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HEALTH EFFECTS ASSESSMENT

SUMMARY TABLES

FY-1997 Update

Office of Research and Development
Office of Emergency and Remedial Response
U.S. Environmental Protection Agency
Washington, DC 20460

HEALTH EFFECTS ASSESSMENT SUMMARY TABLES
FY-1997 UPDATE

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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 National Center for 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 an FY97 Update of the Health Effects Assessment Summary Tables (HEAST) prepared by EPA's National Center for Environmental Assessment, Cincinnati, OH (NCEA-CIN) for use at both Superfund and RCRA sites. It is intended to replace former editions and supplements of the HEAST. The HEAST will be updated annually if sufficient new data exist.

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. Although these entries in the HEAST have undergone review and have the concurrence of individual Agency Program Offices, and each is supported by an Agency reference, they have not 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. Until recently, IRIS evaluations were conducted by the Agency's Work Group Review process. To improve IRIS and to make it more useful, EPA requested and received public comment. As a consequence the Agency has initiated an IRIS Pilot program to replace the Reference Dose/Reference Concentration (RfD/RfC) and the Carcinogen Risk Assessment Verification Endeavor (CRAVE) Work Groups.

The Pilot will produce new or updated toxicological reviews and IRIS entries containing Agency consensus scientific positions on potential adverse human health effects that may result from chronic exposure to environmental contaminants.

The Pilot process consists of (1) a call for public involvement for interested parties to have some level of input into IRIS technical information, (2) a search of the relevant literature,

(3) development of toxicological reviews and draft IRIS summaries, (4) internal peer review within EPA, (5) external peer review by experts selected for each substance outside EPA, (6) consensus review and management approval within EPA, (7) preparation of final IRIS summaries and supporting documents, and (8) entry of summaries into the IRIS database.

Currently, the Pilot process, which has been underway since early FY 1996, is being applied to a select group of chemical substances chosen on the basis of the Agency's need for new or updated hazard or dose-response information. These assessments will be included in IRIS, and do not appear in the HEAST.

There are two exceptions to the above discussion. The HEAST also contains information on chemicals included under the National Ambient Air Quality Standards (NAAQS) and the Drinking Water Criteria Document (DWCD) series. In each of these cases, the chemicals are subject to extensive scientific peer review for quality assurance.

CHEMICAL STATUS DEFINITIONS

Chemicals previously reviewed by the Agency for consensus 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 earlier 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 receives Agency consensus as "verified" after all available information has been reviewed and a value has been calculated using current methodology. Verified values are entered on IRIS.

Some numbers that have achieved unanimous consensus by the previous Agency Work Groups may appear on the HEAST as "provisional" values.

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

Not verifiable: A toxicity value is "not verifiable" if all available data on a chemical was determined by the Agency to be 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 it is undergoing the Pilot process of considering all available data. All Pilot chemicals will have this status until the toxicity value is placed on IRIS.

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

Note: In all cases, the status of a chemical may change as new data become available, and the assessment is revisited, reviewed and verified through the Pilot Proces previously described.

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 earlier "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 Table 4. 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. 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 the RISK INFORMATION HOTLINE 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. Do not consult 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

Questions regarding the contents of the chemical toxicity and carcinogenicity tables on the HEAST (e.g., chemicals not covered, chemicals with pending [RfD]s) may be directed to

EPA's Superfund Health Risk Technical Support Center (TSC) in Cincinnati, OH at (513)569-7300 [FAX#: (513)569-7159]. Requests should include 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.

Written requests should be mailed to:

Superfund Health Risk Technical Support Center
US EPA
26 W. Martin Luther King Dr.
National Center for Environmental Assessment
MS - G44
Cincinnati, OH 45268

Radionuclide Carcinogenicity

Questions concerning radionuclide carcinogenicity should first be addressed by contacting the appropriate Regional Radiation Program Manager. A revised 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 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 (703)603-9230.

Drinking water documents are available by calling the Water Resource Center at (202)260-7786.

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 International Consultants, Inc. (513)569-7300 to be put on the mailing list. 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 Roy Smith of EPA's Office of Air Quality Planning and Standards at (919)541-5632.

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 1997 Health Effects Assessment Summary Table annual update from NTIS refer to the following order number:

PB97-921199: FY97 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 methodology practiced by the RfD/RfC Work Group.

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: 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 were not 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 to be reviewed and revised when necessary to reflect current information.

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: CARCINOGENICITY
Table 3 lists carcinogenicity values that were calculated by the CRAVE Work Group using Agency methodology.

HEAST TABLE 3 REFERENCES: CARCINOGENICITY
The references for Table 3 are numerically coded to associate each toxicity value clearly with its corresponding reference.

HEAST TABLE 4: RADIONUCLIDE CARCINOGENICITY - SLOPE FACTORS (In Units of Picocuries)
Table 4 lists ingestion, inhalation and external exposure carcinogenicity slope factors for radionuclides in units of picocuries and a factor to convert into the International System (SI) activity units of becquerels (Bq).

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

I. 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 FY97 ANNUAL HEAST

GENERAL CHANGES -- CHEMICAL TOXICITY AND CARCINOGENICITY

The changes in this version of the HEAST reflect changes in IRIS through July 1, 1997.

CHEMICAL-SPECIFIC CHANGES -- CHEMICAL TOXICITY AND CARCINOGENICITY

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

Benzo[b]fluoranthene 000205-99-2
Removed from Table 1 due to incomplete subchronic [RfC] assessment.

Bis(2-chloroisopropyl)ether 039638-32-9
Removed general comment from Table 1.

Chlorobenzene 000108-92-7
Removed from Table 1 the subchronic [RfD] comment due to incomplete assessment.

Dichloroethane, 1,2- 000107-06-2
Removed from Table 1 due to incomplete subchronic [RfC] and [RfD] assessments.

Manganese 007439-96-5
Removed the subchronic oral water [RfD] from Table 1 and citation 010850 from References to Table 1.

Trichloroethane, 1,1,1- 000071-55-6
Removed subchronic [RfC] comment from Table 1 due to incomplete assessment.

Uranium, Soluble Salts No CAS #
Removed from Table 1 due to incomplete assessment.

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

Dicyclopentadiene 000077-73-6
Changed target organ from liver to kidney.

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

Allyl chloride 000107-05-1
The general comment, "contact the Health Assessment Section" is removed from Table 3.

Arsenic, inorganic 007440-38-2
Removed inhalation [slope] factor value and comment from Table 3.

Benzo(b)fluoranthene 000205-99-2
Removed general comment.

Benzo(k)fluoranthene 000207-08-9
Removed the general comment, "contact the Health Assessment Section".

Chloromethyl methyl ether 000107-30-2
The general comment, "contact the Health Assessment Section" is removed.

Chrysene 000218-01-9
The general comment, "contact the Health Assessment Section" is removed.

Dibenzo[a,h]anthracene 000053-70-3
The general comment, "contact the Health Assessment Section" is removed.

Dichloroethane, 1,2- 000107-06-2
The inhalation [slope] factor and comment are removed from Table 3.

Dimethylbenz[a]anthracene, 7,12- 000057-97-6
Removed from Table 3. The general comment, "contact the Health Assessment Section" is removed from Table 3 References.

Methylcolanthracene, 3- 000056-49-5
Removed from Table 3 and from Table 3 References.

Nitroso-n-ethylurea, N- 000759-73-9
Removed the general comment, "contact the Health Assessment Section".

Polychlorinated biphenyls 001336-36-3
Added general comment: Carcinogenicity information was changed on IRIS.

D. CHEMICAL-SPECIFIC CHANGES ON HEAST TABLE 4: RADIONUCLIDE CARCINOGENICITY -- SLOPE FACTORS

No changes made to Table 4.

CHEMICAL SPECIFIC CHANGES MADE IN THE NOVEMBER 1995 SUPPLEMENT TO THE MAY 1995 HEAST ANNUAL UPDATE

The following changes were made in the November 1995 supplemental edition of the May 1995 HEAST Annual Update. Because some users may have been unaware of the publication of the November 1995 supplement, the following information should be noted.

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

Antimony trioxide 001309-64-4
An indicator was added to show that an inhalation RfC has been added to IRIS. The chronic inhalation RfC was adopted as the subchronic inhalation [RfC].

Boron, elemental 007440-42-8
The subchronic oral [RfD] was removed because the chronic oral RfD on which it was based is under review by the RfD/RfC Work Group.

Carbon disulfide 000075-15-0
An indicator was added to show that an inhalation RfC has been added to IRIS. The chronic inhalation RfC was adopted as the subchronic inhalation [RfC].

Hydrogen sulfide 007783-06-4
After a reevaluation of uncertainty factors by the RfD/RfC Work Group, the chronic inhalation RfC was modified to estimate the subchronic inhalation [RfC].

Mercuric chloride 007487-94-7
After a reevaluation of uncertainty factors by the RfD/RfC Work Group, The chronic oral RfD was modified to estimate the subchronic oral [RfD].

Phosphine 007803-51-2
An indicator was added to show that an inhalation RfC has been added to IRIS. The chronic inhalation RfC was modified to estimate the subchronic inhalation [RfC].

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

There were no changes to Table 2.

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

Arsenic, inorganic 007440-38-2
Indicators were added to show that an oral slope factor and an oral unit risk have been added to IRIS.

Bis(2-chloro-1-methylethyl) ether 000108-60-1
A typographical error in the CAS Registry Number has been corrected. There were no other changes to the record.

D. CHEMICAL-SPECIFIC CHANGES ON HEAST TABLE 4: RADIONUCLIDE CARCINOGENICITY - SLOPE FACTORS

For the November 1995 Supplement of the HEAST for radionuclides, EPA's Office of Radiation and Indoor Air (ORIA) has:

✓ corrected the factor in Table 4 for converting radionuclide slope factors from the customary units of picocuries (Ci) to the International System (SI) units of becquerels (Bq). (To convert radionuclides slope factors into the SI units of Bq, users should multiply each value in Table 4 by 27.03, not by 3.70E-02, the conversion factor provided in the May 1995 update.)

✓ added ingestion, inhalation, and external exposure slope factors for californium (Cf-252), iridium (Ir-192), thallium (Tl-207), and silver (Ag-110m+D).

✓ removed the ingestion, inhalation, and external slope factors for Cm-243+D and Pu-241+D. (EPA/ORIA re-evaluated the derivation and use of "+D" slope factors for decay chains that include a parent radionuclide (e.g., Cm-243 or Pu-241) with a radioactive half-life much shorter than the half-life of its immediate decay product (e.g., Pu-239 in the case of Cm-243 and Am-241 in the case of Pu-241). ORIA concluded that using "+D" slope factors for these types of radionuclides and decay chains may significantly underestimate radiation exposure and risk at certain sites, because such factors cannot be derived to cover all possible equilibrium conditions in the environment. At sites contaminated with these types of radionuclides, ORIA recommends that users (1) determine the radioactivity concentrations of the parent and each decay product radionuclides separately, (2) apply the appropriate slope factors in Table 4 for each radionuclide individually, and (3) add the individual risks from each radionuclide to calculate the collective risk posed by the site.)

✓ corrected the external slope factor values for Ac-227+D, Ce-144+D, Pu-244+D, Th-228+D, Th-229+D, and U-238+D in Table 4.

✓ corrected the branching factor for Ce-144 to Pr-144 from 9% to 98%, and corrected the half-life for Ra-228 from 8 years to 6 years in Exhibit 1.

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. 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 CARCINOGENICITY)

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):

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

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 Dose]
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
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).

FIGURE 1

Example Data and References for Chemical Toxicity

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

January 1992

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
GLYCIDALDEHYDE		000765-34-4								
NOAEL	10 PPM	RAT				1E-2	4E-3	1E-3	IRIS	005968
	INHALATION	12 WEEKS	WHOLE BODY		DECREASED WEIGHT GAIN	300	300	3000		
	INTERMITTENT		BLOOD		HEMATOPOIETIC EFFECTS					
			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 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, National Center for Environmental Assessment, CINCINNATI, OH FOR THE OFFICE OF SOLID WASTE AND EMERGENCY RESPONSE, WASHINGTON, DC

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 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/m^3), 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 m^3/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/m^3 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:

$$mg/L \text{ in water} = \frac{\text{oral [RfD] (in mg/kg/day)} \times 70 \text{ kg}}{2 \text{ L/day}}$$

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 were not 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 $\mu g/m^3$

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

[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^3/day) or the water consumption rate (2 L/day), respectively, for risk associated with unit concentration in air or water. Hence,

$$\text{risk per } \mu g/m^3 \text{ (air)} = (\text{risk per mg/kg/day}) \times \frac{1}{70\text{ kg}} \times 20\ m^3/\text{day} \times 10^{-3}\ (\text{mg}/\mu g)$$

$$\text{risk per } \mu g/L \text{ (water)} = (\text{risk per mg/kg/day}) \times \frac{1}{70\text{ kg}} \times 2\ L/\text{day} \times 10^{-3}\ (\text{mg}/\mu g)$$

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).

FIGURE 2
Example Data and References for Carcinogenicity

HEAST TABLE 3: CARCINOGENICITY

January 1992

CHEMICAL ROUTE	EXPERIMENT LENGTH	SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)$^{-1}$	REFERENCE
DIMETHYLHYDRAZINE,1,2- 000077-78-1										
ORAL DRINKING WATER	LIFETIME				B2	3.7E+1	3.7E+1	1.1E-3	1 1E-2	009993
		MOUSE	CARDIOVASCULAR SYSTEM	TUMORS						
Inhalation [Slope] Comment			BASED ON ROUTE TO ROUTE EXTRAPOLATION							

REFERENCES FOR HEAST TABLE 3: CARCINOGENICITY

January 1992

DIMETHYLHYDRAZINE, 1,2- 000077-78-1

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

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 risk] for air. Hence, the air concentration (in $\mu g/m^3$) corresponding to an upper-bound increased lifetime cancer risk of $1x10^{-5}$ 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 $1x10^{-5}$ is calculated as follows:

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

USER'S GUIDE: RADIONUCLIDE CARCINOGENICITY

Introduction

EPA classifies all radionuclides as Group A carcinogens. HEAST Table 4 lists ingestion, inhalation and external exposure cancer slope factors for radionuclides in units of picocuries (pCi).[1] Ingestion and inhalation slope factors are central estimates in a linear model of the age-averaged, lifetime attributable radiation cancer incidence (fatal and nonfatal cancer) risk per unit of activity inhaled or ingested, expressed as risk/pCi. External exposure slope factors are central estimates of lifetime attributable radiation 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 soil. When combined with site-specific media concentration data and appropriate exposure assumptions[2], slope factors can be used to estimate lifetime cancer risks to members of the general population due to radionuclide exposures.

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[1]Slope factors are reported in Table 4 in the customary units of picocuries (1 pCi = 10^{-12} curies (Ci) = 3.7×10^{-2} nuclear transformations per second) for consistency with the system used for radionuclides in the IRIS database. If required, slope factors in Table 4 can be converted into the International System (SI) units of becquerels (1 Bq = 1 nuclear transformation per second) by multiplying each inhalation, ingestion, or external exposure value by 27.03. 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.

[2]Agency standardized default exposure scenarios and assumptions for use in baseline risk assessment are provided in EPA (1991), *Risk Assessment Guidance for Superfund, Vol. I, 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 (703) 603-8917.

Radiation Effects

Ionizing radiation has been shown to be a carcinogen, a mutagen, and a teratogen. Radiation can induce cancers in nearly any tissue or organ in both humans and animals, and the probability of cancer induction increases with increasing radiation dose. Cancer induction is a delayed response that has 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 ORIA's approach and assumptions is provided in *Estimating Radiogenic Cancer Risks* (EPA 402-R-93-076).

Radionuclide slope factors are calculated for each radionuclide individually, based on its unique chemical, metabolic and radioactive properties. The calculation uses dose estimates from EPA's computer code RADRISK[3], vital statistics from the *U.S. Decennial Life Tables for 1979-1981* (described in EPA 402-R-93-076), and cancer risk estimates based largely on the results of the NAS BEIR V report[4], ICRP Publication 60[5],

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[3]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.

[4]National Academy of Sciences (1990). Health Effects of Exposure to Low Levels of Ionizing Radiation, BEIR V, Committee on the Biological Effects of Ionizing Radiations, National Research Council, Washington, D.C.

[5]International Commission on Radiological Protection (1991), 1990 Recommendations of the International Commission on Radiological Protection, ICRP Publication 60, Pergamon Press, New York, NY.

and U.S. Nuclear Regulatory Commission (NRC) analyses[6]. Ingestion and inhalation slope factors for radionuclides account for:

- 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 1980. (The expected lifetime for an individual in this population is about 74 years.) Consequently, radionuclide ingestion and inhalation slope factors are not expressed as a function of body weight and time, and do not require corrections for GI 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 Table 4 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.***

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[6]U.S. Nuclear Regulatory Commission (1991, 1993), Health Effects Models for Nuclear Power Plant Accident Consequence Analysis, NUREG/CR-4214. Addenda documenting the scientific basis for radiogenic risk models published in 1991 (for low-LET radiation) and 1993 (for alpha radiation). See EPA 402-R-93-076 for discussion of these models.

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.[7]

Because of the radiation risk models employed for both internal and external exposures, slope factors for radionuclides are characterized as central estimates in a linear model of the age-averaged lifetime total radiation cancer incidence risk per unit intake or exposure.

About the Information Provided in Table 4

Table 4 lists ingestion, inhalation and external exposure slope factors for principal radionuclides, and provides key parameter values used in the derivation of slope factor values. Radionuclides are presented alphabetically by element and atomic weight.

Selected radionuclides and radioactive decay chain products are designated in HEAST Table 4 with the suffix "+D" (e.g., U-238+D, Ra-226+D, Cs-137+D) to indicate that cancer risk estimates for these radionuclides include the contributions from their short-lived decay products, assuming equal activity concentrations (i.e., secular equilibrium) with the principal or parent nuclide in the environment.[8] Decay chains are identified in Exhibit 1.

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[7]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," ORNL-5974, Oak Ridge National Laboratory, Oak Ridge, TN.

[8]There is one exception to the assumption of secular equilibrium. For the inhalation slope factor for Rn-222+D reported in HEAST Table 4, ORIA assumes a 50% equilibrium value for radon decay products (Po-218, Pb-214, Bi-214 and Po-214) in air.

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 $2x10^{-6}$ (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 Managers, ORIA's National Air and Radiation Environmental Laboratory (NAREL) in

Montgomery, Alabama, ORIA's Las Vegas Laboratory (ORIA-LV) in Las Vegas, Nevada, or the ORIA contact at EPA headquarters in Washington, D.C., listed in Exhibit 2.

A Chemical Abstract System Reference Number (CASRN) is assigned to each radionuclide for identification and reporting accuracy during risk assessments, and radioactive half-lives are provided for reference.

The designations "D", "W", and "Y" presented in Table 4 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_1" 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 Table 4 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).

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 methodologies are refined, EPA will revise and update the slope factors in Table 4.

HEAST users with questions about radionuclide slope factor values and their use in radiation risk assessments should contact Michael Boyd of the Remedial Guidance Section of the Radiation Assessment Branch of ORIA at (202) 233-9395. Written requests for assistance can be sent by fax to (202) 233-9650.

