534-52-1	ETHYLENE GLYCOL MONOBUTYL ETHER	000111-76-2
	• •	000540-59-0	DINITRO-P-CRESOL, 2,6-	000609-93-8	ETHYLENE OXIDE	000075-21-8
	DICHLOROETHYLENE, 1,2-C-	000156-59-2	DINITROBENZENE, 1,2-	000528-29-0	ETHYLENE THIOUREA	000096-45-7
	DICHLOROETHYLENE, 1,2-T-	000156-60-5	DINITROBENZENE, 1,3-	000099-65-0	ETHYLTOLUENE, M-	000620-14-4
	DICHLOROPHENOL, 2,3-	000576-24-9	DINITROBENZENE, 1,4-	000100-25-4	ETHYLTOLUENE, O-	000611-14-3
	DICHLOROPHENOL, 2,4-	000120-83-2	DINITROPHENOL, 2,3-	000066-56-8	ETHYLTOLUENE, P-	000622-96-8
	DICHLOROPHENOL, 2,5-	000583-78-8	DINITROPHENOL, 2,4-	000051-28-5	FLUORANTHENE	000206-44-0
	DICHLOROPHENOL, 2,5-	000087-65-0	DINITROPHENOL, 2,5-	000329-71-5	FLUORENE	000086-73-7
		000095-77-2	DINITROPHENOL, 2,6-	000573-56-8	FLUORINE / (SOLUBLE FLUORIDE)	007782-41-4
	DICHLOROPHENOL, 3,4-			000586-11-8	FLURIDONE	059756-60-4
	DICHLOROPHENOL, 3,5-	000591-35-5	DINITROPHENOL, 3,5-	000000 11 0	LAUTAME	

### CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY NAME) Continued

FOLPET 000133-07-3 MERCURY, ELEMENTAL VAPOR 007439-97-6 MOLINATE 002212-67-1 007439-97-6 007439-98-7 000050-00-0 **FORMALDEHYDE** MERCURY, INORGANIC MOLYBDENLIM 010599-90-3 FORMALDEHYDE CYANOHYDRIN 000107-16-4 **MERPHOS** 000150-50-5 MONOCHLORAMINE 000091-20-3 000064-18-6 MERPHOS OXIDE 000078-48-8 NAPHTHALENE FORMIC ACID 000126-98-7 000130-15-4 FLIRAM 000110-00-9 METHACRYLONITRILE NAPHTHOQUINONE, 1.4-002429-74-5 000067-45-8 METHOMYL 016752-77-5 NIAGARA BLUE 48 **FURAZOL IDONE** 007440-02-0 000098-01-1 METHOXY-5-NITROANILINE, 2-000099-59-2 NICKEL (METALLIC) **FURFURAL** 000531-82-8 METHOXYCHLOR 000072-43-5 NICKEL CYANIDE 000557-19-7 **FURIUM** 000765-34-4 000109-86-4 **NICKEL REFINERY DUST** NO CASRN METHOXYETHANOL, 2-**GLYCIDALDEHYDE** 000076-44-8 METHOXYETHANOL ACETATE, 2-000110-49-6 NICKEL SUBSULFIDE 012035-72-2 **HEPTACHLOR** 000100-54-9 001024-57-3 METHYL-4-CHLOROPHENOXY) BUTYRIC ACID, 4-(2-**NICOTINONITRILE** HEPTACHLOR EPOXIDE 010102-43-9 HEPTANE. N-000142-82-5 000094-81-5 NITRIC OXIDE 014797-65-0 000087-82-1 METHYL-4-CHLOROPHENOXY) PROPIONIC ACID, 2-(2-MITRITE **HEXABROMOBENZENE** 000088-74-4 000118-74-1 000093-65-2 NITROANILINE, 2-**HEXACHLOROBENZENE** 000087-68-3 NITROANILINE, M-000099-09-2 METHYL-4-CHLOROPHENOXY ACETIC ACID, 2-**HEXACHLOROBUTADIENE** 000094-74-6 NITROANILINE, P-000100-01-6 000319-84-6 HEXACHLOROCYCLOHEXANE, ALPHA-000319-85-7 000099-55-8 NITROBENZENE 000098-95-3 HEXACHLOROCYCLOHEXANE, BETA-METHYL-5-NITROANILINE, 2-000079-20-9 000067-20-9 000319-86-8 NITROFURANTOIN HEXACHLOROCYCLOHEXANE, DELTA-METHYL ACETATE 000059-87-0 HEXACHLOROCYCLOHEXANE, EPSILON-006108-10-7 METHYL ACRYLATE 000096-33-3 NITROFURAZONE 010102-44-0 000058-89-9 METHYL CHLOROCARBONATE 000079-22-1 NITROGEN DIOXIDE HEXACHLOROCYCLOHEXANE, GAMMA-NO CASRN 000608-73-1 METHYL ETHYL KETOME 000078-93-3 **NITROGEN OXIDES** HEXACHLOROCYCLOHEXANE-TECHNICAL 000077-47-4 METHYL ETHYL KETONE PEROXIDE 001338-23-4 NITROMETHANE 000075-52-5 - HEXACHLOROCYCLOPENTADIENE 000060-34-4 MITROPHEMOLS NO CASRN N HEXACHLOROETHANE 000067-72-1 METHYL HYDRAZINE NITROPROPANE. 2-000070-30-4 METHYL ISOBUTYL KETONE 000108-10-1 000079-46-9 **HEXACHLOROPHENE** 000624-83-9 NITROSO-DI-N-BUTYLAMINE, N-000924-16-3 000124-09-4 METHYL ISOCYANATE HEXAMETHYLENE DIAMINE 022967-92-6 NITROSO-DI-N-PROPYLAMINE, N-000621-64-7 000110-54-3 METHYL MERCURY HEXANE, N-000080-62-6 NITROSO-N-ETHYLUREA. N-000759-73-9 000591-78-6 METHYL METHACRYLATE HEXANONE, 2-000298-00-0 000684-93-5 NITROSO-N-METHYLUREA, N-000302-01-2 METHYL PARATHION HYDRAZINE 010034-93-2 METHYL STYRENE (MIXED ISOMERS) 025013-15-4 NITROSODIETHANOLAMINE, N-001116-54-7 HYDRAZINE SULFATE 000055-18-5 000098-83-9 007783-06-4 METHYL STYRENE, ALPHA NITROSODIETHYLAMINE. N-HYDROGEN SULFIDE 000062-75-9 000123-31-9 METHYLANILINE, 2-000095-53-4 NITROSODIMETHYLAMINE, N-**HYDROQUINONE** 000156-10-5 METHYLANILINE HYDROCHLORIDE, 2-000636-21-5 NITROSODIPHENYLAMINE, P-000193-39-5 INDENO[1,2,3-CD] PYRENE 000086-30-6 000056-49-5 NITROSODIPHENYLAMINE, N-007439-89-6 METHYLCHOLANTHRACENE, 3-000108-87-2 NITROSOMETHYLETHYLAMINE, N-010595-95-6 000078-83-1 METHYLCYCLOHEXANE I SOBUTYL ALCOHOL 004549-40-0 000078-59-1 METHYLENE-BIS(2-CHLOROANILINE), 4,41-NITROSOMETHYLVINYLAMINE. N I SOPHORONE 000930-55-2 000101-14-4 NITROSOPYRROLIDINE, N-033820-53-0 I SOPROPAL I N 000099-08-1 METHYLENE-BIS(N, N'-DIMETHYL)ANILINE, 4,41-NITROTOLUENE, M-000078-97-7 LACTONITRILE NITROTOLUENE, O-000088-72-2 000101-61-1 LEAD 007439-92-1 000099-99-0 000074-95-3 NITROTOLUENE. P-NO CASRN METHYLENE BROMIDE LEAD ALKYLS OCTABROMODIPHENYL ETHER 032536-52-0 000330-55-2 METHYLENE CHLORIDE / (DICHLOROMETHANE) LINURON **OCTAMETHYLPYROPHOSPHORAMIDE** 000152-16-9 000075-09-2 000121-75-5 MALATHION 020816-12-0 OSMIUM TETROXIDE 000108-31-6 METHYLENE-BIS(BENZENEAMINE), 4.4- / MALEIC ANHYDRIDE 000101-77-9 OZONE 010028-15-6 000123-33-1 (METHYLENE DIANILINE, 4.4-) MALEIC HYDRAZIDE 000123-63-7 **PARALDEHYDE** METHYLENEDIPHENYL ISOCYANATE, 4,4- / MALONONITRILE 000109-77-3 (DIPHENYLMETHANE DIISOCYANATE) 000101-68-8 000056-38-2 008018-01-7 **PARATHION** MANCOZEB PARTICULATE MATTER NO CASRN 051218-45-2 012427-38-2 **METOLACHLOR** MANES 001114-71-2 007439-96-5 **METRIBUZIN** 021087-64-9 PEBULATE MANGANESE 040487-42-1 002385-85-5 PENDIMETHALIN 000950-10-7 MIREX **MEPHOSFOLAN** 

### CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY NAME) continued

PENTABROMO-6-CHLOROCYCLOHEXANE, 1,2,3,4,5-000506-64-9 002687-25-4 SILVER CYANIDE TOLUENEDIAMINE, 2,3-000087-84-3 000122-34-9 TOLUENEDIAMINE, 3,4-000496-72-0 SIMAZINE PENTABROMODIPHENYL ETHER 032534-81-9 000143-33-9 000108-44-1 SODIUM CYANIDE TOLUIDINE, M-000106-49-0 PENTACHLOROBENZENE 000608-93-5 SODIUM DIETHYLDITHIOCARBAMATE 000148-18-5 TOLUIDINE. P-025329-35-5 013718-26-8 008001-35-2 **PENTACHLOROCYCLOPENTADIENE** SODIUM METAVANADATE TOXAPHENE 002303-17-5 **PENTACHLORONITROBENZENE** 000082-68-8 STRONTIUM, STABLE 007440-24-6 TRIALLATE 000615-54-3 **PENTACHLOROPHENOL** 000087-86-5 000057-24-9 TRIBROMOBENZENE, 1,2,4-STRYCHNINE PENTACHLOROPROPENE, 1,1,2,3,3,-001600-37-9 STYRENE 000100-42-5 TRICHLORO-1,2,2-TRIFLUOROETHANE, 1,1,2-000109-66-0 000110-61-2 000076-13-1 PENTANE. N-SUCCINONITRILE PHENANTHRENE 000085-01-8 007446-09-5 TRICHLORO-2'-HYDROXYDIPHENYLETHER, 2,2,4'-SULFUR DIOXIDE PHENOL 000108-95-2 SULFUR OXIDES NO CASRN 003380-34-5 000634-93-5 000108-45-2 007664-93-9 PHENYLENEDIAMINE, M-SULFURIC ACID TRICHLOROANILINE, 2,4,6-PHENYLENEDIAMINE. O-000095-54-5 TCDD, 2,3,7,8-001746-01-6 TRICHLOROANILINE HYDROCHLORIDE, 2.4.6-033663-50-2 000106-50-3 003383-96-8 PHENYLENEDIAMINE. P-**TEMEPHOS** 000120-82-1 PHENYLMERCURIC ACETATE 000062-38-4 013071-79-9 TRICHLOROBENZENE, 1,2,4-**TERBUFOS** 000090-43-7 000100-21-0 TRICHLOROCYCLOPENTADIENE 077323-84-3 PHENYLPHENOL. 2-TEREPHTHALIC ACID 000071-55-6 000298-02-2 021232-47-3 TRICHLOROETHANE, 1,1,1-**TETRACHLOROAZOXYBENZENE** PHORATE 000079-00-5 **PHOSGENE** 000075-44-5 TETRACHLOROBENZENE, 1,2,4,5-000095-94-3 TRICHLOROETHANE, 1,1,2-000079-01-6 000695-77-2 PHOSPHINE 007803-51-2 **TETRACHLOROCYCLOPENTADIENE** TRICHLOROETHYLENE 000075-69-4 PHOSPHORUS, WHITE 007723-14-0 TETRACHLOROETHANE, 1,1,1,2-000630-20-6 TRICHLOROFLUOROMETHANE 015950-66-0 NO CASRN 000079-34-5 TRICHLOROPHENOL, 2,3,4-PHOTOCHEMICAL OXIDANTS TETRACHLOROETHANE, 1,1,2,2-000933-78-8 PHTHALIC ACID, M-000121-91-5 TETRACHLOROETHYLENE 000127-18-4 TRICHLOROPHENOL, 2,3,5-071753-42-9 TRICHLOROPHENOL, 2,3,6-000933-75-5 PHTHALIC ACID, O-000088-99-3 **TETRACHLOROHYDRAZOBENZENE** 000095-95-4 000100-21-0 004901-51-3 TRICHLOROPHENOL, 2,4,5-PHTHALIC ACID, P-TETRACHLOROPHENOL. 2.3.4.5-000088-06-2 000085-44-9 000058-90-2 TRICHLOROPHENOL, 2.4.6-PHTHALIC ANHYDRIDE TETRACHLOROPHENOL, 2,3,4,6-000609-19-8 000935-95-5 NO CASRN TETRACHLOROPHENOL, 2,3,5,6-TRICHLOROPHENOL, 3,4,5-POLYBROMINATED BIPHENYLS TRICHLOROPHENOXY) PROPIONIC ACID, 2(2,4,5-POLYCHLORINATED BIPHENYLS 001336-36-3 TETRACHLOROPROPENE, 1,1,2,3-010436-39-2 000093-72-1 000151-50-8 TETRACHLOROTOLUENE, PARA, ALPHA, ALPHA, ALPHA-POTASSIUM CYANIDE TRICHLOROPHENOXY ACETIC ACID, 2,4,5-000506-61-6 005216-25-1 POTASSIUM SILVER CYANIDE 0000093-76-5 000961-11-5 026399-36-0 TETRACHLOROVINPHOS / (STIROPHOS) PROFLURALIN 007789-89-1 003689-24-5 023950-58-5 TRICHLOROPROPANE, 1,1,1-PRONAMIDE TETRAETHYL DITHIOPYROPHOSPHATE 000598-77-6 001314-32-5 TRICHLOROPROPANE, 1,1,2-**PROPACHLOR** 001918-16-7 THALLIC OXIDE TRICHLOROPROPANE, 1,2,2-003175-23-3 000563-68-8 000139-40-2 THALLIUM (I) ACETATE PROPAZINE 000096-18-4 000107-12-0 006533-73-9 TRICHLOROPROPANE, 1,2,3-**PROPIONITRILE** THALLIUM (I) CARBONATE 000096-19-5 TRICHLOROPROPENE, 1,2,3-007791-12-0 PROPYL ALCOHOL, N-000071-23-8 THALLIUM (I) CHLORIDE 002077-46-5 TRICHLOROTOLUENE, 2,3,6-PROPYLENE GLYCOL 000057-55-6 THALLIUM (I) NITRATE 010102-45-1 002014-83-7 007446-18-6 TRICHLOROTOLIJENE, ALPHA, 2, 6-PROPYLENE GLYCOL MONOMETHYL ETHER 000107-98-2 THALLIUM (I) SULFATE 001582-09-8 NO CASRN TRIFLURALIN PROPYLENE OXIDE 000075-56-9 THALLIUM (IN SOLUBLE SALTS) 012039-52-0 TRIMETHYL PHOSPHATE 000512-56-1 000129-00-0 THALLIUM SELENITE PYRENE NO CASRN 000110-86-1 THIOCYANOMETHYLTHIO)BENZOTHIAZOLE, 2-( TRIMETHYLBENZENES PYRIDINE 000099-35-4 021564-17-0 TRINITROBENZENE, 1,3,5-QUINOLINE 000091-22-5 NO CASRN 013196-18-4 TRINITROPHENOLS 000121-82-4 THIOFANOX RDX / (CYCLONITE) 000479-45-8 TRINITROPHENYLMETHYLNITRAMINE RONNEL 000299-84-3 THIRAM 000137-26-8 000118-96-7 NO CASRN TRINITROTOLUENE, 2,4,6-007783-00-8 TIN AND COMPOUNDS SELENIOUS ACID URANIUM, SOLUBLE SALTS NO CASRN 007782-49-2 TOLUENE 000108-88-3 SELENIUM 000095-80-7 **VANADIUM** 007440-62-2 007446-34-6 TOLUENE-2,4-DIAMINE SELENIUM SULFIDE 001314-62-1 000095-70-5 **VANADIUM PENTOXIDE** 000630-10-4 TOLUENE-2.5-DIAMINE SELENOUREA 036907-42-3 007440-22-4 000823-40-5 VANADIUM SULFATE TOLUENE-2.6-DIAMINE SILVER