Exhibit 1. Radionuclide Decay Chains Considered Explicitly in HEAST Table 4[9]

Principal Radionuclide [a]			Terminal Nuclide or Radionuclide [c]	
Nuclide	Half-life (yr)	Associated Decay Chain [b]	Nuclide	Half-life (yr)
Ac-227+D	22	[Th-227 (98.62%, 19 d)] Fr-223 (1.38%, 22 min) Ra-223 (11 d) Rn-219 (4 s) Po-215 (2 ms) Pb-211 (36 min) Bi-211 (2 min) [Tl-207 (99.72%, 5 min) Po-211 (0.28%, 0.5 s)]	Pb-207	*
Ag-108m+D	127	-[d] Ag-108 (8.90%, 2 min)	Pd-108 (91.1%) [Cd-108 (97.65%) Pd-108 (2.35%)]	* * *
Ag-110m+D	0.7	- Ag-110 (1.33%, 25 s)	Cd-110 (98.67%) [Cd-110 (99.7%) Pd-110 (0.3%)]	* * *
Am-243+D	7.4×10^3	Np-239 (2 d)	Pu-239	2.4×10^4
Ce-144+D	0.8	[Pr-144 (98.22%, 17 min) Pr-144m (1.78%, 7 min)]	Nd-144	*
Cs-137+D	30	Ba-137m (94.6%, 3 min)	Ba-137	*
Np-237+D	2.1×10^6	Pa-233 (27 d)	U-233	1.6×10^5
Pb-210+D	22	Bi-210 (5 d) Po-210 (138 d)	Pb-206	*
Pu-244+D	8.3×10^7	U-240 (14 h) Np-240m (7 4 min)	Pu-240	6.5×10^3
Ra-226+D	1.6×10^3	Rn-222 (4 d) Po-218 (3 min) [Pb-214 (99.98%, 27 min) At-218 (0.02%, 2 s)] Bi-214 (99.99%, 20 min) [Po-214 (99.98%, 1.64×10^{-4} s) Tl-210 (0.02%, 1 min)]	Pb-210	22
Ra-228+D	6	Ac-228 (6 h)	Th-228	2
Ru-106+D	1	Rh-106 (30 s)	Pd-106	*

[9]Source: International Commission on Radiological Protection (1983). Radionuclide Transformations: Energy and Intensity of Emission, ICRP Publication 38, Annals of the ICRP, Vols. 11-13, Pergamon Press, New York, NY.

Exhibit 1. Radionuclide Decay Chains Considered Explicitly in HEAST Table (Continued)

Principal Radionuclide [a]			Terminal Nuclide or Radionuclide [c]	
Nuclide	Half-life (yr)	Associated Decay Chain [b]	Nuclide	Half-life (yr)
Sb-125+D	3	Te-125m (22.8%, 58 d)	Te-125	*
Sr-90+D	29	Y-90 (64 h)	Zr-90	*
Th-228+D	2	Ra-224 (4 d) Rn-220 (56 s) Po-216 (0.2 s) Pb-212 (11 h) Bi-212 (61 min) [Po-212 (64.07%, 0.3 μs) Tl-208 (35.93%, 3 min)]	Pb-208	*
Th-229+D	7.3×10^3	Ra-225 (15 d) Ac-225 (10 d) Fr-221 (5 min) At-217 (32 ms) Bi-213 (46 min) [Po-213 (97.8%, 4 μs) Tl-209 (2.2%, 2 min)] Pd-209 (3 h)	Bi-209	*
U-235+D	7.0×10^8	Th-231 (26 h)	Pa-231	3.3×10^4
U-238+D	4.5×10^9	Th-234 (24 d) [Pa-234m (99.80%, 1 min) Pa-234 (0.33%, 7 h)]	U-234	2.4×10^5

(a) Radionuclides with half-lives greater than six months. "+D" designates principal radionuclides with associated decay chains.
(b) The chain of decay products of a principal radionuclide extending to (but not including) the next principal radionuclide or a stable radionuclide. Half-lives are given in parentheses. Branches are indicated by square brackets with branching percentages in parentheses.
(c) The principal radionuclide or stable nuclide that terminates an associated decay chain. Stable nuclides are indicated by an asterisk (*) in place of a half-life.
(d) A hyphen indicates that there are no associated decay products.

Exhibit 2. EPA Radition Program Managers

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CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ACENAPHTHENE		**000083-32-9**							
NOAEL 175 MG/KG/DAY		MOUSE							
	ORAL: GAVAGE	90 DAYS	LIVER	HEPATOTOXICITY		6E-1 300		IRIS	010165
ACENAPTHYLENE		**000208-96-8**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005202
ACEPHATE		**030560-19-1**							
LOAEL 2 PPM		RAT							
	ORAL DIET	13 WEEKS	BRAIN	DECREASED CHOLINESTERASE ACTIVITY		4E-3 30		IRIS	005833
	GENERAL COMMENT	ALSO SEE HEAST TABLE 3. CARCINOGENICITY							
ACETONE		**000067-64-1**							
NOEL 100 MG/KG/DAY		RAT							
	ORAL GAVAGE	90 DAYS	LIVER	INCREASED WEIGHT		1E+0		IRIS	005204
			KIDNEY	INCREASED WEIGHT		100			
			KIDNEY	NEPHROTOXICITY					
ACETONE CYANOHYDRIN / (2-METHYLLACTONITRILE)		**000075-86-5**							
NOAEL 8 75 MG/(KG-DAY)		RAT							
	ORAL GAVAGE	90 DAYS	LIVER	INCREASED RELATIVE WEIGHT		8E-3 300		8E-4 3000	005776

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)

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ACETONITRILE		**000075-05-8**							
NOAEL 100 PPM		MOUSE							
	INHALATION·	92 DAYS	ERYTHROCYTES	DECREASED CELL COUNT		6E-2		IRIS	005210
	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							
	ORAL· DIET	17 WEEKS		NONE OBSERVED		1E+0		IRIS	005212
						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							
	ORAL. WATER	90 DAYS						2E-2	010390
								1000	
GENERAL COMMENT		ALSO SEE HEAST TABLE 3 CARCINOGENICITY							
							IRIS		010856

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ACRYLAMIDE		**000079-06-1**								
NOEL 0.2 MG/KG/DAY	ORAL. DRINKING WATER	RAT	90 DAYS	NERVE	DAMAGE		2E-3 100		IRIS	005835
GENERAL COMMENT. ALSO SEE HEAST TABLE 3· CARCINOGENICITY								IRIS		010876
CHRONIC RfC COMMENT· THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (09/20/90) BY THE RfD/RfC WORK GROUP										
ACRYLIC ACID		**000079-10-7**								
NOAEL 53 MG/KG/DAY	ORAL DRINKING WATER	RAT	2 GENERATION	WHOLE BODY	DECREASED PUP WEIGHT		5E-1 100		IRIS	005836
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD ON IRIS WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]										
LOAEL 5 PPM	INHALATION· INTERMITTENT	MOUSE	13 WEEKS	NASAL MUCOSA	LESIONS	3E-3 100		IRIS		010346
ACRYLONITRILE		**000107-13-1**								
NOAEL 1 MG/(KG-DAY)	ORAL. GAVAGE	MOUSE	60 DAYS	TESTES	DECREASED SPERM COUNTS		1E-2		1E-3	010939
				TESTES	SEMINIFEROUS TUBULE DEGENERATION		100		1000	
SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL [RfD] UNDER REVIEW BY THE RfD/RfC WORK GROUP WAS MODIFIED TO ESTIMATE THE SUBCHRONIC ORAL [RfD]										
GENERAL COMMENT. ALSO SEE HEAST TABLE 3 CARCINOGENICITY										
ADIPONITRILE		**000111-69-3**								
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005157

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ALACHLOR			**015972-60-8**							
NOEL 1 MG/KG/DAY		DOG								
	ORAL CAPSULE		1 YEAR	BLOOD SITES, MULTIPLE	ANEMIA HEMOSIDEROSIS		1E-2 100		IRIS	005837
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]										
GENERAL COMMENT: ALSO SEE HEAST TABLE 3 CARCINOGENICITY.										
ALDICARB			**000116-06-3**							
NOAEL 0.01 MG/KG-DAY		HUMAN								
	ORAL		ACUTE	CENTRAL NERVOUS SYSTEM	SWEATING		1E-3 10		IRIS	010960
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	LIVER	LESIONS		3E-5 1000		IRIS	005159
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										005838

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ALLYL ALCOHOL			**000107-18-6**							
NOEL	50 PPM	RAT								
	ORAL: DRINKING WATER		15 WEEKS	LIVER	EFFECTS		5E-2		IRIS	005839
				KIDNEY	EFFECTS		100			
ALLYL CHLORIDE			**000107-05-1**							
NOAEL	17 MG/CU M	RABBIT								
	INHALATION INTERMITTENT		5 MONTHS	NERVOUS SYSTEM	NEUROTOXICITY	1E-2 300		IRIS		010369
	GENERAL COMMENT: ALSO SEE HEAST TABLE 3· CARCINOGENICITY									
ALUMINUM			**007429-90-5**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005162
ALUMINUM PHOSPHIDE			**020859-73-8**							
NOAEL	0.43 MG/KG/DAY	RAT								
	ORAL: DIET		2 YEARS	WHOLE BODY	ALTERED WEIGHT		4E-4		IRIS	010255
				UNSPECIFIED	ALTERED CLINICAL PARAMETERS		100			
AMETRYN			**000834-12-8**							
NOEL	10 MG/KG/DAY	RAT								
	ORAL. GAVAGE		13 WEEKS	LIVER	EFFECTS		9E-2 100		IRIS	005841
AMINO-2-NAPHTHOL, 1-			**002834-92-6**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005842
AMINO-2-NAPHTOL HYDROCHLORIDE, 1-			**001198-27-2**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.								005843

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
AMINOPHENOL, M-		**000591-27-5**								
NOAEL 1300 PPM		RAT								
	ORAL: DIET		13 WEEKS	WHOLE BODY	ALTERED WEIGHT		7E-1		7E-2	005844
				THYROID	ALTERED WEIGHT		100		1000	
AMINOPHENOL, O-		**000095-55-6**								
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.								005845
AMINOPHENOL, P-		**000123-30-8**								
	GENERAL COMMENT·	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.								005846
AMINOPYRIDINE, 4-		**000504-24-5**								
NOAEL 3 PPM		RAT								
	ORAL: DIET		90 DAYS	LIVER	INCREASED WEIGHT		2E-4		2E-5	005847
				BRAIN	INCREASED WEIGHT		1000		10000	
AMMONIA		**007664-41-7**								
NOAEL 34 MG/L		HUMAN								
	ORAL DRINKING WATER			SENSORY	TASTE THRESHOLD		34 MG/L 1		34 MG/L 1	005166
	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: INTERMITTENT			NASAL CAVITY	RHINITIS	1E-1		IRIS		010392
				LUNGS	PNEUMONIA	30				
				LUNGS	LESIONS					

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ANILINE		**000062-53-3**								
NOAEL 19 MG/CU M		MOUSE								
	INHALATION: INTERMITTENT		20-26 WEEKS	SPLEEN	PATHOLOGY	1E-2 300		IRIS		010370
		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		90 DAYS		NONE OBSERVED		3E+0 300		IRIS	010166
CHRONIC [RfC] COMMENT THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (08/04/94) BY THE RfD/RfC WORK GROUP										010964
ANTIMONY PENTOXIDE		**001314-60-9**								
LOAEL 0 46 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER		LIFETIME	WHOLE BODY	INCREASED MORTALITY		5E-4		5E-4	005174
				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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ANTIMONY POTASSIUM TARTRATE 000304-61-0										
LOAEL 0.91 MG/KG/DAY		RAT								
	ORAL. DRINKING WATER		LIFETIME	WHOLE BODY	INCREASED MORTALITY		9E-4		9E-4	005234
				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										
LOAEL 0 44 MG/KG/DAY		RAT								
	ORAL DRINKING WATER		LIFETIME	WHOLE BODY	INCREASED MORTALITY		4E-4		4E-4	005238
				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								
	ORAL DRINKING WATER		LIFETIME	WHOLE BODY	INCREASED MORTALITY		4E-4		4E-4	005242
				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										
BMC 0.87 MG/CU M		RAT								
	INHALATION, INTERMITTENT		1 YEAR	LUNG	PULMONARY TOXICITY	2E-4		IRIS		010974
				LUNG	INTERSTITIAL INFLAMMATION, CHRONIC	30				
CHRONIC RfC COMMENT· A BENCHMARK DOSE APPROACH WAS USED RATHER THAN A NOAEL/LOAEL TO DERIVE THE RfC.										
SUBCHRONIC [RfC] COMMENT. THE CHRONIC INHALATION RfC IS ADOPTED AS THE SUBCHRONIC INHALATION [RfC]										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ANTIMONY, METALLIC		**007440-36-0**								
LOAEL 0 35 MG SB/KG/DAY		RAT								
	ORAL DRINKING WATER		LIFETIME	WHOLE BODY	INCREASED MORTALITY		4E-4		IRIS	005170
				BLOOD	ALTERED CHEMISTRIES		1000			
ARAMITE		**000140-57-8**								
NOAEL 100 PPM		RAT								
	ORAL DIET		104 WEEKS	LIVER	INCREASED WEIGHT				5E-2	005850
									100	
NOAEL 500 PPM		DOG								
	ORAL DIET		52 WEEKS	LIVER	DEGENERATION		1E-1			005849
							I00			

GENERAL COMMENT ALSO SEE HEAST TABLE 3 CARCINOGENICITY

AROCLOR 1248		**012672-29-6**								
									IRIS	
CHRONIC RfD COMMENT THE CHRONIC ORAL RfD IS CONSIDERED NOT VERIFIABLE (07/20/93) BY THE RfD/RfC WORK GROUP										010940
AROCLOR 1254		**011097-69-1**								
LOAEL 0 005 MG/KG/DAY										
	ORAL. CAPSULE		MONKEY >5 YEARS	IMMUNE SYSTEM	TOXICITY		5E-5 100		IRIS	010963

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ARSENIC, INORGANIC		**007440-38-2**							
NOAEL 0 009 MG/L		HUMAN							
	ORAL		SKIN	KERATOSIS		3E-4		IRIS	010434
			SKIN	HYPERPIGMENTATION		3			
SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
GENERAL COMMENT ALSO SEE HEAST TABLE 3. CARCINOGENICITY									
ATRAZINE		**001912-24-9**							
NOEL 3.5 MG/KG/DAY		RAT							
	ORAL DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		3.5E-2		IRIS	010855
						100			
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
GENERAL COMMENT ALSO SEE HEAST TABLE 3: CARCINOGENICITY.									
BARIUM		**007440-39-3**							
NOAEL 0.21 MG/KG/DAY		HUMAN							
	ORAL WATER	10 WEEKS	CARDIOVASCULAR SYSTEM	INCREASED BLOOD PRESSURE		7E-2		IRIS	010348
						3			
SUBCHRONIC [RfC] COMMENT ALSO SEE HEAST TABLE 2 ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
CHRONIC [RfC] COMMENT· ALSO SEE HEAST TABLE 2 ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BENEFIN			**001861-40-1**							
NOAEL 25 MG/KG/DAY		DOG								
	ORAL: DIET		1 YEAR	ERYTHROCYTE	DECREASED COUNT		3E-1 100		IRIS	005852
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
BENZAL CHLORIDE			**000098-87-3**							
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.										005853
BENZALDEHYDE			**000100-52-7**							
NOEL 200 MG/KG/DAY		RAT								
	ORAL· GAVAGE		13 WEEKS	KIDNEY	EFFECTS		1E+0		IRIS	005854
				FORESTOMACH	LESIONS		100			
SUBCHRONIC [RfD] CPMMENT THE CHRONIC ORAL RfD WAS MODIFIED TO ESTIMATE 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.										
BENZENETHIOL / (THIOPHENOL)			**000108-98-5**							
LOAEL 0.1 MG/(KG-DAY)		RAT								
	ORAL· GAVAGE		90 DAYS	LIVER	CENTRILOBULAR EOSINOPHILIC CHANGES		1E-4 1000		1E-5 10,000	010942
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL [RfD] UNDER REVIEW BY THE RfD/RfC WORKGROUP WAS MODIFIED TO ESTIMATE THE SUBCHRONIC ORAL [RfD]										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BENZIDINE		**000092-87-5**							
LOAEL 2 7 MG/KG/DAY		MOUSE							
	ORAL: DRINKING WATER	33 MONTHS	BRAIN	CELLULAR CHANGES		3E-3		IRIS	005830
			LIVER	CELLULAR CHANGES		1000			
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
GENERAL COMMENT ALSO SEE HEAST TABLE 3· CARCINOGENICITY									
							IRIS		010877
CHRONIC RfC COMMENT· THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (03/28/91) BY THE RfD/RfC WORK GROUP.									
BENZO[A]ANTHRACENE		**000056-55-3**							
CHRONIC [RfC] COMMENT THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (08/04/94) BY THE RfD/RfC WORK GROUP									010965
GENERAL COMMENT· ALSO SEE HEAST TABLE 3· CARCINOGENICITY									
BENZOIC ACID		**000065-85-0**							
NOAEL 312 MG/DAY		HUMAN							
	ORAL DIET			NONE OBSERVED		4E+0 1		IRIS	005260
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									
BENZYL ALCOHOL		**000100-51-6**							
LOAEL 286 MG/KG/DAY		RAT							
	ORAL GAVAGE	103 WEEKS	FORESTOMACH	EPITHELIAL HYPERPLASIA				3E-1 1000	005855
NOAEL 143 MG/KG/DAY		RAT							
	ORAL. GAVAGE	13 WEEKS	WHOLE BODY	DECREASED WEIGHT		1E+0 100			005856

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BERYLLIUM		**007440-41-7**								
NOAEL 0 54 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER		LIFETIME		NONE OBSERVED		5E-3 100		IRIS	005262
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
GENERAL COMMENT. ALSO SEE HEAST TABLE 3: CARCINOGENICITY										
BIPHENYL, 1,1'		**000092-52-4**								
NOAEL 50 MG/KG/DAY		RAT								
	ORAL· DIET		700 DAYS	KIDNEY	DAMAGE		5E-2 1000		IRIS	005857
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]										
								IRIS		010878
CHRONIC RfC COMMENT THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (09/20/90) BY THE RfD/RfC WORK GROUP										
BIS(2-CHLOROISOPROPYL) ETHER		**039638-32-9**								
NOAEL 35 8 MG/KG/DAY		MOUSE								
	ORAL DIET		2 YEARS	ERYTHROCYTES	DECREASED HEMOGLOBIN		4E-2 1000		IRIS	010257
BIS(2-ETHYLHEXYL) PHTHALATE / (DEHP)		**000117-81-7**								
									IRIS	010859
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										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BISPHENOL A		**000080-05-7**								
									IRIS	005268
NOAEL 750 PPM		RAT								
	ORAL		13 WEEKS	WHOLE BODY	DECREASED WEIGHT		6E-1 100			005266
BORON TRIFLUORIDE		**007637-07-2**								
NOAEL 6 MG/CU M		RAT								
	INHALATION INTERMITTENT		13 WEEKS	KIDNEY	NECROSIS	7E-3 300		7E-4 3000		010395
BORON, ELEMENTAL		**007440-42-8**								
NOAEL 8 8 MG/KG/DAY		DOG								
	ORAL: DIET		2 YEARS	TESTIS	LESIONS				IRIS	005272

SUBCHRONIC [RfD] COMMENT. THE SUBCHRONIC ORAL [RfD] WAS REMOVED BECAUSE THE CHRONIC ORAL RfD UPON WHICH IT WAS BASED IS UNDER REVIEW BY THE RfD/RfC WORK GROUP.
CHRONIC RfD COMMENT: THE CHRONIC ORAL RfD, WHILE STILL ON IRIS, IS BEING RECONSIDERED BY THE RfD/RfC WORK GROUP

LOAEL 4.5 MG/CU M		HUMAN								
	INHALATION· INTERMITTENT			RESPIRATORY TRACT BRONCHUS	IRRITATION BRONCHITIS	2E-2 100		2E-2 100		005269

SUBCHRONIC [RfC] COMMENT THE SUBCHRONIC INHALATION [RfC] IS SPECIFICALLY FOR ANHYDROUS BORAX
CHRONIC [RfC] COMMENT THE CHRONIC INHALATION [RfC] IS SPECIFICALLY FOR ANHYDROUS BORAX

BROMINATED DIBENZO-P-DIOXINS

GENERAL COMMENT DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. 005858

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BROMINATED DIBENZOFURANS										
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005859
BROMOACETONE		**000598-31-2**								
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.								005860
BROMOCHLOROETHANES										
	GENERAL COMMENT·	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005861
BROMODICHLOROMETHANE		**000075-27-4**								
LOAEL 17.9 MG/KG/DAY		MOUSE								
	ORAL GAVAGE		102 WEEKS	KIDNEY	CYTOMEGALY		2E-2 1000		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								

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BROMOFORM		**000075-25-2**								
NOEL 17.9 MG/KG/DAY		RAT								
	ORAL GAVAGE		13 WEEKS	LIVER	EFFECTS		2E-1 100		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
BROMOPHOS		**002104-96-3**								
NOAEL 5 MG/KG/DAY		RAT								
	ORAL DIET		3 GENERATIONS	BLOOD	DECREASED CHOLINESTERASE ACTIVITY		5E-2 100		5E-3 1000	005865
				LIVER	DECREASED CHOLINESTERASE ACTIVITY					
SUBCHRONIC [RfD] COMMENT. BASED ON A REPRODUCTION STUDY										
CHRONIC [RfD] COMMENT BASED ON A REPRODUCTION STUDY										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BROMOXYNIL			**001689-84-5**							
NOEL 5 MG/KG/DAY		RAT								
	ORAL: DIET		2 YEARS		NONE OBSERVED		2E-2 300		IRIS	005866
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
BROMOXYNIL OCTANOATE			**001689-99-2**							
NOEL 7.3 MG/KG/DAY		RAT								
	ORAL: DIET		2 YEARS		NONE OBSERVED		2E-2 300		IRIS	005867
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
BUSAN 77			**031512-74-0**							
	GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.									005868
BUSAN 90			**002491-38-5**							
	GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.									005869
BUTANOL, 1-			**000071-36-3**							
NOAEL 125 MG/KG/DAY		RAT								
	ORAL: GAVAGE		13 WEEKS	CENTRAL NERVOUS SYSTEM	HYPOACTIVITY		1E+0 100		IRIS	005870
				CENTRAL NERVOUS SYSTEM	ATAXIA					

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
BUTYL BENZYL PHTHALATE, N-		**000085-68-7**								
NOAEL 159 MG/KG/DAY		RAT								
	ORAL: DIET		26 WEEKS	LIVER	ALTERED WEIGHT		2E+0 100		IRIS	005616
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-2 100		IRIS	005871
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-2 300		3E-3 3000	005873
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	

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CALCIUM CYANIDE		**000592-01-8**							
NOAEL 19.1 MG/KG/DAY		RAT							
	ORAL: DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT		4E-2		IRIS	010258
			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].
CHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.

CAPROLACTAM		**000105-60-2**							
								IRIS	005284
NOAEL 50 MG/KG/DAY		RAT							
	ORAL: DIET	90 DAYS	KIDNEY	EFFECTS		5E-1			005282
						100			
							IRIS		010966
CAPTAFOL		**002425-06-1**							
LOAEL 2 MG/KG/DAY		DOG							
	ORAL: CAPSULE	12 MONTHS	KIDNEY	EFFECTS		2E-3		IRIS	005874
			BLADDER	EFFECTS		1000			

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CAPTAN			**000133-06-2**							
NOEL	12.5 MG/KG/DAY	RAT								
	ORAL: DIET			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.

CARBARYL			**000063-25-2**							
NOAEL	9.6 MG/KG/DAY	RAT								
	ORAL: DIET		2 YEARS	KIDNEY LIVER	TOXICITY TOXICITY		1E-1 100		IRIS	005876

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.

CARBOFURAN			**001563-66-2**							
NOEL	0.5 MG/KG/DAY	DOG								
	ORAL: DIET		1 YEAR	BLOOD TESTIS UTERUS	CHOLINESTERASE INHIBITION EFFECTS EFFECTS		5E-3 100		IRIS	005877

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CARBON DISULFIDE		**000075-15-0**							
NOEL	11 MG/KG/DAY	RABBIT							
	INHALATION: INTERMITTENT		FETUS	TOXICITY		1E-1 100		IRIS	010259
	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.								
BMC	19.7 MG/CU M	HUMAN OCCUPATIONAL							
	INHALATION: INTERMITTENT	12.1 +/- 6.9 YEARS	PERIPHERAL NERVOUS SYSTEM	DYSFUNCTION	7E-1 30		IRIS		010975
	CHRONIC RfC COMMENT: A BENCHMARK DOSE APPROACH WAS USED RATHER THAN A NOAEL/LOAEL TO DERIVE THE RfC.								
	SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RfC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].								
CARBON MONOXIDE		**000630-05-0**							
	CHRONIC [RfC] COMMENT: REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS.								010493
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 WATER	90 DAYS	LIVER	EFFECTS		2E-2 1000		IRIS	005290
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLORDANE		**000057-74-9**								
NOEL 0.055 MG/KG/DAY		RAT								
	ORAL: DIET		130 WEEKS	LIVER	HYPERTROPHY		6E-5 1000		IRIS	005296
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 THYROID NERVE	DECREASED WEIGHT EFFECTS MYELIN DEGENERATION		5E-2 500		IRIS	010261
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
CHLORO-1,3-BUTADIENE, 2- / (CHLOROPRENE)		**000126-99-8**								
NOAEL 32 PPM		RAT								
	INHALATION		90 DAYS	OLFACTORY EPITHELIUM	DEGENERATION	7E-2 30		7E-3 300		010515
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)										
CHLORO-M-CRESOL, P-		**000059-50-7**								
NOAEL 200 MG/KG/DAY		RAT								
	ORAL: GAVAGE		28 DAYS	WHOLE BODY	DECREASED WEIGHT GAIN		2E+0 100			005366
CHLOROACETALDEHYDE		**000107-20-0**								
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.										005342

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLOROACETIC ACID			**000079-11-8**							
LOAEL 30 MG/KG		RAT								
	ORAL: GAVAGE		13 WEEKS	HEART	MYOCARDITIS		2E-2 1000		2E-3 10000	005346
CHLOROANILINE, 2-			**000095-51-2**							
	GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005347
CHLOROANILINE, 3-			**000108-42-9**							
	GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005348
CHLOROANILINE, 4-			**000106-47-8**							
LOAEL 12.5 MG/KG/DAY		RAT								
	ORAL: DIET		78 WEEKS	SPLEEN	PROLIFERATIVE LESIONS		4E-3 3000		IRIS	005349
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
CHLOROBENZENE			**000108-90-7**							
									IRIS	010863
	SUBCHRONIC [RfC] 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).									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLOROBENZILATE		**000510-15-6**								
NOEL 5 MG/KG/DAY		RABBIT								
	ORAL: GAVAGE		13 DAYS	GASTRO-INTESTINAL SYSTEM	DECREASED STOOL QUANTITY		2E-2 300		IRIS	010260
				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								
	ORAL: DIET		5 MONTHS		NONE OBSERVED		2E+0 100		2E-1 1000	005360
CHLOROBENZOTRIFLUORIDE, 4-		**000098-56-6**								
NOAEL 15 MG/KG/DAY		RAT								
	ORAL: GAVAGE			KIDNEY	TUBULAR DEGENERATION		2E-1 100		2E-2 1000	005364

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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLOROBUTANE, 1-		**000109-69-3**							
NOAEL 43 MG/KG/DAY		RAT							
	ORAL: GAVAGE	103 WEEKS	WHOLE BODY	INCREASED MORTALITY				4E-1	005808
			CENTRAL NERVOUS SYSTEM	EFFECTS				100	
			BLOOD	HEMATOLOGIC EFFECTS					
NOAEL 86 MG/KG/DAY		RAT							
	ORAL: GAVAGE	13 WEEKS	WHOLE BODY	DECREASED WEIGHT GAIN		9E-1			005806
			CENTRAL NERVOUS SYSTEM	EFFECTS		100			
			SPLEEN	HEMATOPOIESIS					
CHLOROBUTANE, 2-		**000078-86-4**							
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.							005809
CHLOROCYCLOPENTADIENE		**041851-50-7**							
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005297
CHLOROFORM		**000067-66-3**							
LOAEL 12.9 MG/KG/DAY		DOG							
	ORAL: CAPSULE	7.5 YEARS	LIVER	LESIONS		1E-2		IRIS	005372
						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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLOROMETHANE / (METHYL CHLORIDE) 000074-87-3									
									010005
SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.									
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.									
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 WATER		REPRODUCTION	REPRODUCTIVE EFFECTS		5E-2 100		IRIS	010436
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: INTERMITTENT	4 WEEKS	LIVER	EFFECTS	1E+0 100		1E-1 1000		010444
CHLOROTOLUENE, M- 000108-41-8									
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.									005880

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLOROTOLUENE, O-		**000095-49-8**								
NOAEL 20 MG/KG/DAY		RAT								
	ORAL: GAVAGE	103 DAYS		WHOLE BODY	DECREASED WEIGHT GAIN		2E-1 100		IRIS	010167
CHLOROTOLUENE, P-		**000106-43-4**								
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.										010200
CHLORPYRIFOS		**002921-88-2**								
NOEL 0.03 MG/KG/DAY		HUMAN								
	ORAL: CAPSULE	20 DAYS OR 9 DAYS		BLOOD	DECREASED CHOLINESTERASE ACTIVITY		3E-3 10		IRIS	005881
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
CHLORPYRIFOS METHYL		**005598-13-0**								
NOAEL 1 MG/KG/DAY		RAT								
	ORAL: DIET	3 GENERATIONS		REPRODUCTION	DECREASED FERTILITY		1E-2 100		1E-2 100	005882
		RAT 2 YEARS		LIVER	EFFECTS					
CHLOROTHALONIL		**001897-45-6**								
NOEL 1.5 MG/KG/DAY		DOG								
	ORAL: DIET	2 YEARS		KIDNEY	EFFECTS		1.5E-2 100		IRIS	005883
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLORTHIOPHOS		**060238-56-4**							
NOAEL 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 [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
CHROMIUM(III)		**016065-83-1**							
NOEL 5 % (CR2O3)	ORAL: DIET	RAT 840 DAYS		NONE OBSERVED		1E+0 1000		IRIS	005731
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
CHRONIC [RfC] COMMENT: INHALATION ISSUES ARE UNDER REVIEW BY THE RFD/RFC WORK GROUP.									
SUBCHRONIC [RfC] COMMENT: CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER: (513) 569-7300.									
CHROMIUM(VI)		**018540-29-9**							
NOAEL 2.4 MG/KG/DAY	ORAL: DRINKING WATER	RAT 1 YEAR		NONE OBSERVED		2E-2 100		IRIS	005522
CHRONIC [RfC] COMMENT: INHALATION ISSUES ARE UNDER REVIEW BY THE RFD/RFC WORK GROUP.									
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
COPPER		**007440-50-8**								
LOAEL 5.3 MG		HUMAN								
	ORAL		SINGLE DOSE	GASTRO-INTESTINAL SYSTEM	IRRITATION		1.3 MG/L		1.3 MG/L	005374

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
				KIDNEY	HISTOPATHOLOGY		100			
				WHOLE BODY	DECREASED WEIGHT					
				ORGANS	DECREASED WEIGHT					

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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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
				NERVOUS SYSTEM	NEUROTOXICITY		100			
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								
	ORAL: GAVAGE		GESTATION DAYS 6-18	CENTRAL NERVOUS SYSTEM	HYPOACTIVITY		5E-3 1000		5E-3 1000	010516
				RESPIRATORY SYSTEM	DISTRESS					
				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.										
								IRIS		010890
CHRONIC RfC COMMENT: THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (12/11/91) BY THE RfD/RfC WORK GROUP.										
CUMENE 000098-82-8										
NOAEL 154 MG/KG/DAY		RAT								
	ORAL: GAVAGE		194 DAYS	KIDNEY	INCREASED WEIGHT		4E-1 300		IRIS	005392
NOAEL 105.1 PPM		RAT								
	INHALATION: INTERMITTENT		4 WEEKS	CENTRAL NERVOUS SYSTEM	INVOLVEMENT	9E-2 1000		9E-3 10000		005908
				NOSE	IRRITATION					

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CYANAZINE		**021725-46-2**								
NOEL 0.625 MG/KG/DAY		DOG								
	ORAL: DIET		1 YEAR	WHOLE BODY	DECREASED WEIGHT		2E-3		2E-3	010411
				BLOOD	INCREASED PLATELET COUNT		300		300	
				BLOOD	ALTERED CLINICAL CHEMISTRY PARAMETERS					

SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].
CHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS WITHDRAWN FROM IRIS (07/01/92). UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

CYANIDE		**000057-12-5**								
NOAEL 10.8 MG/KG/DAY		RAT								
	ORAL: DIET		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								
	ORAL: DIET		2 YEARS	WHOLE BODY	DECREASED WEIGHT		4E-2		IRIS	010263
				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].

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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	005400
NOAEL 30 MG/KG/DAY		RAT							
	ORAL: DIET	90 DAYS	WHOLE BODY	DECREASED WEIGHT GAIN		3E-1			005398
			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)								

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DACTHAL		**001861-32-1**							
NOAEL 1	MG/KG/DAY	RAT							
	ORAL: DIET	2 YEARS	LUNG	EFFECTS		1E-2		IRIS	005888
			LIVER	EFFECTS		100			
			KIDNEY	EFFECTS					
			THYROID	EFFECTS					
			THYROID HORMONES	EFFECTS					
	SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								
DALAPON		**000075-99-0**							
NOEL	8.45 MG/KG/DAY	RAT							
	ORAL DIET	2 YEARS	KIDNEY	INCREASED RELATIVE WEIGHT		3E-2		IRIS	005889
						300			
	SUBCHRONIC [RfD] COMMENT: CALCULATED BY ANALOGY TO DALAPON SODIUM BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								
DDT		**000050-29-3**							
NOEL	0.05 MG/KG/DAY	RAT							
	ORAL. DIET	27 WEEKS	LIVER	LESIONS		5E-4		IRIS	005408
						100			
	SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD] GENERAL COMMENT· ALSO SEE HEAST TABLE 3. CARCINOGENICITY.								

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DECABROMODIPHENYL ETHER		**001163-19-5**								
NOEL 1 MG/KG/DAY		RAT								
	ORAL: DIET		2 YEARS	LIVER	INCREASED WEIGHT		1E-2 100		IRIS	005891
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
GENERAL COMMENT ALSO SEE HEAST TABLE 3: CARCINOGENICITY.										
DI-N-OCTYL PHTHALATE		**000117-84-0**								
LOAEL 175 MG/KG/DAY		RAT								
	ORAL: DIET		7-12 MONTHS	KIDNEY	INCREASED WEIGHT		2E-2		2E-2	010275
				LIVER	INCREASED WEIGHT		1000		1000	
				LIVER	INCREASED SGOT ACTIVITY					
				LIVER	INCREASED SGPT ACTIVITY					
DIAZINON		**000333-41-5**								
NOAEL 0.09 MG/KG/DAY		RAT								
	ORAL. DIET		35-42 DAYS	BLOOD	DECREASED CHOLINESTERASE ACTIVITY		9E-4 100		9E-4 100	005892
DIBENZOFURAN		**000132-64-9**								
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005409
DIBROMOBENZENE, 1,4-		**000106-37-6**								
NOAEL 10 MG/KG/DAY		RAT								
	ORAL· GAVAGE		45 OR 90 DAYS	LIVER	INCREASED RELATIVE WEIGHT		1E-1		IRIS	005893
				LIVER	ALTERED ENZYME ACTIVITIES		100			

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DIBROMOCHLOROMETHANE		**000124-48-1**								
NOEL 21 4 MG/KG/DAY		RAT								
	ORAL: GAVAGE		13 WEEKS	LIVER	LESIONS		2E-1 100		IRIS	005894
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY										
DIBROMOETHANE, 1,2-		**000106-93-4**								
LOAEL 88 PPB		HUMAN								
	INHALATION: INTERMITTENT			SPERM	EFFECTS	2E-3 100		2E-4 1000		010854
SUBCHRONIC [RfC] COMMENT THE CHRONIC INHALATION [RfC] WAS MODIFIED TO ESTIMATE THE SUBCHRONIC INHALATION [RfC] CHRONIC [RfC] COMMENT. UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE. GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY										
DIBUTYL PHTHALATE		**000084-74-2**								
NOAEL 125 MG/KG/DAY		RAT								
	ORAL DIET		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										
DICAMBA		**001918-00-9**								
NOAEL 3 MG/KG/DAY		RABBIT								
	ORAL GAVAGE		GESTATION DAYS 6-18	FETUS	DECREASED WEIGHT		3E-2 100		IRIS	010945
				FETUS	INCREASED POST-IMPLANTATION LOSSES					
				FETUS	INCREASED MORTALITY					
				DAM	DECREASED WEIGHT GAIN					
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DICHLOROBENZENE, 1,2-		**000095-50-1**							
								IRIS	010864
GENERAL COMMENT: ALSO SEE HEAST TABLE 2: ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
DICHLOROBENZENE, 1,3-		**000541-73-1**							
									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**							
NOAEL 75 MG/CU M		RAT							
	INHALATION. INTERMITTENT	MULTI-GENERATION	LIVER	INCREASED WEIGHT IN MALE PARENTS	2 5E+0 30		IRIS		010840
SUBCHRONIC [RfC] COMMENT. THE CHRONIC INHALATION RfC WAS MODIFIED TO ESTIMATE THE SUBCHRONIC INHALATION [RfC].									
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.									
DICHLOROBUTENES									
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005415
DICHLORODIFLUOROMETHANE		**000075-71-8**							
								IRIS	005498
NOAEL 90 MG/KG/DAY		DOG							
	ORAL: DIET	90 DAYS		NONE OBSERVED		9E-1 100			005496
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)									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DICHLOROETHANE, 1,1- 000075-34-3									
NOEL 115 MG/KG/DAY	INHALATION INTERMITTENT	RAT 13 WEEKS		NONE OBSERVED		1E+0 100		1E-1 1000	005790
DICHLOROETHYLENE, 1,1- 000075-35-4									
LOAEL 9 MG/KG/DAY	ORAL: DRINKING WATER	RAT 2 YEARS	LIVER	LESIONS		9E-3 1000		IRIS	005419
DICHLOROETHYLENE, 1,2- (MIXED ISOMERS) 000540-59-0									
LOAEL 50 PPM	ORAL· DRINKING WATER	RAT 2 YEARS	LIVER	LESIONS		9E-3 1000		9E-3 1000	010509

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.

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 [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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DICHLOROETHYLENE, 1,2-C- 000156-59-2										
NOAEL 32 MG/KG/DAY		RAT								
	ORAL: GAVAGE		90 DAYS	BLOOD	DECREASED HEMATOCRIT		1E-1		1E-2	005420
				BLOOD	DECREASED HEMOGLOBIN		300		3000	
CHRONIC [RfD] COMMENT: UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.										
DICHLOROETHYLENE, 1,2-T- 000156-60-5										
NOAEL 17 MG/KG/DAY		MOUSE								
	ORAL: DRINKING WATER		90 DAYS	BLOOD	INCREASED ALKALINE PHOSPHATASE		2E-1		IRIS	005895
							100			
DICHLOROPHENOL, 2,3- 000576-24-9										
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005315
DICHLOROPHENOL, 2,4- 000120-83-2										
NOEL 3 PPM		RAT								
	ORAL: DRINKING WATER		2 GENERATIONS	IMMUNE SYSTEM	ALTERED IMMUNE FUNCTION		3E-3		IRIS	005314
							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
DICHLOROPHENOL, 2,6- 000087-65-0										
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005317

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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	TOXICITY		1E-2		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 SYSTEM	HEMORRHAGE		8E-2		IRIS	005890
						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		RAT							
	INHALATION.	13 WEEKS	NASAL MUCOSA	HYPERPLASIA	1 3E-2		IRIS		005898
	INTERMITTENT				100				
	GENERAL COMMENT	ALSO SEE HEAST TABLE 3: CARCINOGENICITY.							