NAME/CASRN-4

### CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY NAME) Continued

VERNAM / (VERNOLATE)	001929-77-7
VINYL-1-CYCLOHEXENE, 4-	000100-40-3
VINYL ACETATE	000108-05-4
VINYL CHLORIDE	000075-01-4
WARFARIN	000081-81-2
XYLENE, M-	000108-38-3
XYLENE, MIXTURE	001330-20-7
XYLENE, O-	000095-47-6
XYLENE, P-	000106-42-3
ZINC (METALLIC)	007440-66-6
ZINC CYANIDE	000557-21-1
ZINC PHOSPHIDE	001314-84-7
ZINEB	012122-67-7

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	000050-32-8	BENZO (A) PYRENE	000074-83-9	BROMOMETHANE	000079-22-1	METHYL CHLOROCARBONATE
	000050-00-0	FORMALDEHYDE	000074-87-3	CHLOROMETHANE / (METHYL CHLORIDE)	000080-62-6	METHYL METHACRYLATE
	000050-29-3	DDT	000074-95-3	METHYLENE BROWIDE	000080-05-7	BISPHENOL A
	000051-28-5	DINITROPHENOL, 2.4-	000074-11-3	CHLOROBENZOIC ACID. P-	000081-81-2	WARFARIN
	000053-70-3	DIBENZO(A, H) ANTHRACENE	000075-35-4	DICHLOROETHYLENE. 1.1-	000082-68-8	PENTACHLORONITROBENZENE
	000055-18-5	NITROSODIETHYLAMINE. N-	000075-60-5	CACODYLIC ACID	000083-32-9	ACENAPHTHEME
	000056-53-1	DIETHYLSTILBESTROL	000075-01-4	VINYL CHLORIDE	000084-66-2	DIFTHY! PHTHALATE
	000056-38-2	PARATHION	000075-21-8	ETHYLENE OXIDE	000084-74-2	DIRITY! PHTHALATE
	000056-55-3	BENZO [A] ANTHRACENE	000075-15-0	CARBON DISULFIDE	000085-01-8	PHENANTHRENE
	000056-23-5	CARBON TETRACHLORIDE	000075-86-5	ACETOME CYANOHYDRIN	000085-68-7	RIITYI RENZYI PHTHALATE N-
	000056-49-5	METHYLCHOLANTHRACENE. 3-	000075-09-2	METHYLENE CHLORIDE /	000085-44-9	PHTHALIC ANHYDRIDE
	000057-74-9	CHLORDANE		(DICHIOPOMETHANE)	000085-73-7	FILMPENE
	000057-12-5	CYANIDE	000075-27-4	RPOMODICAL OPOMETHANE	A-02-0000	NITPOSODIDHENYI AMINE N-
	000057-14-7	DIMETHYLHYDRAZINE 1 1-	000075-00-3	FTHYI CHIOPINE	000086-74-8	CAPRAZOI F
	000057-55-6	PROPYLENE GLYCOL	000075-69-4	TRICHI OROFI HOROMETHANE	000087-68-3	HEYACHI ORORUTAD LENE
	000057-97-6	DIMETHYLBENZ (A) ANTHRACENE 7 12-	000075-25-2	RECHOEOROI ECONOTIETHARE	000007-84-3	DENTARDOMO-A-CHI OPOCYCI OHEYANE
	000057-24-9	STRYCHNINE	000075 25 2	CHI OPAI	000001 04 3	1 2 3 4 5-
	000051-89-9	HEXACHI OROCYCI OHEXANE GAMMA-	000075-71-8	DICHI DODI ELLIDDOMETHANE	000087-65-0	DICHIODODHENOI 2 K-
	000058-90-2	TETRACHI OROPHENOL 2 3 4 6-	000075 71 0	DAI ADON	000007-86-5	DENTACHI OPODHENOL
	000059-87-0	NITROFURAZONE	000075-05-8	ACETOMITRILE	000007-82-1	HEYARDOMORENZENE
	000059-50-7	CHLORO-M-CRESOL P-	000075-29-6	CHI OPOPPOPANE 2-	000007 02 7	PHTHALIC ACID O-
•	000060-34-4	METHYL HYDRAZINE	000075-44-5	PHOSCENE	000000 77 3	NITPOTOLIENE O-
7	000060-57-1	DIFLORIN	000075-52-5	NITROMETHANE	000088-06-2	TRICHIOPOPHENOL 2 4 6-
•	000060-29-7	ETHYL ETHER	000075-34-3	DICHLOROFTHANE 1 1-	000088-74-4	MITROANILINE 2-
	000060-51-5	DIMETHOATE	000075-56-9	PROPYLENE OXIDE	000088-73-3	CHI OPONITRORENZENE O-
	000062-38-4	PHENYLMERCURIC ACETATE	000076-13-1	TRICHIORO-1 2 2-TRIFILIOROFTHANE	000088-85-7	DINOSER
	000062-53-3	ANILINE		1.1.2-	000090-43-7	PHENYLPHENOL 2-
	000062-75-9	NITROSODIMETHYLAMINE. N-	000076-44-8	HEPTACHLOR	000091-66-7	DIETHYLANILINE. N.N-
	000063-25-2	CARBARYL	000077-78-1	DIMETHYLSULFATE	000091-94-1	DICHLOROBENZIDINE, 3.3'-
	000064-18-6	FORMIC ACID	000077-47-4	HEXACHLOROCYCLOPENTAD I ENE	000091-22-5	QUINOLINE
	000065-85-0	BENZOIC ACID	000077-73-6	DICYCLOPENTADIENE	000091-20-3	NAPHTHALENE
	000066-56-8	DINITROPHENOL, 2.3-	000078-99-9	DICHLOROPROPANE, 1.1-	000092-52-4	BIPHENYL. 1.1'
	000067-45-8	FURAZOLIDONE	000078-87-5	DICHLOROPROPANE, 1,2-	000092-87-5	SENZIDINE
	000067-72-1	HEXACHLOROETHANE	000078-86-4	CHLOROBUTANE, 2-	000093-76-5	TRICHLOROPHENOXY ACETIC ACID.
	000067-66-3	CHLOROFORM	000078-48-8	MERPHOS OXIDE		2.4.5-
	000067-20-9	NITROFURANTOIN	000078-97-7	LACTONITRILE	000093-71-0	ALLIDOCHLOR
	000067-64-1	ACETONE	000078-83-1	I SOBUTYL ALCOHOL	000093-72-1	TRICHLOROPHENOXY) PROPIONIC ACID.
	000068-12-2	DIMETHYLFORMAMIDE, N.N-	000078-93-3	METHYL ETHYL KETONE		2(2.4.5-
	000070-30-4	HEXACHLOROPHENE	000078-59-1	ISOPHORONE	000093-65-2	METHYL-4-CHLOROPHENOXY) PROPIONIC
	000071-23-8	PROPYL ALCOHOL. N-	000079-06-1	ACRYLAMIDE		ACID. 2-(2-
	000071-36-3	BUTANOL, 1-	000079-20-9	METHYL ACETATE	000094-82-6	DICHLOROPHENOXY) BUTYRIC ACID, 4-
	000071-55-6	TRICHLOROETHANE, 1,1,1-	000079-34-5	TETRACHLOROETHANE, 1,1,2,2-		(2.4- / (2.4-DB)
	000071-43-2	BENZO [A] PYRENE FORMALDEHYDE DDT DINITROPHENOL, 2,4- DIBENZO [A, H] ANTHRACENE NITROSODIETHYLAMINE, N- DIETHYLSTILBESTROL PARATHION BENZO [A] ANTHRACENE CARBON TETRACHLORIDE METHYLCHOLANTHRACENE, 3- CHLORDANE CYANIDE DIMETHYLHYDRAZINE, 1,1- PROPYLENE GLYCOL DIMETHYLBENZ [A] ANTHRACENE, 7,12- STRYCHNINE HEXACHLOROCYCLOHEXANE, GAMMA- TETRACHLOROPHENOL, 2,3,4,6- NITROFURAZONE CHLORO-M-CRESOL, P- METHYL HYDRAZINE DIELDRIN ETHYL ETHER DIMETHOATE PHENYLMERCURIC ACETATE ANILINE NITROSODIMETHYLAMINE, N- CARBARYL FORMIC ACID BENZOIC ACID DINITROPHENOL, 2,3- FURAZOLIDONE HEXACHLOROETHANE CHLOROFORM NITROFURANTOIN ACETONE DIMETHYL FORMAMIDE, N,N- HEXACHLOROETHANE CHLOROFORM NITROFURANTOIN ACETONE DIMETHYLFORMAMIDE, N,N- HEXACHLOROETHANE, 1,1,1- BENZENE ENDRIN DDE METHOXYCHLOR DDD	000079-46-9	NITROPROPANE, 2-	000094-75-7	DICHLOROPHENOXY ACETIC ACID, 2,4-
	000072-20-8	ENDRIN	000079-01-6	TRICHLOROETHYLENE	000094-74-6	METHYL-4-CHLOROPHENOXY ACETIC
	000072-55-9	DDE	000079-00-5	TRICHLOROETHANE, 1,1,2-		ACID, 2-
	000072-43-5	METHOXYCHLOR	000079-11-8	CHLOROACETIC ACID		•
	000072-54-8	DOD	000079-10-7	ACRYLIC ACID		
	· <del>-</del>					

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	000094-81-5	METHYL-4-CHLOROPHENOXY) BUTYRIC	000100-40-3	VINYL-1-CYCLOHEXENE, 4-	000108-95-2	PHENOL
		ACID, 4-(2-	000100-25-4	DINITROBENZENE, 1,4-	000108-88-3	TOLUENE
	000095-53-4	METHYLANILINE, 2-	000100-44-7	BENZYL CHLORIDE	000108-98-5	BENZENETHIOL / (THIOPHENOL)
	000095-51-2	CHLOROANILINE, 2-	000100-41-4	ETHYL BENZENE	000108-42-9	CHLOROANILINE, 3-
	000095-70-5	TOLUENE-2.5-DIAMINE	000100-54-9	NICOTINONITRILE	000108-93-0	CYCLOHEXANOL
	000095-95-4	TRICHLOROPHENOL, 2.4.5-	000100-21-0	PHTHALIC ACID. P-	000108-45-2	PHENYLENEDIAMINE. M-
	000095-50-1	DICHLOROBENZENE. 1.2-	000100-01-6	NITROANILINE. P-	000108-91-8	CYCLOHEXYLAMINE
	000095-48-7	CRESOL. Q- / (2-METHYLPHENOL)	000101-55-3	BROMOPHENYL PHENYL ETHER. 4-	000108-39-4	CRESOL, M- / (3-METHYLPHENOL)
	000095-80-7	TOLLIENE-2.4-DIAMINE	000101-14-4	METHYLENE-BIS(2-CHLOROANILINE).	000108-41-8	CHLOROTOLUENE. M-
	000095-69-2	CHLORO-2-METHYLANILINE. 4-		4.41-	000108-05-4	VINYL ACETATE
	000095-87-4	DIMETHYLPHENOL 2.5-	000101-68-8	METHYLENEDIPHENYL ISOCYANATE 4.4-	000108-31-6	MALEIC ANHYDRIDE
	000095-49-8	CHLOROTOLLIENE. O-		/ (DIPHENYLMETHANE DIISOCYANATE)	000108-44-1	TOLUIDINE. M-
	000095-47-6	XYLENE O-	000101-61-1	METHYLENE-RISON NI-DIMETHYL DANIL IN	000108-43-0	CHLOROPHENOL 3-
	000075 47 5	TETRACHI ORORENZENE. 1.2.4.5-		F. 4.41-	000108-90-7	CHLOROBENZENE
	000095-57-8	CHI OROPHENOL . 2-	000103-33-3	AZOBENZENE	000109-86-4	METHOXYETHANOL . 2-
	000075-54-5	PHENYLENEDIAMINE O-	000103-69-5	ETHYLANILINE N-	000109-69-3	CHLOROBLITANE . 1-
	000075 37 3	DICHLOPOPHENOL 3 4-	000105-60-2	CAPPOLACTAM	000109-78-4	ETHYLENE CYANOHYDRIN
	000075 17 2	DIMETHYL PHENOL 3 4-	000105-67-9	DIMETHYLPHENOL 2 4-	000109-66-0	PENTANE. N-
	000075-68-1	DIMETHYLANTI THE 2 4-	000106-48-9	CHI OROPHENOL 4-	000109-77-3	MALONONITRILE
	000075-55-6	AMIMOPHEMOL O-	000106-37-6	DIRROMORENZENE 1 4-	000110-54-3	HEXAME. N-
_	000075 55 0	ETHYLENE THICHDEA	000106-93-4	DIRROMOETHANE 1 2-	000110-86-1	PYRIDINE
<b>P</b>	0-84-200000	RITYPOLACTONE CAMMA-	000106-67-8	CHI OPPOANTI INF 4-	000110-61-2	SUCCINOMITRILE
<u>ن</u>	000090-40-0	DIRDOMO-T-CHI ODODDODANE 1 2	4-54-A01000	CHI OPOTOLLIENE P-	000110-69-6	METHOXYETHANOL ACETATE 2-
တ	000076-12-5	TRICHI COCOBCOEME 1 2 3-	000106-43-4	CRESCI D- / (A-METHYL PHENOL)	000110 47 5	STHONYETHANOL 2-
	000096-17-3	METHYL ACOVIATE	000106-47-5	EDICHI ODOHADDIN	000110-00-9	FIRAM
	000096-33-3	TRICHIOPODPODANE 1 2 %.	000100-07-0	NICHIADARENZENE 1 4-	000110-00-7	ETHOMYETHANOL ACETATE 2-
	00007-43-2	ETHYL METHAPOVIATE	000100 40 7	BUTANIENE 1 %	000111-90-0	DIETHYLENE GLYCOL MONOETHYL ETHER
	000097-03-2	DENTAL CHICOTOE	000100-99-0	STHOUGHT PODECAMONTE 2-	000111-76-2	ETHYLENE GLYCOL MONOBUTYL ETHER