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DICHLOROPROPANE, 1,3-		**000142-28-9**							
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.									005899
DICHLOROPROPANE, 2,2-		**000594-20-7**							
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.									005900
DICHLOROPROPENE, 1,3- / (TELONE II)		**000542-75-6**							
NOEL 3 MG/KG/DAY		RAT							
	ORAL. DIET	90 DAYS	ORGANS	INCREASED WEIGHT		3E-3 1000		IRIS	005901
GENERAL COMMENT: ALSO SEE HEAST TABLE 3 CARCINOGENICITY.									
NOAEL 5 PPM		MOUSE							
	INHALATION:	2 YEARS	NASAL MUCOSA	HYPERTROPHY	2E-2		IRIS		010351
	INTERMITTENT		NASAL MUCOSA	HYPERPLASIA	30				
SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RfC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].									
DICHLORPROP		**000120-36-5**							
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.									005896
DICYCLOPENTADIENE		**000077-73-6**							
NOEL 32 MG/KG/DAY		RAT							
	ORAL: DIET	3 GENERATIONS		NONE OBSERVED		3E-1 100		3E-2 1000	005425

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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DIELDRIN		**000060-57-1**							
NOAEL 0.005 MG/KG/DAY	ORAL: DIET	RAT 2 YEARS	LIVER	LESIONS		5E-5 100		IRIS	005429
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
GENERAL COMMENT: ALSO SEE HEAST TABLE 3. CARCINOGENICITY.									
DIETHYL PHTHALATE		**000084-66-2**							
NOAEL 750 MG/KG/DAY	ORAL: DIET	RAT 16 WEEKS	WHOLE BODY ORGANS	DECREASED GROWTH DECREASED WEIGHT		8E+0 100		IRIS	005620
DIETHYL-P-NITROPHENYL PHOSPHATE		**000311-45-5**							
GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005922
DIETHYLANILINE, N,N-		**000091-66-7**							
GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005903
DIETHYLENE GLYCOL MONOBUTYL ETHER		**000112-34-5**							
NOAEL 18 PPM		RAT		NONE OBSERVED	2E-1 100		2E-2 1000		005482
CHRONIC [RfC] COMMENT: UNDER REVIEW									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DIETHYLENE GLYCOL MONOETHYL ETHER 000111-90-0									
NOEL	200 MG/KG/DAY	RAT							
	ORAL. DRINKING WATER		KIDNEY	HISTOPATHOLOGY				2E+0 100	005478
	CHRONIC [RfD] COMMENT. BASED ON A 3-GENERATION REPRODUCTION STUDY.								
NOEL	500 MG/KG/DAY	RAT							
	ORAL· DIET	90 DAYS	KIDNEY	IMPAIRED FUNCTION		5E+0			005476
			TESTIS	INCREASED WEIGHT		100			
DIETHYLFORMAMIDE 000617-84-5									
NOEL	0.546 MG/DAY, 5 DAYS/WEEK		RAT						
	ORAL GAVAGE	73 WEEKS		NONE OBSERVED		1.1E-2 100		1 1E-2 100	010437
DIETHYLHYDRAZINE, 1,2- 001615-80-1									
	GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ALSO SEE HEAST TABLE 3. CARCINOGENICITY								005921
DIMETHOATE 000060-51-5									
NOEL	0.05 MG/KG/DAY	RAT							
	ORAL. DIET	2 YEARS	BRAIN	DECREASED CHOLINESTERASE ACTIVITY		2E-4 300		IRIS	005923
	SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								
DIMETHYLANILINE, N,N- 000121-69-7									
LOAEL	22.32 MG/KG/DAY	MOUSE							
	ORAL· GAVAGE	13 WEEKS	SPLEEN	EFFECTS		2E-2 1000		IRIS	005924

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DIMETHYLFORMAMIDE, N,N- 000068-12-2										
NOAEL 96 MG/KG/DAY		RAT								
	ORAL· DIET		119 DAYS	LIVER	EFFECTS		1E+0 100		1E-1 1000	005925
LOAEL 22 MG/CU M		HUMAN								
	INHALATION: INTERMITTENT			LIVER GASTRO INTESTINAL SYSTEM	EFFECTS EFFECTS	3E-2 300		IRIS		010352
SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RFC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RFC].										
DIMETHYLPHENOL, 2,3- 000526-75-0										
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005926
DIMETHYLPHENOL, 2,4- 000105-67-9										
NOAEL 50 MG/KG/DAY		MOUSE								
	ORAL: GAVAGE		90 DAYS	NERVOUS SYSTEM BLOOD	EFFECTS ALTERATIONS		2E-1 300		IRIS	010266
DIMETHYLPHENOL, 2,5- 000095-87-4										
GENERAL COMMENT· DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005928
DIMETHYLPHENOL, 2,6- 000576-26-1										
NOEL 0 6 MG/KG/DAY		RAT								
	ORAL		8 MONTHS	WHOLE BODY ORGANS, MAJOR	INCREASED WEIGHT LESIONS		6E-3 100		IRIS	005431

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DIMETHYLPHENOL, 3,4-		**000095-65-8**							
NOEL 1.4 MG/KG/DAY	ORAL	RAT 8 MONTHS	WHOLE BODY	DECREASED WEIGHT		1E-2		IRIS	005437
			ORGANS, MAJOR	LESIONS		100			
			CARDIOVASCULAR SYSTEM	ALTERED BLOOD PRESSURE					
DIMETHYLPHTHALATE		**000131-11-3**							
CHRONIC RfD COMMENT THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (02/16/94) BY THE RfD/RfC WORK GROUP									010267
							IRIS		010894
CHRONIC RfC COMMENT: THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (07/26/90) BY THE RfD/RfC WORK GROUP.									
DIMETHYLTEREPHTHALATE		**000120-61-6**							
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]									
DIMETHYLUREA, N,N-		**000598-94-7**							
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005931
DINITRO-O-CRESOL, 4,6-		**000534-52-1**							
CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (02/11/93) BY THE RfD/RfC WORK GROUP.									010470
DINITRO-P-CRESOL, 2,6-		**000609-93-8**							
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005934

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DINITROBENZENE, 1,2-		**000528-29-0**								
NOAEL 0.4 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER		16 WEEKS	SPLEEN	INCREASED WEIGHT		4E-3 100		4E-4 1000	010201
SUBCHRONIC [RfD] COMMENT: DERIVED BY ANALOGY TO 1,3-DINITROBENZENE.										
CHRONIC [RfD] COMMENT DERIVED BY ANALOGY TO 1,3-DINITROBENZENE										
DINITROBENZENE, 1,3-		**000099-65-0**								
NOAEL 0.4 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER			SPLEEN	INCREASED WEIGHT		1E-3 100		IRIS	010471
DINITROBENZENE, 1,4-		**000100-25-4**								
NOAEL 0.4 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER		16 WEEKS	SPLEEN	INCREASED WEIGHT		4E-3 100		4E-4 1000	010202
SUBCHRONIC [RfD] COMMENT DERIVED BY ANALOGY TO 1,3-DINITROBENZENE										
CHRONIC [RfD] COMMENT· DERIVED BY ANALOGY TO 1,3-DINITROBENZENE.										
DINITROPHENOL, 2,3-		**000066-56-8**								
GENERAL COMMENT DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005936
DINITROPHENOL, 2,4-		**000051-28-5**								
LOAEL 2 MG/KG/DAY		HUMAN								
	ORAL			EYE	CATARACT		2E-3 1000		IRIS	010438
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										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DINITROPHENOL, 2,5-		**000329-71-5**							
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005937
DINITROPHENOL, 2,6-		**000573-56-8**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005938
DINITROPHENOL, 3,5-		**000586-11-8**							
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005939
DINITROTOLUENE, 2,3-		**000602-01-7**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005940
DINITROTOLUENE, 2,4-		**000121-14-2**							
NOAEL 0.2 MG/KG/DAY		DOG							
	ORAL GELATIN CAPSULE	UP TO 2 YEARS	CENTRAL NERVOUS SYSTEM	NEUROTOXICITY		2E-3 100		IRIS	005941
			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							
							IRIS		010896
	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**							
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							005942

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DINITROTOLUENE, 2,6-		**000606-20-2**								
NOAEL 4 MG/KG/DAY		DOG								
	ORAL: DIET		13 WEEKS	WHOLE BODY	MORTALITY		1E-2		1E-3	005943
				CENTRAL NERVOUS SYSTEM	NEUROTOXICITY		300		3000	
				BLOOD	HEINZ BODIES					
				BLOOD	METHEMOGLOBINEMIA					
				BILE DUCT	HYPERPLASIA					
				KIDNEY	HISTOPATHOLOGY					

CHRONIC [RfD] COMMENT: UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE
GENERAL COMMENT ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

DINITROTOLUENE, 3,4- **000610-39-9**

GENERAL COMMENT DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005944

DINOSEB **000088-85-7**

LOAEL 1 MG/KG/DAY		RAT								
	ORAL: DIET		29 WEEKS	FETUS	DECREASED WEIGHT		1E-3		IRIS	005945
							1000			

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 3-GENERATION REPRODUCTION STUDY.

DIPHENYLAMINE, N,N- **000122-39-4**

NOEL 2.5 MG/KG/DAY		DOG								
	ORAL· DIET		2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		2 5E-2		IRIS	005946
				LIVER	INCREASED WEIGHT		100			
				KIDNEY	INCREASED WEIGHT					

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DIRECT LIGHTFAST BLUE		**004399-55-7**							
GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005947
DISULFOTON		**000298-04-4**							
LOAEL 0.04 MG/KG/DAY		RAT							
	ORAL DIET	2 YEARS	EYE	DEGENERATION		4E-5		IRIS	010412
			BLOOD	CHOLINESTERASE INHIBITION		1000			
SUBCHRONIC [RfD] COMMENT·	THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]								
ENDOSULFAN		**000115-29-7**							
NOAEL 15 PPM		RAT							
	ORAL: DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		6E-3		IRIS	010926
			KIDNEY	GLOMERULONEPHROSIS		100			
			BLOOD VESSEL	ANEURYSMS					
NOAEL 10 PPM		DOG							
	ORAL: DIET	1 YEAR	WHOLE BODY	DECREASED WEIGHT GAIN					010938
SUBCHRONIC [RfD] COMMENT	THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHORNIC ORAL [RfD] BASED ON CO-CRITICAL RAT AND DOG STUDIES.								
ENDOTHALL		**000145-73-3**							
NOEL 2 MG/KG/DAY		DOG							
	ORAL DIET	2 YEARS	STOMACH	INCREASED WEIGHT		2E-2		IRIS	005948
			SMALL INTESTINE	INCREASED WEIGHT		100			
SUBCHRONIC [RfD] COMMENT·	THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ENDRIN		**000072-20-8**							
NOEL 0 025 MG/KG/DAY		DOG							
	ORAL: DIET	2 YEARS	CENTRAL NERVOUS SYSTEM	CONVULSIONS		3E-4 100		IRIS	005445
			LIVER	LESIONS					
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
EPICHLOROHYDRIN		**000106-89-8**							
LOAEL 37.8 MG/CU M		RAT							
	INHALATION: INTERMITTENT	136 WEEKS	KIDNEY	LESIONS		2E-3 1000		2E-3 1000	010440
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. INTERMITTENT	90 DAYS	NASAL EPITHELIUM	LESIONS			1E-2 100	IRIS	010492
EPTC		**000759-94-4**							
NOEL 2.5 MG/KG/DAY		RAT							
	ORAL DIET	2 GENERATIONS	HEART	DEGENERATIVE CARDIOMYOPATHY		2 5E-2 100		IRIS	005959
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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
ETHOXYETHANOL PHOSPHATE, 2- 068554-00-7									
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005955
ETHOXYETHANOL, 2- 000110-80-5									
LOAEL 357 MG/KG/DAY		RAT							
	ORAL: GAVAGE	103 WEEKS	WHOLE BODY	DECREASED WEIGHT				4E-1 1000	005470
NOEL 50 uL/KG/DAY		RAT							
	ORAL· GAVAGE	21 DAYS	FETUS	SKELETAL MALFORMATIONS		5E-1 100			005468
SUBCHRONIC [RfD] COMMENT BASED ON A REPRODUCTION STUDY WITH EXPOSURES DURING DAYS 1-21 OF GESTATION									
NOAEL 380 MG/CU M		RABBIT							
	INHALATION· INTERMITTENT	13 WEEKS	BLOOD	ALTERED HEMATOLOGY	2E+0 30		IRIS		010441
ETHOXYETHYL METHACRYLATE, 2- 002370-63-0									
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005954

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ETHYL ACETATE		**000141-78-6**								
NOEL	900 MG/KG/DAY	RAT								
	ORAL. GAVAGE		90 DAYS	WHOLE BODY	INCREASED MORTALITY		9E+0		IRIS	005957
				WHOLE BODY	DECREASED WEIGHT		100			
ETHYL BENZENE		**000100-41-4**								
									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.

ETHYL CHLORIDE		**000075-00-3**								
NOAEL	1504 PPM	MOUSE								
	INHALATION		10 DAYS	FETUS	DEVELOPMENTAL TOXICITY	1E+1		IRIS		010371
	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								
	ORAL GAVAGE		90 DAYS	LIVER	EFFECTS		2E+0		IRIS	010396
							300			

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ETHYL METHACRYLATE		**000097-63-2**								
NOEL 7 5 MG/KG/DAY	ORAL DRINKING WATER	RAT	2 YEARS	KIDNEY	INCREASED RELATIVE WEIGHT		9E-2 100		9E-2 100	005961
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**								
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.										010472
ETHYLANILINE, N-		**000103-69-5**								
GENERAL COMMENT· DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005958
ETHYLENE CYANOHYDRIN		**000109-78-4**								
NOEL 30 MG/KG/DAY	ORAL DRINKING WATER	RAT	90 DAYS	HEART	DECREASED WEIGHT		3E-1		3E-1	005780
				BRAIN	DECREASED WEIGHT		100		100	
ETHYLENE DIAMINE		**000107-15-3**								
NOAEL 22 6 MG/KG/DAY	ORAL DIET	RAT	3 MONTHS	HEART	DECREASED WEIGHT		2E-1		2E-2	005796
				BLOOD	HEMATOLOGIC CHANGES		100		1000	
								IRIS		010898
CHRONIC RfC COMMENT THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (12/18/90) BY THE RfD/RfC WORK GROUP.										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ETHYLENE GLYCOL		**000107-21-1**							
								IRIS	005454
NOEL	200 MG/KG/DAY	RAT							
	ORAL: DIET		FETUS	FETOTOXICITY		2E+0			005452
						100			
	SUBCHRONIC [RfD] COMMENT BASED ON A REPRODUCTION STUDY WITH EXPOSURES DURING DAYS 6-15 OF GESTATION.								
ETHYLENE GLYCOL MONOBUTYL ETHER		**000111-76-2**							
NOAEL	121 MG/CU M	RAT							
	INHALATION:	13 WEEKS	BLOOD	ALTERED HEMATOLOGY	2E-1		2E-2		010353
	INTERMITTENT				100		1000		
	CHRONIC [RfC] COMMENT UNDER REVIEW. CURRENT NUMBER SUBJECT TO CHANGE.								
ETHYLENE THIOUREA		**000096-45-7**							
LOAEL	0.25 MG/KG/DAY	RAT							
	ORAL: DIET	24 MONTHS	THYROID	HYPERPLASIA		8E-5		IRIS	010397
						3000			
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]								
	GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY								010899
	CHRONIC [RfC] COMMENT. THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (08/12/92) BY THE RfD/RfC WORK GROUP								
ETHYLTOLUENE, M-		**000620-14-4**							
	GENERAL COMMENT DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005963
ETHYLTOLUENE, O-		**000611-14-3**							
	GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005962

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ETHYLTOLUENE, P-		**000622-96-8**								
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005964
FLUORANTHENE		**000206-44-0**								
NOAEL 125 MG/KG/DAY		MOUSE								
	ORAL· GAVAGE		90 DAYS	KIDNEY	NEPHROPATHY		4E-1		IRIS	010168
				LIVER	WEIGHT CHANGES		300			
				BLOOD	HEMATOLOGICAL CHANGES					
CHRONIC [RfC] COMMENT THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (08/04/94) BY THE RfD/RfC WORK GROUP.										010967
FLUORENE		**000086-73-7**								
NOAEL 125 MG/KG/DAY		MOUSE								
	ORAL GAVAGE		13 WEEKS	ERYTHROCYTES	DECREASED COUNTS		4E-1		IRIS	010169
							300			
FLUORINE / (SOLUBLE FLUORIDE)		**007782-41-4**								
NOAEL 0 06 MG/KG/DAY		HUMAN								
	ORAL: DRINKING WATER			TOOTH	FLUOROSIS		6E-2		IRIS	005965
							1			
FLURIDONE		**059756-60-4**								
NOEL 200 PPM		RAT								
	ORAL: DIET		2 YEARS	KIDNEY	GLOMERULONEPHRITIS		8E-2		IRIS	005966
				TESTIS	ATROPHY		100			
				WHOLE BODY	DECREASED WEIGHT					
				ORGANS	DECREASED WEIGHT					
SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
FOLPET		**000133-07-3**								
NOEL	10 MG/KG/DAY	DOG								
	ORAL: CAPSULE		1 YEAR	WHOLE BODY	ALTERED WEIGHT GAIN		1E-1		IRIS	005967
				BLOOD	ALTERED CHEMISTRY		100			

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

FORMALDEHYDE		***000050-00-0***								
NOAEL	15 MG/KG/DAY	RAT								
	ORAL: WATER		2 YEARS	GASTRO-INTESTINAL TRACT	LESIONS		2E-1		IRIS	010398
							100			

GENERAL COMMENT ALSO SEE HEAST TABLE 3: CARCINOGENICITY.

FORMALDEHYDE CYANOHYDRIN		**000107-16-4**								
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005782

FORMIC ACID		**000064-18-6**								
NOAEL	200 MG/KG/DAY	RAT								
	ORAL WATER		MULTI-GENERATION	WHOLE BODY	DECREASED GROWTH		2E+0		2E+0	010268
							100		100	

CHRONIC [RfD] COMMENT BASED ON A MULTI-GENERATION STUDY. WITHDRAWN FROM IRIS (12/01/90) UNDER REVIEW.
CURRENT NUMBER SUBJECT TO CHANGE

FURAN		**000110-00-9**								
NOAEL	1.4 MG/KG/DAY	MOUSE								
	ORAL GAVAGE		13 WEEKS	LIVER	LESIONS		1E-2		IRIS	005462
							100			

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
FURFURAL		**000098-01-1**								
LOAEL 7 9 MG/KG/DAY		RAT								
	ORAL: GAVAGE		13 WEEKS	LIVER	HEPATOTOXICITY		3E-2 300		IRIS	005466
GLYCIDALDEHYDE		**000765-34-4**								
NOAEL 29 MG/CU M		RAT								
	INHALATION. INTERMITTENT		12 WEEKS	WHOLE BODY KIDNEY	DECREASED WEIGHT EFFECTS	1E-2 300	4E-3 300	1E-3 3000	IRIS	005968
HEPTACHLOR		**000076-44-8**								
NOEL 0.15 MG/KG/DAY		RAT								
	ORAL. DIET		2 YEARS	LIVER	INCREASED WEIGHT		5E-4 300		IRIS	005506
HEPTACHLOR EPOXIDE		**001024-57-3**								
LOAEL 0.0125 MG/KG/DAY		DOG								
	ORAL DIET		60 WEEKS	LIVER	INCREASED RELATIVE WEIGHT		1 3E-5 1000		IRIS	010399

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)

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.

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

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
HEPTANE, N-		**000142-82-5**								
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT										005969
HEXABROMOBENZENE		**000087-82-1**								
NOAEL 2 MG/KG/DAY		RAT								
	ORAL: DIET		12 WEEKS	LIVER	INDUCED CARBOXYLESTERASE ACTIVITY		2E-2 100		IRIS	005970
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.										
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 1000	010927
CHRONIC [RfD] COMMENT WITHDRAWN FROM IRIS (05/01/93) UNDER REVIEW. CURRENT NUMBER SUBJECT TO CHANGE.										
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
HEXACHLOROCYCLOHEXANE, EPSILON-		**006108-10-7**								
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT. ALSO SEE HEAST TABLE 3 CARCINOGENICITY.										010496

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
HEXACHLOROCYCLOHEXANE, GAMMA- 000058-89-9									
NOAEL 0.33 MG/KG/DAY		RAT							
	ORAL: DIET	12 WEEKS	LIVER	TOXICITY		3E-3		IRIS	005537
			KIDNEY	TOXICITY		100			
GENERAL COMMENT: ALSO SEE HEAST TABLE 3 CARCINOGENICITY.									010903
CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (05/27/92) BY THE RfD/RfC WORK GROUP.									
HEXACHLOROCYCLOPENTADIENE 000077-47-4									
NOAEL 7.1 MG/KG/DAY		RAT							
	ORAL	13 WEEKS	FORESTOMACH	LESIONS		7E-2		IRIS	005299
						100			
NOAEL 0.15 PPM		RAT							
	INHALATION:	13 WEEKS	NASAL CAVITY	SQUAMOUS METAPLASIA	7E-4		7E-5		010445
	INTERMITTENT				100		1000		
HEXACHLOROETHANE 000067-72-1									
NOAEL 1 MG/KG/DAY		RAT							
	ORAL: DIET	16 WEEKS	KIDNEY	DEGENERATION		1E-2		IRIS	005518
						100			
GENERAL COMMENT: ALSO SEE HEAST TABLE 3 CARCINOGENICITY.									010904
CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (11/05/92), BY THE RfD/RfC WORK GROUP.									
HEXACHLOROPHENE 000070-30-4									
LOAEL 0 75 MG/KG/DAY		DOG							
	ORAL: DIET	13 WEEKS	NERVOUS SYSTEM	EFFECTS		3E-3		IRIS	005972
						300			

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
HEXAMETHYLENE DIAMINE		**000124-09-4**							
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005973
HEXANE, N-		**000110-54-3**							
LOAEL 570 MG/KG/DAY		RAT							
	ORAL		NERVOUS SYSTEM	NEUROPATHY		6E-1		6E-2	005974
			TESTIS	ATROPHY		1000		10000	
LOAEL 73 MG/CU M		HUMAN							
	INHALATION		NERVOUS SYSTEM	NEUROTOXICITY	2E-1		IRIS		010273
	INTERMITTENT				300				
SUBCHRONIC [RfC] COMMENT· THE CHRONIC INHALATION RfC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC]									
HEXANONE, 2-		**000591-78-6**							
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005976
HYDROGEN SULFIDE		**007783-06-4**							
NOAEL 3 1 MG/KG/DAY		PIG							
	ORAL· FOOD	105 DAYS	GASTRO-INTESTINAL SYSTEM	DISTURBANCE		3E-2		IRIS	010269
						100			
NOAEL 42 MG/CU M		MOUSE							
	INHALATION:	13 WEEKS	NASAL MUCOSA	INFLAMMATION	1E-2		IRIS		010354
	INTERMITTENT				100				

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
HYDROQUINONE			**000123-31-9**							
NOAEL 4.29 MG/KG/DAY		HUMAN								
	ORAL		3-5 MONTHS	BLOOD	HEMATOLOGICAL EFFECTS		4E-1 10		4E-2 100	005526
								IRIS		010905
	CHRONIC RfC COMMENT	THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (10/01/90) BY THE RfD/RfC WORK GROUP								
IRON			**007439-89-6**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005527
ISOBUTYL ALCOHOL			**000078-83-1**							
NOEL 316 MG/KG/DAY		RAT								
	ORAL	GAVAGE	13 WEEKS	NERVOUS SYSTEM	HYPOACTIVITY		3E+0		IRIS	005977
				NERVOUS SYSTEM	ATAXIA		100			
ISOPHORONE			**000078-59-1**							
NOEL 150 MG/KG/DAY		DOG								
	ORAL	CAPSULE	90 DAYS	KIDNEY	LESIONS		2E+0 100		IRIS	005910
	GENERAL COMMENT	ALSO SEE HEAST TABLE 3: CARCINOGENICITY								
								IRIS		010906
	CHRONIC RfC COMMENT	THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (11/15/90) BY THE RfD/RfC WORK GROUP								
ISOPROPALIN			**033820-53-0**							
NOEL 15 MG/KG/DAY		RAT								
	ORAL	DIET	90 DAYS	BLOOD	HEMATOLIGICAL EFFECTS		1.5E-1		IRIS	005978
				ORGANS, UNSPECIFIED	ALTERED WEIGHTS		100			
LACTONITRILE			**000078-97-7**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005783

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
LEAD		**007439-92-1**							010447
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 ALSO SEE HEAST TABLE 3: CARCINOGENICITY.									
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, TRIETHYL LEAD, TRIMETHYL LEAD, TRIMETHYLETHYL LEAD; TRIPROPYL LEAD									
LINURON		**000330-55-2**							
LOAEL 0.625 MG/KG/DAY		DOG							
	ORAL: DIET	2 YEARS	BLOOD	HEMATOLOGICAL EFFECTS		2E-3 300		IRIS	005990
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
GENERAL COMMENT. ALSO SEE HEAST TABLE 3 CARCINOGENICITY									
MALATHION		**000121-75-5**							
NOEL 0 23 MG/KG/DAY		HUMAN							
	ORAL CAPSULE	47 DAYS	BLOOD	HEMATOLOGICAL EFFECTS		2E-2 10		IRIS	005991
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
CHRONIC [RfC] COMMENT· UNDER REVIEW									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
MALEIC ANHYDRIDE		**000108-31-6**							
NOAEL 10 MG/KG/DAY		RAT							
	ORAL: DIET	2 YEARS	KIDNEY	LESIONS		1E-1 100		IRIS	005992
SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
MALEIC HYDRAZIDE		**000123-33-1**							
LOAEL 500 MG/KG/DAY		RAT							
	ORAL DIET	28 MONTHS	KIDNEY	ALTERED FUNCTION		5E-1 1000		IRIS	005993
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
MALONONITRILE		**000109-77-3**							
LOAEL 0.21 MG/KG/DAY		RAT							
	ORAL· GAVAGE	120 DAYS	LIVER	EFFECTS		2E-4		2E-5	005994
			SPLEEN	EFFECTS		1000		10000	
MANCOZEB		**008018-01-7**							
NOEL 2 9 MG/KG/DAY		RAT							
	ORAL: DIET	90 WEEKS	THYROID	GOITROGENIC EFFECTS		3E-2 100		3E-2 100	005995
MANEB		**012427-38-2**							
NOEL 5 MG/KG/DAY		MONKEY							
	ORAL· DIET	6 MONTHS	THYROID	INCREASED WEIGHT		5E-2 100		IRIS	005996

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
MANGANESE		**007439-96-5**							
NOAEL 0.14 MG/KG/DAY		HUMAN							
	ORAL: DIET	CHRONIC	CENTRAL NERVOUS SYSTEM	EFFECTS		1 4E-1 1		IRIS	010851
SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL FOOD RfD WAS ADOPTED AS THE SUBCHRONIC ORAL FOOD [RfD]. SEE IRIS FOR SPECIFIC DIETARY INFORMATION.							IRIS		010959
SUBCHRONIC [RfC] COMMENT. A SUBCHRONIC [RfC] HAS NOT BEEN DERIVED FOR MANGANESE.									
MEPHOSFOLAN		**000950-10-7**							
NOEL 0.09 MG/KG/DAY		RAT							
	ORAL: DIET	17 WEEKS	LIVER	ALTERED WEIGHT		9E-4		9E-5	005997
			KIDNEY	ALTERED WEIGHT		100		1000	
			BLOOD	DECREASED CHOLINESTERASE ACTIVITY					
			ERYTHROCYTES	DECREASED CHOLINESTERASE ACTIVITY					
			BRAIN	DECREASED CHOLINESTERASE ACTIVITY					
MERCURIC CHLORIDE		**007487-94-7**							
		RAT							
	ORAL; SUBCUTANEOUS		IMMUNE SYSTEM	AUTOIMMUNE EFFECTS		3E-3 100		IRIS	005800
MERCURY, ELEMENTAL		**007439-97-6**							
NOAEL 0.009 MG/CU M		HUMAN							
	INHALATION.		NERVOUS SYSTEM	NEUROTOXICITY	3E-4 30		3E-4 30		010270
CHRONIC [RfC] COMMENT UNDER REVIEW. CURRENT NUMBER SUBJECT TO CHANGE.									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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					
										010907

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].

METHACRYLONITRILE		**000126-98-7**								
NOAEL 3 2 PPM		DOG								
	INHALATION		90 DAYS	LIVER	INCREASED SGOT		1E-3		IRIS	005812
	INTERMITTENT			LIVER	INCREASED SGPT		300			
				CENTRAL NERVOUS SYSTEM	LOSS OF HINDLIMB MOTOR CONTROL					
				BRAIN	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
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHANOL		**000067-56-1**							
NOEL	500 MG/KG/DAY	RAT							
	ORAL: GAVAGE	90 DAYS	BLOOD	INCREASED ALKALINE PHOSPHATASE		5E+0		IRIS	010271
			BLOOD	INCREASED SGPT		100			
			BRAIN	DECREASED WEIGHT					
METHOMYL		**016752-77-5**							
NOEL	2.5 MG/KG/DAY	DOG							
	ORAL: DIET	24 MONTHS	KIDNEY	LESIONS		2.5E-2		IRIS	005802
						100			
	SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								
METHOXYCHLOR		**000072-43-5**							
NOEL	5 01 MG/KG/DAY	RABBIT							
	ORAL: GAVAGE	13 DAYS	REPRODUCTION	LOSS OF LITTERS		5E-3		IRIS	010357
						1000			
	SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].								
							IRIS		010909
	CHRONIC RfC COMMENT: THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (11/07/91) BY THE RfD/RfC WORK GROUP								
METHOXYETHANOL ACETATE, 2-		**000110-49-6**							
									010497
	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)								

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHOXYETHANOL, 2-		**000109-86-4**							
NOAEL 93 MG/CU M		RABBIT							
	INHALATION:	13 WEEKS	TESTICLE	EFFECTS	2E-1		IRIS		010372
	INTERMITTENT				100				

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**							
NOEL 1156 MG/KG/DAY		RAT							
	ORAL: GAVAGE	90 DAYS	LIVER	INCREASED ALKALINE PHOSPHATASE		1E+1		1E+0	010002
			LIVER	INCREASED SGPT		100		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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHYL ETHYL KETONE		**000078-93-3**							
NOAEL 1711 MG/KG/DAY	ORAL DRINKING WATER	RAT MULTI-GENERATION	FETUS	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	INHALATION: INTERMITTENT	MOUSE 10 DAYS	FETUS	DECREASED BIRTH WEIGHT	1E+0 3000		IRIS		010845
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		8E-1		8E-2	010949
			LIVER	INCREASED RELATIVE WEIGHT IN FEMALES		300		3000	
			LIVER	INCREASED ABSOLUTE WEIGHT IN FEMALES					
			KIDNEY	INCREASED RELATIVE WEIGHT IN FEMALES					
			KIDNEY	INCREASED ABSOLUTE WEIGHT IN FEMALES					
			KIDNEY	INCREASED URINARY PROTEIN 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.									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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.										
METHYLMERCURY		**022967-92-6**								
CRITICAL ORAL DOSE		HUMAN								
0.001 MG/KG/DAY				DEVELOPMENTAL NEUROLOGICAL ABNORMALITIES IN HUMAN INFANTS			1E-4 10	IRIS		010970
SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD] CHRONIC [RfD] COMMENT. A BENCHMARK DOSE APPROACH WAS USED RATHER THAN A NOAEL/LOAEL TO DERIVE THE RfD.										
METHYL METHACRYLATE		**000080-62-6**								
NOEL 7 5 MG/KG/DAY		RAT								
	ORAL: WATER		24 MONTHS	KIDNEY	INCREASED RELATIVE WEIGHT		8E-2 100		8E-2 100	010014
METHYL PARATHION		**000298-00-0**								
									IRIS	010015
NOAEL 2.5 PPM		RAT								
	ORAL· DIET		90 DAYS	ERYTHROCYTES	CHOLINESTERASE INHIBITION		2E-3 100			010846
METHYL STYRENE (MIXED ISOMERS)		**025013-15-4**								
										010500
CHRONIC [RfD] COMMENT. UNDER REVIEW 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 STYRENE, ALPHA		**000098-83-9**								
										010499
GENERAL COMMENT· ALSO SEE HEAST TABLE 2 ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)										

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHYL-4-CHLOROPHENOXY) BUTYRIC ACID, 4-(2- 000094-81-5										
NOEL 12 MG/KG/DAY		RAT								
	ORAL: DIET		13 WEEKS	LIVER	EFFECTS		1E-1		IRIS	010008
				KIDNEY	EFFECTS		100			
		DOG								
	ORAL. DIET		13 WEEKS	LIVER	EFFECTS					
				KIDNEY	EFFECTS					
METHYL-4-CHLOROPHENOXY) PROPIONIC ACID, 2-(2- 000093-65-2										
NOEL 3 MG/KG/DAY		RAT								
	ORAL: DIET		90 DAYS	KIDNEY	ALTERED WEIGHT		1E-2		IRIS	010009
							300			
METHYL-4-CHLOROPHENOXYACETIC ACID, 2- 000094-74-6										
NOEL 0.15 MG/KG/DAY		DOG								
	ORAL: DIET		52 WEEKS	KIDNEY	EFFECTS		5E-4		IRIS	010007
				LIVER	EFFECTS		300			
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
METHYLCYCLOHEXANE 000108-87-2										
NOAEL 287 MG/CU M		RAT								
	INHALATION		1 YEAR	KIDNEY	MINERALIZATION	3E+0		3E+0		010431
	INTERMITTENT			KIDNEY	PAPILLARY HYPERPLASIA	100		100		
METHYLENE BROMIDE 000074-95-3										010501
	GENERAL COMMENT: ALSO SEE HEAST TABLE 2 ALTERNATE METHODS -- SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHYLENE CHLORIDE (DICHLOROMETHANE) 000075-09-2									
NOAEL 5 85 MG/KG/DAY		RAT							
	ORAL. DRINKING WATER	24 MONTHS	LIVER	TOXICITY		6E-2 100		IRIS	005553
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
NOAEL 694 8 MG/CU M		RAT							
	INHALATION: INTERMITTENT	2 YEARS	LIVER	TOXICITY	3E+0 100		3E+0 100		005552
GENERAL COMMENT ALSO SEE HEAST TABLE 3· CARCINOGENICITY.									
METHYLENE-BIS(2-CHLOROANILINE), 4,4'- 000101-14-4									
LOAEL 7.3 MG/KG/DAY		DOG							
	ORAL	9 YEARS	LIVER	EFFECTS		7E-4		7E-4	010413
			BLADDER	EFFECTS		10000		10000	
									010933
CHRONIC [RfC] COMMENT THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE 02/10/93) BY THE RfD/RfC WORK GROUP.									
METHYLENEDIPHENYL ISOCYANATE, 4,4- (DIPHENYLMETHANE DIISOCYANATE) 000101-68-8									
NOAEL 0.2 MG/CU M		RAT							
	INHALATION· INTERMITTENT	24 MONTHS	NASAL CAVITY	LESIONS	2E-5 300		IRIS		010449
SUBCHRONIC [RfC] COMMENT THE CHRONIC INHALATION RfC ON IRIS WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC].									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METOLACHLOR		**051218-45-2**							
NOAEL 300 PPM		RAT							
	ORAL· DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		1 5E-1 100		1.5E-1 100	010950

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

METRIBUZIN		**021087-64-9**							
NOAEL 100 PPM		DOG							
	ORAL: DIET	2 YEARS	LIVER	EFFECTS				IRIS	010928
			KIDNEY	EFFECTS					
			WHOLE BODY	MORTALITY					
			WHOLE BODY	DECREASED WEIGHT					

SUBCHRONIC [RfD] COMMENT· THE SUBCHRONIC ORAL [RfD] WAS REMOVED BECAUSE THE CHRONIC ORAL RfD UPON WHICH IT WAS BASED IS UNDER REVIEW BY THE RfD/RfC WORK GROUP.
CHRONIC RfD COMMENT. THE CHRONIC ORAL RfD, WHILE STILL ON IRIS, IS BEING RECONSIDERED BY THE RfD/RfC WORK GROUP

MIREX		**002385-85-5**							
NOAEL 0.07 MG/KG/DAY		RAT							
	ORAL DIET	2 YEARS	LIVER	CYTOMEGALY		2E-4		IRIS	010841
			LIVER	FATTY METAMORPHOSIS		300			
			LIVER	ANGIECTASIS					
			THYROID	CYSTIC FOLLICLES					

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
MOLINATE				**002212-67-1**						
NOEL 0 2 MG/KG/DAY		RAT								
	ORAL. GAVAGE			REPRODUCTIVE SYSTEM	TOXICITY		2E-3 100		IRIS	010017
SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].										
CHRONIC [RfD] COMMENT BASED ON A REPRODUCTION STUDY.										
MOLYBDENUM				**007439-98-7**						
LOAEL 0.14 MG/KG/DAY		HUMAN								
	ORAL: WATER, DIET			URINE	INCREASED URIC ACID		5E-3		IRIS	010489
				JOINTS	PAIN, SWELLING		30			
				BLOOD	DECREASED COPPER LEVELS					
SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RFD].										
MONOCHLORAMINE				**010599-90-3**						
NOAEL 9.5 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER		2 YEARS	WHOLE BODY	WEIGHT CHANGES		1E-1		IRIS	010517
				LIVER	WEIGHT CHANGES		100			
				KIDNEY	WEIGHT CHANGES					
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 CYANIDE				**000557-19-7**						
CHRONIC [RfD] COMMENT· THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/20/93) BY THE RfD/RfC WORK GROUP										010953

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
NICKEL, SOLUBLE SALTS		**VARIOUS**							
NOAEL 100 PPM		RAT							
	ORAL, DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT		2E-2		IRIS	005579
			ORGANS, MAJOR	DECREASED WEIGHT		300			

SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].
CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] WAS DERIVED FROM NICKEL MOIETY OF ADMINISTERED NICKEL CHLORIDE.

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
CHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD HAS BEEN PERMANENTLY WITHDRAWN (09/01/94) FROM IRIS.

NITRITE **014797-65-0**

NOEL 10 PPM		HUMAN							
	ORAL: WATER		BLOOD	METHEMOGLOBINEMIA		1E-1		IRIS	010021
						10			

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**

010936

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

LOAEL 9.8 MG/CU M		RAT							
	INHALATION:	4 WEEKS	BLOOD	HEMATOLOGICAL EFFECTS	2E-3		2E-4		010935
	INTERMITTENT				1000		10000		

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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
NITROBENZENE			**000098-95-3**							
LOAEL 25 MG/CU M		MOUSE								
	INHALATION·	90 DAYS		BLOOD	HEMATOLOGICAL EFFECTS		5E-3		IRIS	005589
	INTERMITTENT			ADRENAL	LESIONS		1000			
				KIDNEY	LESIONS					
				LIVER	LESIONS					
		RAT								
	INHALATION:	90 DAYS		BLOOD	HEMATOLOGICAL EFFECTS					
	INTERMITTENT			ADRENAL	LESIONS					
				KIDNEY	LESIONS					
				LIVER	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								
	CHRONIC [RfC] COMMENT:	ALSO SEE HEAST TABLE 2· ALTERNATE METHODS SUBCHRONIC AND CHRONICTOXICITY (OTHER THAN CARCINOGENICITY)								
	CHRONIC [RfD] COMMENT	THE ORAL RFD, WHILE STILL AVAILABLE ON IRIS, IS BEING RECONSIDERED BY THE RFD WORKGROUP. BASED ON ROUTE TO ROUTE EXTRAPOLATION								
	GENERAL COMMENT:	ALSO SEE HEAST TABLE 3: CARCINOGENICITY								
NITROFURANTOIN			**000067-20-9**							
NOAEL 300 PPM		MOUSE								
	ORAL· DIET	13 WEEKS		TESTIS	DAMAGE		7E-1		7E-2	005593
							100		1000	