	000098-01-1	ENDERDAL CHLOKIDE	Z-C1-201000	VVIEWE D.	A-44-L	RIS/2-CHI OPOETHY! ) FTHER
	000096-01-1	METHYL STYDENE ALDUA	000100-42-3	DUENVIEWEDIAMINE D.	000111-40-3	ANIPONITALLE
	000076-05-7	MITAGENICAE	000106-20-3	TOURDING D.	000111 07 5	DIETHYLENE GLYCOL MONOBUTYL ETHER
	000000-93-9	MI I KUBENZENE	000106-49-0	ETHOVYETHANOL ACRYLATE 2-	000112-34-3	ENDOGUI FAN
	000000 04 3	ACCOUNTYOUT	000100-74-1	ACOMEIN	000113-27-7	ALDICADD
	000000 54 4	ALE I UP RENURE	000107-02-0	ACKULEIN	000118-06-3	NI-M-OCTY! DUTUALATE
	000000000	CHLURUSENZUIRIFLUURIUE, 4-	000107-30-2	ACONIONITORIE	000117-04-0	BIS(2-ETHYLHEXYL) PHTHALATE /
	000096-07-7	SENZULKICHLUKIDE	000107-13-1	ALLYL CHIODIDE	000117-01-7	PIS(2-EINICHEAIL) PHINALAIC /
	000099-55-8	MEINTL-D-NIIKOANILINE, 2-	000107-03-1	ALLIL CHLOKIDE	000119.75-2	CHICRANTI
	000099-59-2	METHOXY-5-NITROANILINE, 2-	000107-00-2	DICHEURUETRANE, 1,2"	000110-73-2	TOTALTRATOLLICAE 2 / 4-
	000099-35-4	TRINITROBENZENE, 1,3,3-	000107-18-8	ALLYL ALCOHOL	000110-70-7	USTATIROTOLUERE, 2,4,0"
	000099-65-0	DINITROBENZENE, 1,3-	000107-21-1	EINTLENE GLYCOL	000110-74-1	MEXACILURUSENZENE
	000099-08-1	NITROTOLUENE, M-	000107-15-3	ETHYLENE DIAMINE	000119-90-4	DIMETHOXIBERZIDIRE, 3,3'-
	000099-99-0	NITROTOLUENE, P-	000107-12-0	PROPIONITRILE	000119-93-7	DIMETHTLEENZIDINE, 3,3'-
	000099-09-2	NITROANILINE, M-	000107-98-2	PROPYLENE GLYCOL MONOMETHYL ETHER	000120-12-7	ANTHRACENE
	000100-52-7	BENZALDEHYDE	000107-16-4	FORMALDEHYDE CYANOHYDRIN	000120-36-5	DICHLORPROP
	000100-51-6	BENZYL ALCOHOL	000107-20-0	CHLOROACETALDEHYDE	000120-83-2	DICHLOROPHENOL, 2,4-
	000100-42-5	STYRENE	000108-10-1	METHYL ISOBUTYL KETONE	000120-82-1	TRICHLOROBENZENE, 1,2,4-
	000100-00-5	CHLORONITROBENZENE, P-	000108-38-3	XYLENE, M-	000120-61-6	DIMETHYLTEREPHTHALATE
	000100-21-0	TEREPHTHALIC ACID	000108-87-2	VINYL-1-CYCLOHEXENE, 4- DINITROBENZENE, 1,4- BENZYL CHLORIDE ETHYL BENZENE NICOTINONITRILE PHTHALIC ACID, P- NITROANILINE, P- BROMOPHENYL PHENYL ETHER, 4- METHYLENE-BIS(2-CHLOROANILINE), 4,4'- METHYLENE-BIS(2-CHLOROANILINE), 4,4'- METHYLENE-BIS(N,N'-DIMETHYL)ANILIN E, 4,4'- AZOBENZENE ETHYLANILINE, N- CAPROLACTAM DIMETHYLPHENOL, 2,4- CHLOROPHENOL, 4- DIBROMOBENZENE, 1,4- DIBROMOBENZENE, 1,4- DIBROMOBENZENE, 1,4- DIBROMOETHANE, 1,2- CHLOROANILINE, 4- CHLOROTOLUENE, P- CRESOL, P- / (4-METHYLPHENOL) EPICHLOROHYDRIN DICHLOROBENZENE, 1,4- BUTADIENE, 1,3- ETHOXYETHANOL DODECANOATE, 2- XYLENE, P- PHENYLENEDIAMINE, P- TOLUIDINE, P- ETHOXYETHANOL ACRYLATE, 2- ACROLEIN CHLOROMETHYL METHYL ETHER ACRYLONITRILE ALLYL CHLORIDE DICHLOROETHANE, 1,2- ALLYL ALCOHOL ETHYLENE GLYCOL ETHYLENE GLYCOL ETHYLENE GLYCOL ETHYLENE GLYCOL MONOMETHYL ETHER FORMALDEHYDE CYANOHYDRIN CHLOROACETALDEHYDE METHYL ISOBUTYL KETONE XYLENE, M- METHYLCYCLOHEXANE	000121-75-5	MALATHION

## CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY CAS REGISTRY NUMBER) continued

	000121-69-7	DIMETHYLANTI INF N.N-	000218-01-9	CHRYSENE	000583-78-8	DICHLOROPHENOL, 2,5-
	000121-82-4	PDX / (CYCLONITE)	000298-02-2	CHRYSENE PHORATE DISULFOTON METHYL PARATHION RONNEL HYDRAZINE ANTIMONY POTASSIUM TARTRATE ALDRIN DIETHYL-P-NITROPHENYL PHOSPHATE HEXACHLOROCYCLOHEXANE, ALPHA- HEXACHLOROCYCLOHEXANE, BETA- HEXACHLOROCYCLOHEXANE, DELTA- DINITROPHENOL, 2,5- LINURON DIAZINON CYANOGEN TRINITROPHENYLMETHYLNITRAMINE TOLLEREDIAMINE 3 4-	000586-11-8	DINITROPHENOL, 3,5-
	000121-73-3	CHICOCHITECHE M-	000298-04-4	DISULFOTON	000591-27-5	AMINOPHENOL, M-
	000121-01-5	DUTUALIC ACID M.	000278-00-0	METHYL PARATHION	000591-35-5	DICHLOROPHENOL, 3,5-
	000121-71-3	DINITEDIO DE 2 A	0.00270	POWNEI	000591-78-6	HEXANONE, 2-
	000121-14-2	DINITROTOLOGNE, 2,4	000277-04-3	HYDDATINE	000592-01-8	CALCIUM CYANIDE
	000122-34-9	SIRALINE N N.	0-13-408000	ANTIMOMY DOTACCIIM TARTRATE	000593-60-2	BROMOETHENE / (VINYL BROMIDE)
	000122-39-4	DIPHENTLAMINE, N.M.	000304-01-0	ALADIN	000575 20 2	DICHLOROPROPANE, 2,2-
	000122-00-7	VIPHENTENTUKAZINE, 1,2	000309-00-2	NIETUVI DANITRODUENVI DUOCDUATE	000574 26 7	DIMETHYLUREA, N.N-
	000123-30-8	ARINOPHERUL, P-	000311-43-3	DIETHIC PARTICULAR ALBUA.	000578-31-2	BROHOACETONE
	000123-73-9	CROTONALDENTDE	000319-04-0	HEXACHLOROCYCLOHEXANE, ALPHA	000576 51 2	TRICHLOROPROPANE, 1,1,2-
	000123-31-9	HYDROQUINONE	000319-83-7	HEXACHLOROCYCLOHEXANE, DELTA-	000390-77-0	DINITROTOLUENE, 2,3-
	000123-33-1	MALEIC HYDRAZIDE	000319-50-5	HEXACHLUROCTCLUMEXAME, DELIA-	000002-01-7	DINITROTOLUENE, 2,6-
	000123-91-1	DIOXANE, 1,4-	000329-71-5	DINITROPHENOL, 2,5-	000000-20-2	PENTACHLOROBENZENE
	000123-63-7	PARALDEHYDE	000330-55-2	LINURON	000000-73-3	HEXACHLOROCYCLOHEXANE-TECHNICAL
	000124-48-1	DIBROMOCHLOROMETHANE	000333-41-5	DIAZINON	000000-73-1	
	000124-09-4	HEXAMETHYLENE DIAMINE	000460-19-5	CYANOGEN	000009-19-0	TRICHLOROPHENOL, 3,4,5-
	000126-99-8	CHLORO-1,3-BUTADIENE, 2- / (CHLOROPRENE)	000479-45-8	TRINITROPHENYLMETHYLNITRAMINE	000609-93-8	DINITRO-P-CRESOL, 2,6-
		(CHLOROPRENE)	000496-72-0	TOLUENEDIAMINE, 3,4-	000610-39-9	DINITROTOLUENE, 3,4-
	000126-98-7	METHACRYLONITRILE	000504-24-5	AMINOPYRIDINE, 4-	000611-14-5	ETHYLTOLUENE, O-
	000127-18-4	TETRACHLOROETHYLENE	000506-68-3	CYANOGEN BROMIDE	000615-54-3	TRIBROMOBENZENE, 1,2,4-
Þ	000129-00-0	PYRENE	000506-64-9	TOLUENEDIAMINE, 3,4- AMINOPYRIDINE, 4- CYANOGEN BROMIDE SILVER CYANIDE	000617-84-5	DIETHYLFORMAMIDE
느	000130-15-4	NAPHTHOQUINONE, 1,4-	000506-61-6	POTASSIUM SILVER CYANIDE	000619-15-8	DINITROTOLUENE, 2,5-
7	000131-11-3	DIMETHYLPHTHALATE	000506-77-4	SILVER CYANIDE POTASSIUM SILVER CYANIDE CHLORINE CYANIDE BUTYLCHLORIDE, T- CHLOROBENZILATE TRIMETHYL PHOSPHATE DIMETHYLPHENOL, 2,3- DINITROBENZENE, 1,2- FURILM	000620-14-4	ETHYLTOLUENE, M-
	000132-64-9	DIBENZOFURAN	000507-20-0	BUTYLCHLORIDE, T-	000621-64-7	NITROSO-DI-N-PROPYLAMINE, N-
	000133-06-2	CAPTAN	000510-15-6	CHLOROBENZILATE	000622-96-8	ETHYLTOLUENE, P-
	000133-07-3	FOLPET	000512-56-1	TRIMETHYL PHOSPHATE	000624-83-9	METHYL ISOCYANATE
	000137-26-8	THIRAM	000526-75-0	DIMETHYLPHENOL, 2,3-	000630-05-0	CARBON MONOXIDE
	000139-40-2	PROPAZINE	000528-29-0	DINITROBENZENE, 1,2-	000630-10-4	
	000140-57-8	ARAMITE	000531-82-8		000630-20-6	TETRACHLOROETHANE, 1,1,1,2-
	000140-88-5	ETHYL ACRYLATE	000532-28-5	BENZALDEHYDE CYANOHYDRIN	000634-93-5	TRICHLOROANILINE, 2,4,6-
	000141-78-6	FTHYL ACETATE	000534-52-1	DINITRO-O-CRESOL, 4,6-	000636-21-5	METHYLANILINE HYDROCHLORIDE, 2-
	000142-82-5	HEPTANE. N-	000540-73-8	DIMETHYLHYDRAZINE, 1,2-		MITROSO-N-METHYLUREA, N-
	000142 32 3	DICHLOPOPROPANE 1 3-	000540-59-0	DICHLOROETHYLENE, 1,2- (MIXED	000695-77-2	TETRACHLOROCYCLOPENTADIENE
	000142 23 7	CHLORO-1,3-BUTADIENE, 2- / (CHLOROPRENE) METHACRYLONITRILE TETRACHLOROETHYLENE PYRENE MAPHTHOQUINONE, 1,4- DIMETHYLPHTHALATE DIBENZOFURAN CAPTAN FOLPET THIRAM PROPAZINE ARAMITE ETHYL ACRYLATE ETHYL ACRYLATE ETHYL ACETATE HEPTANE, N- DICHLOROPROPANE, 1,3- SODIUM CYANIDE ENDOTHALL		I SOMERS)	000759-73-9	NITROSO-N-ETHYLUREA, N-
	000145-73-3	EMPOTHALI	000541-73-1	DICHLOROBENZENE, 1,3-	000759-94-4	EPTC
	000145 15 5	SODIUM DIETHYLDITHIOCARBAMATE	000542-92-7	CYCLOPENTADIENE BARIUM CYANIDE	000764-41-0	DICHLORO-2-BUTENE, 1,4-
	000150-50-5		000542-62-1	RAPILM CYANIDE	000765-34-4	GLYCIDALDEHYDE
		POTASSIUM CYANIDE	000542-88-1	BIS(CHLOROMETHYL) ETHER	000823-40-5	TOLUENE-2,6-DIAMINE
		OCTAMETHYLPYROPHOSPHORAMIDE	000542-75-6	DICHLOROPROPENE, 1,3- / (TELONE	000834-12-8	AMETRYN
		DICHLOROETHYLENE, 1,2-C-			000001 44 T	NITROSO-DI-N-BUTYLAMINE, N-
		· · ·	0005//-02-3	COODED CYANINE	000930-55-2	NITROSOPYRROLIDINE, N-
	000156-10-5	NITROSODIPHENYLAMINE, P-	000344-72-3	NICYEL CYANINE	000933-78-8	TRICHLOROPHENOL, 2,3,5-
	000156-60-5	DICKLORUEINTLENE, 1,2-1-	000557-19-7	TING CVANIDE	000933-75-5	TRICHLOROPHENOL, 2,3,6-
	000193-39-5	INDENU[1,2,3-CD]PYKENE	000557-48 A	TUALITIM / IL ACETATE	000735 77 5	ETHYL-O-XYLENE, 4-
	000205-99-2	BENZO (B) FLUORANTHENE	000577 54 6	DINITEONIEMO 2 4-	000734 30 3	TETRACHLOROPHENOL, 2,3,5,6-
	000206-44-0	FLUORANTHENE	000574 34 3	DINIIKUTENUL, 2,0°	000733 73-3	MEPHOSFOLAN
	000207-08-9	BENZO (K) FLUORANTHENE	000574-74-9	DIUNCHURUPHENUL, 6,3°	000750 10 7	TETRACHLOROVINPHOS / (STIROPHOS)
	000208-96-8	NITROSODIPHENYLAMINE, P- DICHLOROETHYLENE, 1,2-T- INDENO(1,2,3-CD)PYRENE BENZO(B)FLUORANTHENE FLUORANTHENE BENZO(K)FLUORANTHENE ACENAPTHYLENE	UUUD/6-26-1	II) COPPER CYANIDE NICKEL CYANIDE ZINC CYANIDE THALLIUM (I) ACETATE DINITROPHENOL, 2,6- DICHLOROPHENOL, 2,3- DIMETHYLPHENOL, 2,6-	000701 11"7	tribulation of the man

CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE

### CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE (LISTED BY CAS REGISTRY NUMBER) continued

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031512-74-0 BUSAN 77
032534-81-9 PENTABROMODIPHENYL ETHER
032536-52-0 OCTABROMODIPHENYL ETHER
033663-50-2 TRICHLOROANILINE HYDROCHLORIDE,
            2,4,6-
033820-53-0 ISOPROPALIN
036907-42-3 VANADIUM SULFATE
039638-32-9 BIS(2-CHLOROISOPROPYL) ETHER
040487-42-1 PENDIMETHALIN
041851-50-7 CHLOROCYCLOPENTADIENE
051218-45-2 METOLACHLOR
059756-60-4 FLURIDONE
060238-56-4 CHLORTHIOPHOS
068554-00-7 ETHOXYETHANOL PHOSPHATE, 2-
071753-42-9 TETRACHLOROHYDRAZOBENZENE
077323-84-3 TRICHLOROCYCLOPENTADIENE
NO CASRN
            SULFUR OXIDES
NO CASRN
            BROMINATED DIBENZO-P-DIOXINS
NO CASRN
            BROMOCHLOROETHANES
NO CASRN
            BROMINATED DIBENZOFURANS
NO CASRN
            TRIMETHYLBENZENES
NO CASRN
            PARTICULATE MATTER
NO CASRN
            TIN AND COMPOUNDS
            LEAD ALKYLS
NO CASRN
NO CASRN
             NITROPHENOLS
NO CASRN
             NITROGEN OXIDES
             NICKEL REFINERY DUST
NO CASRN
NO CASRN
             DICHLOROBUTENES
             PHOTOCHEMICAL OXIDANTS
NO CASRN
NO CASRN
             THALLIUM (IN SOLUBLE SALTS)
             URANIUM, SOLUBLE SALTS
NO CASRN
             TRINITROPHENOLS
NO CASRN
NO CASRN
             POLYBROMINATED BIPHENYLS
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#### APPENDIX A-IV