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
NITROGEN DIOXIDE		**010102-44-0**							
CHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD HAS BEEN PERMANENTLY WITHDRAWN (09/01/94) FROM IRIS									010402
CHRONIC [RfC] COMMENT REFER TO APPENDIX A: TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS									010912
NITROGEN OXIDES									
CHRONIC [RfC] COMMENT· REFER TO APPENDIX A TECHNICAL INFORMATION, SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS									010170
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. INTERMITTENT	22 MONTHS	LIVER	LESIONS	2E-2 1000		IRIS		010374
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 1000		1E-2 10000	010029
SUBCHRONIC [RfD] COMMENT BASED ON DATA OBTAINED WITH O-NITROTOLUENE									
CHRONIC [RfD] COMMENT BASED ON DATA OBTAINED WITH O-NITROTOLUENE.									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
NITROTOLUENE, O-			**000088-72-2**							
LOAEL 200 MG/KG/DAY		RAT								
	ORAL. GAVAGE	6 MONTHS		SPLEEN	LESIONS		1E-1 1000		1E-2 10000	010028
NITROTOLUENE, P-			**000099-99-0**							
LOAEL 200 MG/KG/DAY		RAT								
	ORAL: GAVAGE	6 MONTHS		SPLEEN	LESIONS		1E-1 1000		1E-2 10000	010030
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		RAT								
	ORAL GAVAGE	90 DAYS		LIVER	HISTOLOGICAL CHANGES		3E-2 100		IRIS	010032
OCTAMETHYLPYROPHOSPHORAMIDE			**000152-16-9**							
NOAEL 0.02 MG/KG/DAY		HUMAN								
	ORAL	AT LEAST 30 DAYS		BLOOD	DECREASED CHOLINESTERASE ACTIVITY		2E-3 10		2E-3 10	010031
OSMIUM TETROXIDE			**020816-12-0**							
CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/22/93) BY THE RfD/RfC WORK GROUP.										010954
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PARATHION		**000056-38-2**							
NOAEL 0 064 MG/KG/DAY		HUMAN							
	ORAL		CHOLINESTERASE	DECREASED CHOLINESTERASE ACTIVITY		6E-3 10		6E-3 10	005598
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL [RfD] WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD] GENERAL COMMENT: ALSO SEE HEAST TABLE 3. CARCINOGENICITY.									
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							
	ORAL: DIET	SUBCHRONIC	BLOOD	INCREASED CLOTTING TIME		5E-2 100		5E-2 100	010036
PENDIMETHALIN		**040487-42-1**							
NOEL 12.5 MG/KG/DAY		DOG							
	ORAL: CAPSULE	2 YEARS	LIVER	EFFECTS		4E-2 300		IRIS	010037
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							
	ORAL GAVAGE	90 DAYS	LIVER	ALTERED ENZYME ACTIVITIES		2E-2 100		IRIS	010038

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PENTACHLOROBENZENE			**000608-93-5**							
LOAEL 8.3 MG/KG/DAY		RAT								
	ORAL· DIET		100 DAYS	LIVER	TOXICITY		8E-3		IRIS	010039
				KIDNEY	TOXICITY		1000			
PENTACHLOROCYCLOPENTADIENE			**025329-35-5**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005302
PENTACHLORONITROBENZENE			**000082-68-8**							
NOEL 0 75 MG/KG/DAY		DOG								
	ORAL: DIET		2 YEARS	LIVER	TOXICITY		3E-3		IRIS	010040
							300			
	SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
	GENERAL COMMENT	ALSO SEE HEAST TABLE 3. CARCINOGENICITY								
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PHENANTHRENE		**000085-01-8**							
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005604
CHRONIC [RfC] COMMENT· THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (08/04/94) BY THE RfD/RfC WORK GROUP.									
PHENOL		**000108-95-2**							
NOAEL 60 MG/KG/DAY		RAT							
	ORAL. GAVAGE		FETUS	DECREASED WEIGHT		6E-1 100		IRIS	005824
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							
	ORAL	90 DAYS	LIVER	LESIONS		6E-2 100		IRIS	010044
PHENYLENEDIAMINE, O-		**000095-54-5**							
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT ALSO SEE HEAST TABLE 3 CARCINOGENICITY.									010042
PHENYLENEDIAMINE, P-		**000106-50-3**							
NOAEL 18.7 MG/KG/DAY		RAT							
	ORAL: DIET	2 YEARS	WHOLE BODY	EFFECTS				1 9E-1 100	010043

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PHENYLMERCURIC ACETATE		**000062-38-4**								
NOAEL 0 0084 MG/KG/DAY		RAT								
	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		RAT								
	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

PHOSPHINE		**007803-51-2**								
NOEL 0 026 MG/KG/DAY		RAT								
	ORAL· DIET		2 YEARS				3E-4 100		IRIS	010174

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

NOAEL 1 4 MG/CU M		MOUSE								
	INHALATION: INTERMITTENT		13 WEEKS	WHOLE BODY	DECREASED WEIGHT	3E-3 100		IRIS		010976

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PHOSPHORUS, WHITE		**007723-14-0**							
								IRIS	010452
	GENERAL COMMENT:	FORMERLY LISTED AS PHOSPHORUS (INORGANIC COMPOUNDS).							
PHOTOCHEMICAL OXIDANTS									
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT.							010172
PHTHALIC ACID, M-		**000121-91-5**							
	GENERAL COMMENT:	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010047
PHTHALIC ACID, O-		**000088-99-3**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010046
PHTHALIC ACID, P-		**000100-21-0**							
NOEL	142 MG/KG/DAY	RAT							
	ORAL, DIET	2 YEARS	BLADDER	HYPERPLASIA		1E+0 100		1E+0 100	010048
PHTHALIC ANHYDRIDE		**000085-44-9**							
LOAEL	1562 MG/KG/DAY	MOUSE							
	ORAL, DIET	104 WEEKS	LUNG	HISTOPATHOLOGY		2E+0		IRIS	010049
			KIDNEY	HISTOPATHOLOGY		1000			
	SUBCHRONIC [RfD] COMMENT:	THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]							
LOAEL	0 1 MG/CU M	HUMAN							
	INHALATION	12 YEARS	NOSE	RHINITIS	1.2E-1		1 2E-1		010847
	INTERMITTENT		LUNGS	BRONCHITIS	300		300		
	SUBCHRONIC [RfC] COMMENT:	THE CHRONIC INHALATION [RfC] WAS ADOPTED AS THE SUBCHRONIC INHALATION [RfC]							

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
POLYBROMINATED BIPHENYLS									
LOAEL 0.07 MG/KG/DAY		RAT							
	ORAL. GAVAGE	25 WEEKS	LIVER	INCREASED WEIGHT		7E-5		7E-6	010050
			LIVER	LESIONS		1000		10000	
GENERAL COMMENT· ALSO SEE HEAST TABLE 3. CARCINOGENICITY.									
POTASSIUM CYANIDE		**000151-50-8**							
NOAEL 27 MG/KG/DAY		RAT							
	ORAL. DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT		5E-2		IRIS	010278
			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									
POTASSIUM SILVER CYANIDE		**000506-61-6**							
NOAEL 82 7 MG/KG/DAY		RAT							
	ORAL· DIET	2 YEARS	WHOLE BODY	DECREASED WEIGHT		2E-1		IRIS	010279
			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									
PROFLURALIN		**026399-36-0**							
NOEL 3 MG/KG/DAY		RAT							
	ORAL. DIET	SUBCHRONIC		NONE OBSERVED		6E-3		6E-3	010051
						500		500	

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PRONAMIDE			**023950-58-5**							
NOEL 7.5 MG/KG/DAY		DOG								
	ORAL: DIET		2 YEARS		NONE OBSERVED		7 5E-2 100		IRIS	010280
	SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
PROPACHLOR			**001918-16-7**							
NOEL 13.3 MG/KG/DAY		RAT								
	ORAL: DIET		90 DAYS	WHOLE BODY	DECREASED WEIGHT GAIN		1.3E-1 100		IRIS	010175
PROPAZINE			**000139-40-2**							
NOEL 5 MG/KG/DAY		RAT								
	ORAL DIET		2 YEARS	WHOLE BODY	DECREASED WEIGHT GAIN		2E-2 300		IRIS	010052
	SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
PROPIONITRILE			**000107-12-0**							
	GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									010053
PROPYL ALCOHOL, N-			**000071-23-8**							
	GENERAL COMMENT· DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005627

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PROPYLENE GLYCOL		**000057-55-6**								
NOEL	50000 PPM	DOG								
	ORAL. DIET		2 YEARS	ERYTHROCYTES	DECREASED COUNT				2E+1	005631
				BLOOD	DECREASED HEMATOCRIT				100	
				BLOOD	DECREASED HEMOGLOBIN					
NOEL	6 %	RAT								
	ORAL: DIET		20 WEEKS	KIDNEY	LESIONS		3E+1			005629
							100			
								IRIS		010914
	CHRONIC RfC COMMENT. THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (04/25/91) BY THE RfD/RfC WORK GROUP									
PROPYLENE GLYCOL MONOETHYL ETHER		**001569-02-4**								
NOEL	680 MG/KG/DAY	RAT								
	ORAL DRINKING WATER		30 DAYS	WHOLE BODY	DECREASED WEIGHT GAIN		7E+0		7E-1	005488
							100		1000	
								IRIS		010915
	CHRONIC RfC COMMENT. THE CHRONIC INHALATION RfC IS CONSIDERED NOT VERIFIABLE (04/25/91) BY THE RfD/RfC WORK GROUP.									
PROPYLENE GLYCOL MONOMETHYL ETHER		**000107-98-2**								
NOEL	947 MG/KG/DAY	RAT								
	ORAL GAVAGE		35 DAYS	LIVER	HISTOPATHOLOGY		7E+0		7E-1	005486
				KIDNEY	HISTOPATHOLOGY		100		1000	
NOAEL	1000 PPM	RAT, RABBIT								
	INHALATION. INTERMITTENT		13 WEEKS	CENTRAL NERVOUS SYSTEM	EFFECTS	2E+1 30		IRIS		010276

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
PROPYLENE OXIDE			**000075-56-9**							
LOAEL 71 MG/CU M		RAT								
	INHALATION:		2 YEARS	EPITHELIUM	UNSPECIFIED	3E-2		IRIS		010375
	INTERMITTENT					100				
SUBCHRONIC [RfC] COMMENT: THE CHRONIC INHALATION RfC WAS ADOPTED AS THE SUBCHRONIC INHALATION [RFC]										
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY										
PYRENE			**000129-00-0**							
NOAEL 75 MG/KG/DAY		MOUSE								
	ORAL: GAVAGE		13 WKS	KIDNEY	EFFECTS		3E-1		IRIS	010176
							300			
CHRONIC [RfC] COMMENT: THE CHRONIC INHALATION [RfC] IS CONSIDERED NOT VERIFIABLE (08/04/94) BY THE RfD/RfC WORK GROUP										010968
PYRIDINE			**000110-86-1**							
NOAEL 1 MG/KG/DAY		RAT								
	ORAL. GAVAGE		90 DAYS	LIVER	INCREASED WEIGHT		1E-2		IRIS	010055
				LIVER	INCREASED RELATIVE WEIGHT		100			
RDX / (CYCLONITE)			**000121-82-4**							
NOEL 0 3 MG/KG/DAY		RAT								
	ORAL		105 WEEKS	PROSTATE	INFLAMMATION		3E-3		IRIS	010056
				PROSTATE	HEMOSIDEROSIS		100			
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]										
GENERAL COMMENT: ALSO SEE HEAST TABLE 3 CARCINOGENICITY.										
RONNEL			**000299-84-3**							
NOAEL 5 MG/KG/DAY		RAT								
	ORAL DIET		2 YEARS	LIVER	EFFECTS		5E-2		5E-2	010057
							100		100	

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
SELENIOUS ACID		**007783-00-8**							
NOAEL 0.046 MG/KG/DAY		HUMAN							
	ORAL: DIET		WHOLE BODY	SELENOSIS, CLINICAL		5E-3 3		IRIS	010504
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
SELENIUM		**007782-49-2**							
NOAEL 0.853 MG/DAY		HUMAN							
	ORAL: DIET		WHOLE BODY	SELENOSIS, CLINICAL		5E-3 3		IRIS	010404
SUBCHRONIC [RfD] COMMENT· THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
SELENOUREA		**000630-10-4**							
NOAEL 0 072 MG/KG/DAY		HUMAN							
	ORAL· DIET		WHOLE BODY	SELENOSIS		5E-3 15		5E-3 15	010473
CHRONIC [RfD] COMMENT WITHDRAWN FROM IRIS (05/01/91) UNDER REVIEW, CURRENT NUMBER SUBJECT TO CHANGE.									
SILVER		**007440-22-4**							
LOAEL 0.014 MG/KG/DAY		HUMAN							
	IV	2-9 YEARS	SKIN	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									

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
SODIUM METAVANADATE		**013718-26-8**							
NOAEL 10 PPM		RAT							
	ORAL: DRINKING WATER	3 MONTHS	KIDNEY	IMPAIRED FUNCTION		1E-2 100		1E-3 1000	005735
STRONTIUM, STABLE		**007440-24-6**							
NOAEL 190 MG/KG/DAY		RAT, YOUNG							
	ORAL· DRINKING WATER	20 DAYS	BONE	RACHITIC CHANGES		6E-1 300		IRIS	010842
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 UNSPECIFIED	TOXICITY HISTOPATHOLOGY		3E-3 1000		IRIS	010285
GENERAL COMMENT THE LOAEL IS ALSO THE FEL.									
STYRENE		**000100-42-5**							
								IRIS	010059
NOAEL 22 PPM		HUMAN							
	INHALATION OCCUPATIONAL		CENTRAL NERVOUS SYSTEM	EFFECTS	3E+0 10		IRIS		010511
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.									
GENERAL COMMENT ALSO SEE HEAST TABLE 3 CARCINOGENICITY									
SUCCINONITRILE		**000110-61-2**							
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005585

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
SULFUR DIOXIDE		**007446-09-5**							
	CHRONIC [RfC] COMMENT	REFER TO APPENDIX A· TECHNICAL INFORMATION. SECTION V ON NATIONAL AMBIENT AIR QUALITY STANDARDS							010505
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 1000	010060
TERBUFOS		**013071-79-9**							
NOAEL 0 0025 MG/KG/DAY		DOG							
	ORAL DIET	6 MONTHS	CHOL1NESTERASE	INHIBITION		2 5E-5 100		2 5E-5 100	010408
TEREPHTHALIC ACID		**000100-21-0**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010474
TETRACHLOROAZOXYBENZENE		**021232-47-3**							
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010064

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TETRACHLOROBENZENE, 1,2,4,5- 000095-94-3										
NOAEL 0.34 MG/KG/DAY		RAT								
	ORAL DIET		13 WEEKS	KIDNEY	LESIONS		3E-3 100		IRIS	010286
TETRACHLOROCYCLOPENTADIENE 000695-77-2										
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005303
TETRACHLOROETHANE, 1,1,1,2- 000630-20-6										
LOAEL 89 3 MG/KG/DAY		RAT								
	ORAL GAVAGE		103 WEEKS	LIVER	LESIONS		3E-2		IRIS	010407
				KIDNEY	LESIONS		3000			
	SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
	GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.									
TETRACHLOROETHYLENE 000127-18-4										
NOAEL 14 MG/KG/DAY		MOUSE								
	ORAL		6 WEEKS	LIVER	HEPATOTOXICITY		1E-1 100		IRIS	005650
	GENERAL COMMENT· ALSO SEE HEAST TABLE 3· CARCINOGENICITY.									
TETRACHLOROHYDRAZOBENZENE 071753-42-9										
	GENERAL COMMENT	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								010065
TETRACHLOROPHENOL, 2,3,4,5- 004901-51-3										
	GENERAL COMMENT.	DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT								005324

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TETRACHLOROPHENOL, 2,3,4,6- 000058-90-2									
NOAEL 25 MG/KG/DAY		RAT							
	ORAL: GAVAGE	90 DAYS	LIVER	INCREASED WEIGHT		3E-1		IRIS	005323
			LIVER	CENTRILOBULAR HYPERTROPHY		100			
TETRACHLOROPHENOL, 2,3,5,6- 000935-95-5									
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005325
TETRACHLOROPROPENE, 1,1,2,3- 010436-39-2									
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									010066
TETRACHLOROVINPHOS / (STIROPHOS) 000961-11-5									
NOEL 3.1 MG/KG/DAY		DOG							
	ORAL: DIET	2 YEARS	LIVER	INCREASED WEIGHT		3E-2		IRIS	010067
			KIDNEY	INCREASED WEIGHT		100			
			WHOLE BODY						
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
GENERAL COMMENT: ALSO SEE HEAST TABLE 3: CARCINOGENICITY.									
TETRAETHYL DITHIOPYROPHOSPHATE 003689-24-5									
NOEL 0.5 MG/KG/DAY		RAT							
	ORAL: DIET	3 MONTHS	ERYTHROCYTES	DECREASED CHOLINESTERASE ACTIVITY		5E-3		IRIS	010287
						100			
			BLOOD	DECREASED CHOLINESTERASE ACTIVITY					
THALLIC OXIDE 001314-32-5									
CHRONIC [RfD] COMMENT: THE CHRONIC ORAL [RfD] IS CONSIDERED NOT VERIFIABLE (07/22/93) BY THE RfD/RfC WORK GROUP									010956

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
THALLIUM (I) ACETATE		**000563-68-8**							
NOAEL 0 26 MG/KG/DAY		RAT							
	ORAL	90 DAYS	LIVER	INCREASED SGOT		9E-4		IRIS	005664
			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 (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES.									
THALLIUM (I) CARBONATE		**006533-73-9**							
NOAEL 0 23 MG/KG/DAY		RAT							
	ORAL	90 DAYS	LIVER	INCREASED SGOT		8E-4		IRIS	005668
			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 (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES.									
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 (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES.									
CHRONIC [RfD] COMMENT. CALCULATED BY ANALOGY TO THALLIUM (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES									
THALLIUM, INSOLUBLE SALTS									
CHRONIC [RfD] COMMENT. REFER TO IRIS FOR OTHER THALLIUM SALTS.									010458

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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 (I) SULFATE BY CORRECTING FOR MOLECULAR WEIGHT DIFFERENCES

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

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					
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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
THIOFANOX		**013196-18-4**							
NOAEL 0 025 MG/KG/DAY		DOG							
	ORAL	8 DAYS	CHOLINESTERASE	DECREASED CHOLINESTERASE ACTIVITY		3E-4 100		3E-4 100	010069
SUBCHRONIC [RfD] COMMENT. BASED ON CONFIDENTIAL BUSINESS INFORMATION DATA.									
CHRONIC [RfD] COMMENT. BASED ON CONFIDENTIAL BUSINESS INFORMATION DATA									
THIRAM		**000137-26-8**							
								IRIS	010459
NOAEL 0.61 MG/KG/DAY		FERRET							
	ORAL	24 WEEKS	REPRODUCTION	IMPAIRED		6E-3 100			010070
TIN AND COMPOUNDS									
NOAEL 2000 PPM		RAT							
	ORAL: DIET	2 YEARS	LIVER KIDNEY	LESIONS LESIONS		6E-1 100		6E-1 100	005688
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									
TOLUENE		**000108-88-3**							
NOAEL 223 MG/KG/DAY		RAT							
	ORAL. GAVAGE	13 WEEKS	LIVER KIDNEY	ALTERED WEIGHT ALTERED WEIGHT		2E+0 100		IRIS	010469
							IRIS		010844
SUBCHRONIC [RfC] COMMENT CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER (513) 569-7300									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TOLUENE-2,5-DIAMINE		**000095-70-5**							
NOAEL 56 MG/KG/DAY		RAT							
	ORAL: DIET	78 WEEKS				6E-1		6E-1	010073
						100		100	
SUBCHRONIC [RfD] COMMENT: DETERMINED FROM DATA OBTAINED WITH THE SULFATE SALT.									
CHRONIC [RfD] COMMENT: DETERMINED FROM DATA OBTAINED WITH THE SULFATE SALT.									
TOLUENE-2,6-DIAMINE		**000823-40-5**							
NOAEL 16 MG/KG/DAY		RAT							
	ORAL DIET	2 YEARS				2E-1		2E-1	010074
						100		100	
SUBCHRONIC [RfD] COMMENT DETERMINED FROM DATA OBTAINED WITH THE DIHYDROCHLORIDE									
CHRONIC [RfD] COMMENT: DETERMINED FROM DATA OBTAINED WITH THE DIHYDROCHLORIDE									
TOLUENEDIAMINE, 2,3-		**002687-25-4**							
GENERAL COMMENT		DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010071
TOLUENEDIAMINE, 3,4-		**000496-72-0**							
GENERAL COMMENT:		DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010072
TOLUIDINE, M-		**000108-44-1**							
GENERAL COMMENT		DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT							010075
TRIALLATE		**002303-17-5**							
NOAEL 1 275 MG/KG/DAY		DOG							
	ORAL. DIET	24 MONTHS	SPLEEN	EFFECTS		1.3E-2		IRIS	010076
			LIVER	EFFECTS		100			
SUBCHRONIC [RfD] COMMENT THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TRIBROMOBENZENE, 1,2,4- 000615-54-3									
NOAEL 5 MG/KG/DAY	ORAL. DIET	RAT 45 OR 90 DAYS	LIVER	ALTERED WEIGHT		5E-2		IRIS	010077
			LIVER	ENZYME INDUCTION		100			
TRICHLORO-1,2,2-TRIFLUOROETHANE, 1,1,2- 000076-13-1									
								IRIS	010460
NOEL 2000 PPM	INHALATION. INTERMITTENT	RAT 24 MONTHS	WHOLE BODY	DECREASED WEIGHT	3E+1 100	3E+0 100	3E+1 100		010376
SUBCHRONIC [RfD] COMMENT BASED ON ROUTE TO ROUTE EXTRAPOLATION USING AN ABSORPTION FACTOR OF 0.2									
TRICHLORO-2'-HYDROXYDIPHENYLETHER, 2,2,4'- 003380-34-5									
NOEL 500 MG/KG/DAY	ORAL	RAT 4 WEEKS	WHOLE BODY	DECREASED WEIGHT		4E+0 100			005492
TRICHLOROBENZENE, 1,2,4- 000120-82-1									
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 WEEKS	LIVER	NON-ADVERSE WEIGHT CHANGES	2E+0 100		2E-1 1000		010958
TRICHLOROCYCLOPENTADIENE 077323-84-3									
GENERAL COMMENT: DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005304

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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 WATER		90 DAYS	BLOOD	CLINICAL CHEMISTRY ALTERATIONS		4E-2 100		IRIS	005702
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 1000			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

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TRICHLOROPROPANE, 1,1,2- 000598-77-6									
NOEL 100 MG/L		RAT							
	ORAL. DRINKING WATER	13 WEEKS	LIVER	HISTOPATHOLOGY		5E-2		IRIS	005708
			KIDNEY	HISTOPATHOLOGY		300			
			THYROID	HISTOPATHOLOGY					
TRICHLOROPROPANE, 1,2,2- 003175-23-3									
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									005706
TRICHLOROPROPANE, 1,2,3- 000096-18-4									
NOAEL 8 MG/KG/DAY		RAT							
	ORAL	120 DAYS	WHOLE BODY	TOXICITY		6E-2		IRIS	005714
			LIVER	LESIONS		100			
			KIDNEY	LESIONS					
			ERYTHROCYTES	DECREASED COUNT					
			BLOOD	DECREASED HEMATOCRIT					
			BLOOD	DECREASED HEMOGLOBIN					
GENERAL COMMENT· ALSO SEE HEAST TABLE 3· CARCINOGENICITY.									
TRICHLOROPROPENE, 1,2,3- 000096-19-5									
NOEL 18 MG/CU M		DOG							
	INHALATION. INTERMITTENT	66 WEEKS	EYE	IRRITATION		5E-3		5E-3	010078
						100		100	

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TRICHLOROTOLUENE, 2,3,6- 002077-46-5									
LOAEL 0.5 PPM		RAT							
	ORAL· DIET	28 DAYS	LIVER	LESIONS		5E-5			005335
			KIDNEY	LESIONS		1000			
			THYROID	LESIONS					
TRICHLOROTOLUENE, ALPHA,2,6- 002014-83-7									
LOAEL 0.5 PPM		RAT							
	ORAL· DIET	28 DAYS	LIVER	LESIONS		5E-5			005339
			KIDNEY	LESIONS		1000			
			THYROID	LESIONS					
TRIFLURALIN 001582-09-8									
NOEL 0.75 MG/KG/DAY		DOG							
	ORAL· DIET	·12 MONTHS	LIVER	INCREASED WEIGHT		7 5E-3		IRIS	010080
			BLOOD	METHEMOGLOBINEMIA		100			

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

TRIMETHYLBENZENES

GENERAL COMMENT· DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT 005727

TRINITROBENZENE, 1,3,5- 000099-35-4									
NOAEL 0 51 MG/KG/DAY		RAT							
	ORAL: WATER	16 WEEKS	SPLEEN	INCREASED WEIGHT		5E-4		IRIS	010081
						1000			

SUBCHRONIC [RfD] COMMENT· CALCULATED FROM DATA OBTAINED WITH 1,3-DINITROBENZENE.
CHRONIC [RfD] COMMENT· CALCULATED FROM DATA OBTAINED WITH 1,3-DINITROBENZENE.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
TRINITROPHENOLS									
GENERAL COMMENT. DATA INADEQUATE FOR QUANTITATIVE RISK ASSESSMENT									010082
TRINITROPHENYLMETHYLNITRAMINE 000479-45-8									
LOAEL 125 MG/KG/DAY		RABBIT							
	ORAL: GAVAGE	9 MONTHS	LIVER	HISTOPATHOLOGICAL EFFECTS		1E-1		1E-2	010377
			KIDNEY	HISTOPATHOLOGICAL EFFECTS		1000		10000	
			SPLEEN	HISTOPATHOLOGICAL EFFECTS					
TRINITROTOLUENE, 2,4,6- 000118-96-7									
LOAEL 0.5 MG/KG/DAY		DOG							
	ORAL: GAVAGE		LIVER	EFFECTS		5E-4		IRIS	010416
						1000			
SUBCHRONIC [RfD] COMMENT: THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									
GENERAL COMMENT ALSO SEE HEAST TABLE 3. CARCINOGENICITY									
VANADIUM 007440-62-2									
NOAEL 5 PPM		RAT							
	ORAL: DRINKING WATER	LIFETIME				7E-3		7E-3	005739
						100		100	
VANADIUM PENTOXIDE 001314-62-1									
NOAEL 17 85 PPM		RAT							
	ORAL: DIET	LIFETIME				9E-3		IRIS	005743
						100			
SUBCHRONIC [RfD] COMMENT. THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
VANADIUM SULFATE		**036907-42-3**								
NOAEL 2.24 MG/KG/DAY		RAT								
	ORAL: DRINKING WATER		LIFETIME				2E-2 100		2E-2 100	005747
VERNAM./ (VERNOLATE)		**001929-77-7**								
NOEL 1 MG/KG/DAY		RAT								
	ORAL: DIET			WHOLE BODY	DECREASED WEIGHT		1E-2 100		IRIS	010083
VINYL ACETATE		**000108-05-4**								
NOAEL 100 MG/KG/DAY		RAT								
	ORAL WATER		2 YEARS	WHOLE BODY	ALTERED WEIGHT		1E+0		1E+0	010417
				KIDNEY	ALTERED WEIGHT		100		100	
NOAEL 176 MG/CU M		MOUSE								
	INHALATION INTERMITTENT		104 WEEKS	NASAL CAVITY	LESIONS	2E-1 30		IRIS		010418

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
WARFARIN		**000081-81-2**							
LOAEL 2 MG/DAY	ORAL	HUMAN	BLOOD	INCREASED PROTHROMBIN TIME		3E-4 100		IRIS	010409

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

XYLENE, M-		**000108-38-3**							
NOAEL 250 MG/KG	ORAL GAVAGE	RAT 103 WEEKS	CENTRAL NERVOUS SYSTEM	HYPERACTIVITY				2E+0 100	005755
			WHOLE BODY	DECREASED WEIGHT					
			WHOLE BODY						

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

010920

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

XYLENE, MIXTURE		**001330-20-7**						IRIS	010872

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

010921

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

XYLENE, O-		**000095-47-6**							
NOEL 250 MG/KG	ORAL GAVAGE	RAT 103 WEEKS	CENTRAL NERVOUS SYSTEM	HYPERACTIVITY				2E+0 100	005751
			WHOLE BODY	DECREASED WEIGHT					

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

010922

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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.									
ZINC (METALLIC)		**007440-66-6**								
LOAEL 1.0 MG/KG/DAY		HUMAN								
	ORAL: DIET		10 WEEKS	BLOOD	DECREASED BLOOD ENZYME		3E-1		IRIS	010937
	SUPPLEMENT						3			
CHRONIC [RfD] COMMENT	THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD].									
ZINC CYANIDE		**000557-21-1**								
NOAEL 24 3 MG/KG/DAY		RAT								
	ORAL. DIET		2 YEARS	WHOLE BODY	DECREASED WEIGHT		5E-2		IRIS	010289
				THYROID	EFFECTS		500			
				NERVE	MYELIN DEGENERATION					
SUBCHRONIC [RfD] COMMENT	CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT.									
CHRONIC [RfD] COMMENT	CALCULATED BY ANALOGY TO FREE CYANIDE BY CORRECTING FOR DIFFERENCES IN MOLECULAR WEIGHT									
ZINC PHOSPHIDE		**001314-84-7**								
LOAEL 3.48 MG/KG/DAY		RAT								
	ORAL: DIET		13 WEEKS	WHOLE BODY	DECREASED WEIGHT		3E-3		IRIS	010290
				WHOLE BODY	DECREASED FOOD INTAKE		1000			
ZINEB		**012122-67-7**								
LOAEL 25 MG/KG/DAY		RAT								
	ORAL. DIET		2 YEARS	THYROID	HYPERPLASIA		5E-2		IRIS	010085
							500			
SUBCHRONIC [RfD] COMMENT·	THE CHRONIC ORAL RfD WAS ADOPTED AS THE SUBCHRONIC ORAL [RfD]									

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July 1997

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US EPA 1987 RfD/RfC WORK GROUP

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US EPA. 1989 RfD/RfC WORK GROUP

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CHLORDANE **000057-74-9**

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US EPA. 1987 RfD/RfC WORK GROUP

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US EPA. 1987. RfD/RfC WORK GROUP

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US EPA 1988 RfD/RfC WORK GROUP.

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US EPA. 1992. RfD/RfC WORK GROUP

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US EPA 1989 RfD/RfC WORK GROUP.

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US EPA. 1990 RfD/RfC WORK GROUP

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US EPA 1986. RfD/RfC WORK GROUP.

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US EPA 1988 RfD/RfC WORK GROUP

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US EPA 1988 RfD/RfC WORK GROUP.

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US EPA. 1990. RfD/RfC WORK GROUP.

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US EPA 1987 RfD/RfC WORK GROUP

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US EPA 1992. RfD/RfC WORK GROUP

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DIETHYLHYDRAZINE, 1,2- **001615-80-1**

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DIMETHOATE **000060-51-5**

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US EPA 1988 RfD/RfC WORK GROUP

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US EPA 1991 RfD/RfC WORK GROUP

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US EPA 1990 RfD/RfC WORK GROUP

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US EPA 1986 RfD/RfC WORK GROUP

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US EPA 1992 RfD/RfC WORK GROUP

NAPHTHOQUINONE, 1,4- 000130-15-4

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US EPA 1991 RfD/RfC WORK GROUP

NICKEL CYANIDE 000557-19-7

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NICKEL, SOLUBLE SALTS **VARIOUS**

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US EPA 1991 RfD/RfC WORK GROUP

NICOTINONITRILE **000100-54-9**

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NITRIC OXIDE **010102-43-9**

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US EPA. 1994 RfD/RfC WORK GROUP.

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US EPA 1988 RfD/RfC WORK GROUP

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US EPA 1986. RfD/RfC WORK GROUP.

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010920 US EPA 1991. RfD/RfC WORK GROUP

XYLENE, MIXTURE **001330-20-7**

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010921 US EPA 1991 RfD/RfC WORK GROUP

XYLENE, O- **000095-47-6**

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US EPA 1986 RfD/RfC WORK GROUP

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

July 1997

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
ACETONE CYANOHYDRIN		**000075-86-5**								
NOEL	4 0 MG/KG/DAY	RAT								
	INHALATION:		14 WEEKS	CENTRAL NERVOUS	EFFECTS	1E-1		1E-2		010432
	INTERMITTENT			SYSTEM		100		1000		

SUBCHRONIC [RfC] COMMENT: 4E-2 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). AN ERROR IN THE UNCERTAINTY FACTOR THAT WAS REPORTED IN HEED (1988) WAS CORRECTED.
CHRONIC [RfC] COMMENT 4E-2 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.

ACETONITRILE		**000075-05-8**								
NOAEL	100 PPM	MOUSE								
	INHALATION		92 DAYS	LIVER	INCREASED RELATIVE WEIGHT	5E-1		5E-2		005208
	INTERMITTENT					300		3000		

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	RAT								
	INHALATION,		4 MONTHS	FETUS	FETOTOXICITY	5E-3		5E-4		005249
	INTERMITTENT					100		1000		

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

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

July 1997

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
CHLORO-1,3-BUTADIENE / (CHLOROPRENE) 000126-99-8									
NOAEL 10 PPM		RAT							
	INHALATION·	2 YEARS	HAIR	ALOPECIA		2E-2		2E-2	005878
	INTERMITTENT		WHOLE BODY	DECREASED WEIGHT GAIN		100		100	

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)

CHLOROBENZENE 000108-90-7									
LOAEL 75 PPM		RAT							
	INHALATION.	120 DAYS	LIVER	EFFECTS			2E-2		005353
	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.

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

DICHLORODIFLUOROMETHANE		**000075-71-8**								
LOAEL 482.3 MG/KG/DAY		GUINEA PIG								
	INHALATION		6 WEEKS	LIVER	LESIONS	2E+0		2E-1		005497
	INTERMITTENT					1000		10000		

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)
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

DICHLOROETHANE, 1,1-		**000075-34-3**								
NOEL 138 MG/KG/DAY		CAT								
	INHALATION		13 WEEKS	KIDNEY	DAMAGE	5E+0		5E-1		005789
	INTERMITTENT					100		1000		

SUBCHRONIC [RfC] COMMENT: 1E+0 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST).
CHRONIC [RfC] COMMENT 1E-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 HEAST TABLE 1. CHRONIC AND SUBCHRONIC TOXICITY AND

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
DICYCLOPENTADIENE		**000077-73-6**							
LOAEL 1 PPM	INHALATION: INTERMITTENT	RAT 90 DAYS	KIDNEY	DYSFUNCTION	2E-3 1000		2E-4 10000		005424

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

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.

FURFURAL		**000098-01-1**							
NOAEL 20 PPM	INHALATION· INTERMITTENT	HAMSTER 13 WEEKS	NASAL CAVITY	OLFACTORY DEGENERATION	5E-1 100		5E-2 1000		005465

SUBCHRONIC [RfC] COMMENT: 1E-1 MG/KG/DAY (SEE APPENDIX A-11, DOSE CONVERSIONS ON HEAST)
CHRONIC [RfC] COMMENT. 1E-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.

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHACRYLONITRILE		**000126-98-7**							
NOEL 3.2 PPM		DOG							
	INHALATION:	90 DAYS	LIVER	INCREASED SGOT	7E-3		7E-4		005811
	INTERMITTENT		LIVER	INCREASED SGPT	300		3000		

SUBCHRONIC [RfC] COMMENT: 2E-3 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST). THESE VALUES DIFFER FROM THOSE IN THE 1987 HEED

CHRONIC [RfC] COMMENT 2E-4 MG/KG/DAY (SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST) THESE VALUES DIFFER FROM THOSE IN THE 1987 HEED.

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.

METHOXYETHANOL ACETATE, 2-		**000110-49-6**							
NOAEL 10 PPM		RABBIT							
	INHALATION	13 WEEKS	TESTIS	DEGENERATION		2E-2		2E-3	010001
	INTERMITTENT					100		1000	

SUBCHRONIC [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).

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

METHOXYETHANOL, 2-		**000109-86-4**							
NOAEL 31 MG/CU M		RABBIT							
	INHALATION.	13 WEEKS	TESTICLE	EFFECTS		1E-2		1E-3	010910
	INTERMITTENT					100		1000	

SUBCHRONIC [RfD] COMMENT. BASED ON ROUTE TO ROUTE EXTRAPOLATION

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

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHYL ACRYLATE		**000096-33-3**								
NOEL 15 PPM		RAT								
	INHALATION:		2 YEARS		NONE OBSERVED		3E-2		3E-2	010003
	INTERMITTENT						100		100	

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 KETONE		**000108-10-1**								
NOEL 50 PPM		RAT								
	INHALATION.		90 DAYS	LIVER	INCREASED WEIGHT	8E-1		8E-2		005562
	INTERMITTENT			KIDNEY	EFFECTS	100		1000		

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHYL STYRENE (MIXED ISOMERS) 025013-15-4									
LOAEL 5 6 MG/KG/DAY		MOUSE							
	INHALATION:	103 WEEKS	NASAL CAVITY	LESIONS		6E-3		6E-3	005567
	INTERMITTENT					1000		1000	

SUBCHRONIC [RfD] COMMENT· BASED ON ROUTE TO ROUTE EXTRAPOLATION WITH AN ABSORPTION FACTOR OF 0.5
CHRONIC [RfD] COMMENT BASED ON ROUTE TO ROUTE EXTRAPOLATION WITH AN ABSORPTION FACTOR OF 0.5.
GENERAL COMMENT: NO PHARMACOKINETIC DATA WERE PROVIDED TO JUSTIFY ROUTE TO ROUTE EXTRAPOLATION FOR THE ORAL [RfD] VALUES.

LOAEL 11 2 MG/KG/DAY		MOUSE							
	INHALATION.	103 WEEKS	NASAL CAVITY	LESIONS	4E-2		4E-2		005566
	INTERMITTENT				1000		1000		

SUBCHRONIC [RfC] COMMENT: 1E-2 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).
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.

METHYL STYRENE, ALPHA 000098-83-9									
NOEL 970 MG/CU M		RAT							
	INHALATION·	197 DAYS	LIVER	INCREASED WEIGHT		7E-1		7E-2	010016
	INTERMITTENT		KIDNEY	INCREASED WEIGHT		100		1000	

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT	LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
METHYLENE BROMIDE		**000074-95-3**								
NOAEL 11 MG/KG/DAY		RAT								
	INHALATION:	90 DAYS		BLOOD	INCREASED CARBOXYHEMOGLOBIN		1E-1		1E-2	010011
	INTERMITTENT						100		1000	

SUBCHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION, INCLUDING AN ABSORPTION FACTOR OF 0.5.
CHRONIC [RfD] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION, INCLUDING AN ABSORPTION FACTOR OF 0.5.
GENERAL COMMENT: NO PHARMACOKINETIC DATA WERE PROVIDED TO JUSTIFY ROUTE TO ROUTE EXTRAPOLATION FOR THE ORAL [RfD] VALUES.

NITROBENZENE		**000098-95-3**								
LOAEL 25 MG/CU M		MOUSE								
	INHALATION:	90 DAYS		BLOOD	HEMATOLOGICAL EFFECTS	2E-2		2E-3		010518
	INTERMITTENT			ADRENAL	LESIONS	1000		10000		
				KIDNEY	LESIONS					
				LIVER	LESIONS					
		RAT								
	INHALATION.	90 DAYS		BLOOD	HEMATOLOGICAL EFFECTS					
	INTERMITTENT			ADRENAL	LESIONS					
				KIDNEY	LESIONS					
				LIVER	LESIONS					

SUBCHRONIC [RfC] COMMENT. 6E-3 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 [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

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

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

CHEMICAL LEVEL	DOSE ROUTE	SPECIES EXPERIMENT LENGTH	TARGET	CRITICAL EFFECT	Subchronic [RfC] (mg/cu m) UF	Subchronic [RfD] (mg/kg/day) UF	Chronic [RfC] (mg/cu m) UF	Chronic [RfD] (mg/kg/day) UF	REFERENCE
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+0 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

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).

US EPA. 1988. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR CYANOHYDRINS. 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.

ACETONITRILE **000075-05-8**

005208 COATE WB. 1983 90-DAY SUBCHRONIC TOXICITY STUDY OF ACETONITRILE IN B6C3F1 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 M, O PROMIN AND A SILAYEV 1977. BARIUM COMPOUNDS AS INDUSTRIAL POISONS (AN EXPERIMENTAL STUDY). J HYG EPIDEM MICROBIOL IMMUNOL. 21: 361.

US EPA 1984. HEALTH EFFECTS ASSESSMENT FOR BARIUM. 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

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.

US EPA 1989. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR 2-CHLORO-1,3-BUTADIENE. 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.

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.

US EPA. 1989. UPDATED HEALTH EFFECTS ASSESSMENT FOR CHLOROBENZENE. 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.

CYCLOPENTADIENE **000542-92-7**

005401 DOW. 1987 UNPUBLISHED DATA. DOW CHEMICAL, USA, MIDLAND, MI.

US EPA. 1987 HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR CYCLOPENTADIENE AND DICYCLOPENTADIENE. 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.