#### IV. EFFECT LEVEL DEFINITIONS

Adverse effect. A biochemical change, functional impairment, or pathologic lesion that either singly or in combination adversely affects the performance of the whole organism, or reduces an organism's ability to respond to an additional environmental challenge.

Frank-effect-level (FEL). The exposure level at which there are statistically or biologically significant increases in frequency or severity of severe effects between the exposed population and its appropriate control group. These severe effects produce an unmistakable adverse health effect (such as severe convulsions or death).

Lowest-observed-adverse-effect level (LOAEL). The lowest exposure level at which there are statistically or biologically significant increases in frequency or severity of adverse effects between the exposed population and its appropriate control group.

Lowest-observed-effect level (LOEL). The lowest exposure level at which there are statistically or biologically significant increases in frequency or severity of any effects between the exposed population and its appropriate control group. The effects that are seen at this level may or may not be considered as adverse.

No-observed-adverse-effect level (NOAEL). An exposure level at which there are no statistically or biologically significant increases in the frequency or severity of adverse effects between the exposed population and its appropriate control; some effects may be produced at this level, but they are not considered to be adverse.

No-observed-effect level (NOEL). An exposure level at which there are no statistically or biologically significant increases in the frequency or severity of any effect between the exposed population and its appropriate control.

Adapted from: U.S. EPA. 1991. Integrated Risk Information System (IRIS). Online. Office of Health and Environmental Assessment, Environmental Criteria and Assessment Office, Cincinnati, OH.

#### APPENDIX A-V

#### V. NATIONAL AMBIENT AIR QUALITY STANDARDS (NAAQS)

The Clean Air Act requires that National Ambient Air Quality Standards (NAAQS) be set and ultimately met for any air pollutant which, if present in air, may reasonably be anticipated to endanger public health or welfare and whose presence in the air results from numerous or diverse mobile and/or stationary sources. Since the primary NAAQS and the inhalation RfC serve essentially the same function, and the primary NAAQS have extensive data bases rigorously reviewed, the primary NAAQS with annual averaging times should be used *in lieu* of an inhalation RfC, except for lead. In deriving a risk assessment number for lead (Pb), the Integrated Exposure Uptake Biokinetics (IEUBK) model should be used instead of the RfC. Primary standards are designed to protect public health and secondary standards are designed to protect public health and secondary standards are designed to protect public welfare. Each primary NAAQS has either one or two averaging times depending on the health effects of the chemical. To date, six NAAQS have been established: Carbon Monoxide (CO), Lead (Pb), Nitrogen Dioxide (NO<sub>2</sub>), Particulate Matter, less than 10  $\mu$ m in size, (PM<sub>10</sub>), Ozone (O<sub>3</sub>) and Sulfur Dioxide (SO<sub>2</sub>). A table of the most recent NAAQS is provided as Table A-V-1.

The process of establishing and revising the NAAQS is detailed by Padgett and Richmond (Journal of the Air Pollution Control Association, 33:13-16, 1983). The primary NAAQS are solely health based and designed to protect the most sensitive group of individuals (but not necessarily the most sensitive members of that group) against adverse health effects. Thus, by definition, the NAAQS primary standards define allowable pollutant concentrations which can be present in the atmosphere without causing adverse effects, and essentially serve the same function as an inhalation RfC in a risk assessment/risk management decision, except for lead. The data bases supporting each of the NAAQS are extensive. More importantly, the NAAQS are set by the USEPA Administrator as mandated by Congress after numerous reviews and a public comment process

#### TABLE A-V-1

### NATIONAL AMBIENT AIR QUALITY STANDARDS<sup>a</sup> (as of December 2, 1991)

Pollutant	Primary Standards <sup>b</sup> Averaging Time		Secondary Standards <sup>b</sup>
Carbon monoxide (CO)	9 ppm (10 mg/m³) 35 ppm (40 mg/m³)	8 hour <sup>c</sup> 1 hour <sup>c</sup>	None
Lead (Pb) (and Lead compounds	1.5 <b>µ</b> g/m³	Quarterly	Same as primary
Nitrogen dioxide (NO <sub>2</sub> ) (Nitrogen oxide) (Nitric oxide)	0.053 ppm (100 <b>µ</b> g/m³)	Annual	Same as primary
Particulate Matter (PM <sub>10</sub> )	50 <b>µ</b> g/m³ 150 <b>µ</b> g/m³	Annual <sup>d</sup> 24 hours <sup>e</sup>	Same as primary
Ozone (O <sub>3</sub> )	0.12 ppm (235 <b>µ</b> g/m³)	1 hour <sup>f</sup>	Same as primary
Sulfur dioxide (SO <sub>2</sub> ) (Sulfur oxide)	0.03 ppm (80 <b>µ</b> g/m³)	Annual	
	0.14 ppm (365 <b>µ</b> g/m³)	24 hours <sup>c</sup>	
		3 hours <sup>c</sup>	0.5 ppm (1300 <b>µ</b> g/m³)

<sup>&</sup>lt;sup>a</sup>Source: U.S. EPA 1991. Subchapter C - Air Programs. Part 50 -National Primary and Secondary Ambient Air Quality Standards. Code of Federal Regulations 50: 693-697. Revised 7/1/91.

<sup>&</sup>lt;sup>b</sup>Primary standards are designed to protect public health; Secondary standards are designed to protect public welfare.

Not to be exceeded more than once per year.

<sup>&</sup>lt;sup>d</sup>The standard is attained when the expected annual arithmetic mean concentration is less than or equal to  $50 \mu g/m^3$ .

<sup>&</sup>lt;sup>e</sup>The standard is attained when the expected number of days per calendar year with a 24-hour average concentration above 150  $\mu$ g/m<sup>3</sup> is equal to or less than 1.

The standard is attained when the expected number of days per calendar year with maximum hourly average concentrations above 0.12 ppm is equal to or less than 1

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