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

US EPA 1984 HEALTH EFFECTS ASSESSMENT FOR 1,1-DICHLOROETHANE. 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.

DICYCLOPENTADIENE **000077-73-6**

005424 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 ID 88-8300464, ODD DOC ID 8EHQ-0283-0364,MICROFICHE NO OTS 204864

US EPA. 1987 HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR CYCLOPENTADIENE AND DICYCLOPENTADIENE 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

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

US EPA. 1985. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR 2- ETHOXYETHANOL ESTERS. 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.

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.

US EPA. 1988. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR FURFURAL. 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

METHACRYLONITRILE 000126-98-7

005811 POZZANI, UC, CR KINKEAD AND JM KING. 1968 THE MAMMALIAN TOXICITY OF METHACRYLONITRILE AM IND HYG ASSOC J. 29(3): 202-210.

US EPA 1987. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR SELECTED NITRILES, 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

METHOXYETHANOL ACETATE, 2- 000110-49-6

010001 MILLER RR, LL CALHOUN AND BL YANO. 1982. ETHYLENE GLYCOL MONOMETHYL ETHER: 13-WEEK VAPOR INHALATION STUDY IN MALE RABBITS REPORT PREPARED FOR THE CHEMICAL MANUFACTURERS ASSOCIATION, MARCH 25, 1982.

US EPA 1987. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR 2- METHOXYETHANOL ACETATE. 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, OC

METHOXYETHANOL, 2- 000109-86-4

010910 MILLER RR, LL CALHOUN, BL YANO 1982. ETHYLENE GLYCOL MONOETHYL ETHER: 13 WEEK VAPOR INHALATION STUDY IN MALE RABBITS. REPORT PREPARED FOR THE CMA MARCH 25, 1982.

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METHYL ACRYLATE 000096-33-3

010003 KLIMISCH HJ AND W REININGHAUS 1984 CARCINOGENICITY OF ACRYLATES LONG-TERM INHALATION STUDIES ON METHYL ACRYLATE (MA) AND N-BUTYL ACRYLATE (BA) IN RATS. TOXICOLOGIST. 4(1) 53

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METHYL ISOBUTYL KETONE **000108-10-1**

005562 UNION CARBIDE CORP. 1983 NINETY-DAY INHALATION STUDY IN RATS AND MICE SPONSORED BY CMA US EPA/OTS PUBLIC FILES 0750507469.

US EPA 1987. HEALTH EFFECTS ASSESSMENT FOR METHYL ETHYL KETONE. 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

METHYL STYRENE (MIXED ISOMERS) **025013-15-4**

005567 MRI (MIDWEST RESEARCH INSTITUTE) 1984. STUDY OF THE INHALATION CARCINOGENICITY (BIOASSAY) OF VINYL TOLUENE IN B6C3F1 MICE

US EPA 1987. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR METHYL STYRENES 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

005566 FINAL CHRONIC REPORT PERFORMED FOR THE NTP UNDER CONTRACT NO N01-ES-38042.

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METHYL STYRENE, ALPHA **000098-83-9**

010016 WOLF MA, VK ROWE, DD MCCOLLISTER, ET AL 1956 TOXICOLOGICAL STUDIES OF CERTAIN ALKYLATED BENZENES AND BENZENE ARCH IND HEALTH 14· 387-398.

US EPA 1987. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR ALPHA- METHYL STYRENE. 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.

METHYLENE BROMIDE **000074-95-3**

010011 KEYES DG, JW HENCK, GC JERSEY, RJ KOCIBA, DJ SCHWETZ AND TD LANDRY 1982 METHYLENE BROMIDE: A 90-DAY REPEATED INHALATION TOXICITY STUDY IN RATS AND DOGS WITH A SUBSEQUENT TWO-YEAR HOLDING PERIOD FOR RATS. TOXICOLOGY RESEARCH LAB, HEALTH AND ENVIRONMENTAL SCIENCES, DOW CHEMICAL CO , MIDLAND, MI.

US EPA 1987 HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR METHYLENE BROMIDE. 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.

NITROBENZENE **000098-95-3**

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.

US EPA. 1987. HEALTH EFFECTS ASSESSMENT FOR NITROBENZENE. 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.

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

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.

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
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 WATER	2 YEARS RAT	MAMMARY	TUMORS	IRIS	IRIS	4.5E+0	IRIS	IRIS	010087
			THYROID	TUMORS						
			UTERUS	TUMORS						
			ORAL CAVITY	TUMORS						
			CENTRAL NERVOUS SYSTEM	TUMORS						
	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
	OCCUPATIONAL	HUMAN	LUNG	TUMORS						
	INHALATION [SLOPE] COMMENT SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
ALACHLOR		**015972-60-8**								
	ORAL: DIET	MULTIPLE SITES	TUMORS		B2	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)									
ALDRIN		**000309-00-2**								
	ORAL: DIET	MOUSE	LIVER	CARCINOMA	IRIS	IRIS	1.7E+1	IRIS	IRIS	005006
	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: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
ANILINE		**000062-53-3**								
					IRIS	IRIS		IRIS		010088
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1. SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
ARAMITE		**000140-57-8**								
	ORAL· DIET	104 WKS / RAT	LIVER	TUMORS	IRIS	IRIS	2.5E-2	IRIS	IRIS	010206
	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)									

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
ARSENIC, INORGANIC		**007440-38-2**								
						IRIS		IRIS		010925
	INHALATION: OCCUPATIONAL	HUMAN	RESPIRATORY SYSTEM	TUMORS	IRIS		IRIS		IRIS	005007
	GENERAL COMMENT: ALSO SEE HEAST TABLE I: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
ASBESTOS		**001332-21-4**								
					IRIS					005010
					IRIS				IRIS	005919
ATRAZINE		**001912-24-9**								
	ORAL: DIET	2 YEARS RAT	MAMMARY GLAND	ADENOMA	C	2.22E-1		6.3E-6		010380
			MAMMARY GLAND	FIBROADENOMA						
			MAMMARY GLAND	ADENOCARCINOMA						
			MAMMARY GLAND	CARCINOSARCOMA						
	ORAL [SLOPE] COMMENT· UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
	GENERAL COMMENT ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
AZOBENZENE		**000103-33-3**								
	ORAL DIET	2 YEARS RAT	ABDOMINAL CAVITY	SARCOMA	IRIS	IRIS	1 1E-1	IRIS	IRIS	010089
	INHALATION [SLOPE] COMMENT. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.									

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
BENZENE		**000071-43-2**								
	INHALATION: OCCUPATIONAL	HUMAN	BLOOD	LEUKEMIA	IRIS	IRIS	2.9E-2	IRIS	IRIS	005011
	ORAL [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION 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)									
BENZIDINE		**000092-87-5**								
					IRIS	IRIS	IRIS	IRIS	IRIS	005014
	GENERAL COMMENT· ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
BENZOTRICHLORIDE		**000098-07-7**								
					IRIS	IRIS		IRIS		010092
BENZO[A]ANTHRACENE		**000056-55-3**								
					IRIS					010182
BENZO[A]PYRENE		**000050-32-8**								
					IRIS	IRIS		IRIS		010508
BENZO[B]FLUORANTHENE		**000205-99-2**								
					IRIS					010183
BENZO[K]FLUORANTHENE		**000207-08-9**								
					IRIS					010090
BENZYL CHLORIDE		**000100-44-7**								
					IRIS	IRIS		IRIS		010093

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
BERYLLIUM		**007440-41-7**								
					IRIS	IRIS		IRIS		005018
	GENERAL COMMENT ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
	INHALATION:				IRIS		8.4E+0		IRIS	005017
	OCCUPATIONAL	HUMAN	LUNG	TUMORS						
	INHALATION [SLOPE] COMMENT. SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST									
BIS(2-CHLOROETHYL) ETHER		**000111-44-4**								
	ORAL	560 DAYS			IRIS	IRIS	1.1E+0	IRIS	IRIS	005076
		MOUSE	LIVER	TUMORS						
	INHALATION [SLOPE] COMMENT: _BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST									
BIS(2-CHLORO-1-METHYLETHYL) ETHER		**000108-60-1**								
	ORAL GAVAGE	2 YEARS			C	7E-2	3.5E-2	2E-6	1E-5	005079
		MOUSE	LIVER	TUMORS						
			LUNG	TUMORS						
	INHALATION [SLOPE] COMMENT. BASED ON ROUTE TO ROUTE EXTRAPOLATION (50% RESPIRATORY ABSORPTION) SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST. GENERAL COMMENT. COMPOUND TESTED CONTAINED 70% BIS(2-CHLORO-1-METHYLETHYL)ETHER AND 30% BIS(2-CHLOROISOPROPYL)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 COMMENT: ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
BIS(CHLOROMETHYL) ETHER		**000542-88-1**								
	INHALATION: INTERMITTENT	10-100 DAYS RAT	RESPIRATORY SYSTEM	TUMORS	IRIS	IRIS	2.2E+2	IRIS	IRIS	005077
	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: INTERMITTENT	2 YEARS RAT	LIVER	TUMORS	B2		1.1E-1		3 2E-5	010094
	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)									
BROMOFORM		**000075-25-2**								
	ORAL · GAVAGE	2 YEARS RAT	INTESTINE, LARGE	ADENOMATOUS POLYP ADENOCARCINOMA	IRIS	IRIS	3.9E-3	IRIS	IRIS	005150
	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).									
BUTADIENE, 1,3-		**000106-99-0**								
	INHALATION: INTERMITTENT	MOUSE	MULTIPLE SITES	TUMORS	IRIS		1.8E+0		IRIS	010477
	INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST									

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
BUTYL BENZYL PHTHALATE, N-		**000085-68-7**								
					IRIS					005122
	GENERAL COMMENT. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
CADMIUM		**007440-43-9**								
					IRIS				IRIS	005019
	ORAL [SLOPE] 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**								
	ORAL: DIET				C	8.6E-3		2.4E-7		010095
		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**								
					B2	3.5E-3		1.0E-7		010184
	ORAL [SLOPE] COMMENT. UNDER REVIEW. NUMBER SUBJECT TO CHANGE									
	GENERAL COMMENT· ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
CARBAZOLE		**000086-74-8**								
	ORAL: DIET	96 WEEKS			B2	2E-2		5.7E-7		010096
		MOUSE	LIVER	TUMORS						
CARBON TETRACHLORIDE		**000056-23-5**								
	ORAL: DIET		LIVER	TUMORS	IRIS	IRIS	5 3E-2	IRIS	IRIS	005022
	INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II DOSE CONVERSIONS ON HEAST									
	INHALATION [UNIT RISK] COMMENT. BASED ON ROUTE TO ROUTE EXTRAPOLATION. INCORPORATES AN ABSORPTION FACTOR OF 0.4.									
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
CHLORANIL		**000118-75-2**								
	ORAL. DIET	82 WEEKS			C	4.03E-1		1.2E-5		010097
		MOUSE	LIVER	TUMORS						
			LUNG	TUMORS						
CHLORDANE		**000057-74-9**								
	ORAL· DIET				IRIS	IRIS	1.3E+0	IRIS	IRIS	005024
		MOUSE	LIVER	CARCINOMA						

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).

CHLORO-2-METHYLANILINE HYDROCHLORIDE, 4-		**003165-93-3**								
	ORAL: DIET	18 MONTHS			B2	4.6E-1		1.3E-5		010419
		MOUSE	CARDIOVASCULAR SYSTEM	HEMANGIOMA						
			CARDIOVASCULAR SYSTEM	HEMANGIOSARCOMA						
CHLORO-2-METHYLANILINE, 4-		**000095-69-2**								
	ORAL· DIET	18 MONTHS			B2	5.8E-1		1 6E-5		010098
		MOUSE	CARDIOVASCULAR SYSTEM	HEMANGIOMA						
			CARDIOVASCULAR SYSTEM	HEMANGIOSARCOMA						

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

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
CHLOROBENZILATE		**000510-15-6**								
	ORAL: GAVAGE, DIET	82 WEEKS MOUSE	LIVER	HEPATOMA	B2	2.7E-1	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 PHARMACOKINETIC DATA TO THE CONTRARY. SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST									
	INHALATION [UNIT RISK] 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		IRIS		005036
	ORAL. GAVAGE	78 WEEKS MOUSE	LIVER	CARCINOMA	IRIS		8.1E-2		IRIS	005035
	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).									
CHLOROMETHANE		**000074-87-3**								
	INHALATION· INTERMITTENT	24 MONTHS MOUSE	KIDNEY	TUMORS	C	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).									
CHLOROMETHYL METHYL ETHER		**000107-30-2**								
					IRIS					005081
CHLORONITROBENZENE, O-		**000088-73-3**								
	ORAL DIET	18 MONTHS MOUSE	LIVER	TUMORS	B2	2.5E-2		7.1E-7		010099

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
CHLORONITROBENZENE, P-		**000100-00-5**								
	ORAL: DIET	18 MONTHS MOUSE	CARDIOVASCULAR SYSTEM	TUMORS	B2	1.8E-2		5.1E-7		010100
CHLOROTHALONIL		**001897-45-6**								
	ORAL: DIET	27-32 MONTHS RAT	KIDNEY	TUMOR	B2	1.1E-2		3 1E-7		010384
	ORAL [SLOPE] COMMENT. UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
CHROMIUM(VI)		**018540-29-9**								
	INHALATION· OCCUPATIONAL	HUMAN	LUNG	TUMORS	IRIS		4 1E+1		IRIS	005091
	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).									
CHRYSENE		**000218-01-9**								
					IRIS					010185
	GENERAL COMMENT: SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
COKE OVEN EMISSIONS		**008007-45-2**								
	INHALATION: OCCUPATIONAL	HUMAN	LUNG	TUMORS	IRIS		2.2E+0		IRIS	005039
	INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.									
	GENERAL COMMENT: FORMERLY LISTED UNDER COAL TARS.									
CREOSOTE, COAL TAR		**008001-58-9**								
					IRIS					005042

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
CRESOL, M- / (3-METHYLPHENOL) 000108-39-4										
					IRIS					010187
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
CRESOL, O- / (2-METHYLPHENOL) 000095-48-7										
					IRIS					010186
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
CRESOL, P- / (4-METHYLPHENOL) 000106-44-5										
					IRIS					010188
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
CROTONALDEHYDE 000123-73-9										
	ORAL· DRINKING WATER	113 WKS RAT	LIVER	TUMOR	IRIS	1.9E+0		5.4E-5		010190
	ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
CYANAZINE 021725-46-2										
	ORAL: DIET	2 YEARS RAT	MAMMARY GLAND	ADENOMA/ CARCINOMA, COMBINED	C	8.4E-1		2 4E-5		010944
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
DDD 000072-54-8										
					IRIS	IRIS		IRIS		010291

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DDE		**000072-55-9**								
					IRIS	IRIS		IRIS		010292
DDT		**000050-29-3**								
	ORAL· DIET				IRIS	IRIS	3.4E-1	IRIS	IRIS	005044
		MOUSE, RAT	LIVER	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).

DECABROMODIPHENYL ETHER		**001163-19-5**								
					IRIS					010102

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

DIALLATE		**002303-16-4**								
	ORAL	19 MONTHS			B2	6.1E-2		1 7E-6		010103
		MOUSE	LIVER	TUMORS						

DIBENZO[A,H]ANTHRACENE		**000053-70-3**								
					IRIS					010191

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DIBROMO-3-CHLOROPROPANE, 1,2	**000096-12-8**									
	ORAL: DIET				B2	1.4E+0		4E-5		010484
			STOMACH	TUMORS						
			KIDNEY	TUMORS						
			LIVER	TUMORS						
	[EPA GROUP] COMMENT: UNDER REVIEW. CLASSIFICATION SUBJECT TO CHANGE									
	INHALATION [SLOPE] COMMENT. SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST.									
	INHALATION·				B2		2 4E-3		6 9E-7	010519
	INTERMITTENT	RAT, MOUSE	NASAL CAVITY	TUMORS						
DIBROMOCHLOROMETHANE		**000124-48-1**								
					IRIS	IRIS		IRIS		010891
	GENERAL COMMENT FORMERLY LISTED AS CHLORODIBROMOMETHANE. ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
DIBROMOETHANE, 1,2-		**000106-93-4**								
					IRIS	IRIS		IRIS		005818
	GENERAL COMMENT FORMERLY LISTED UNDER ETHYLENE DIBROMIDE									
	INHALATION	88-103 WEEKS			IRIS		7 6E-1		IRIS	005071
	INTERMITTENT	RAT	NASAL CAVITY	TUMORS						
	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 FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DICHLORO-2-BUTENE, 1,4-		**000764-41-0**								
	INHALATION INTERMITTENT	90 DAYS RAT	NASAL PASSAGES	TUMORS	B2		9.3E+0		2 6E-3	005053
	INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II: DOSE CONVERSIONS ON HEAST.									
	GENERAL COMMENT. FORMERLY LISTED UNDER DICHLOROBUTENES									
DICHLOROBENZENE, 1,4-		**000106-46-7**								
	ORAL: GAVAGE	103 WEEKS MOUSE	LIVER	TUMORS	C	2.4E-2		6 8E-7		005050
	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 RAT	CIRCULATORY SYSTEM	SARCOMA	IRIS	IRIS		IRIS	IRIS	005058
	GENERAL COMMENT. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DICHLOROETHYLENE, 1,1-		**000075-35-4**								
					IRIS	IRIS		IRIS		005060
	INHALATION	12 MONTHS			IRIS		1.2E+0		IRIS	005059
		MOUSE	KIDNEY	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).									
DICHLOROPROPANE, 1,2-		**000078-87-5**								
	ORAL: GAVAGE				B2	6.8E-2		1.9E-6		005062
		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)									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DICHLOROPROPENE, 1,3- / (TELONE II)		**000542-75-6**								
	ORAL. GAVAGE	104 WEEKS			IRIS	1 8E-1		5E-6		010946
		MOUSE	BLADDER	CARCINOMA						
			RESPIRATORY SYSTEM	ALVEOLAR/ BRONCHIOLAR ADENOMA						
			FORESTOMACH	PAPILLOMA/ CARCINOMA						
	ORAL: GAVAGE	104 WEEKS								
		RAT	LIVER	NEOPLASTIC NODULE/CARCINOMA						
			FORESTOMACH	PAPILLOMA/ CARCINOMA						
	ORAL [SLOPE] COMMENT· THE [SLOPE] IS THE GEOMETRIC MEAN OF SLOPE FACTORS OF COMBINED TUMORS LISTED									
	GENERAL COMMENT ALSO SEE HEAST TABLE 1. SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
	INHALATION:	2 YEARS			IRIS		1 3E-1		3.7E-5	010104
	INTERMITTENT	MOUSE	LUNG	ADENOMA						
	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)									
DIELDRIN		**000060-57-1**								
	ORAL. DIET				IRIS	IRIS	1.6E+1	IRIS	IRIS	005816
		MOUSE	LIVER	CARCINOMA						
	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)									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DIETHYLSTILBESTROL		**000056-53-1**								
	ORAL DIET				A	4.7E+3		1 3E-1		010485
		MOUSE	MAMMARY GLAND	CARCINOMA						
DIMETHOXYBENZIDINE, 3,3'-		**000119-90-4**								
	ORAL: DIET	LIFETIME			B2	1 4E-2		4E-7		010106
		HAMSTER	FORESTOMACH	PAPILLOMA						
DIMETHYLANILINE HYDROCHLORIDE, 2,4-		**021436-96-4**								
	ORAL: DIET	18 MONTHS			C	5 8E-1		1.7E-5		010108
		MOUSE	LUNG	TUMORS						
DIMETHYLANILINE, 2,4-		**000095-68-1**								
	ORAL: DIET	18 MONTHS			C	7.5E-1		2.1E-5		010107
		MOUSE	LUNG	TUMORS						
DIMETHYLBENZIDINE, 3,3'-		**000119-93-7**								
	ORAL: GAVAGE	30 DAYS			B2	9 2E+0		2 6E-4		010109
		RAT	MAMMARY	TUMORS						
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**								
					B2					010962
DIMETHYLSULFATE		**000077-78-1**								
					IRIS					010112

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

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DINITROTOLUENE, 2,4-		**000121-14-2**								
					IRIS	IRIS		IRIS		005066
	GENERAL COMMENT: LISTED AS "DINITROTOLUENE MIXTURE, 2,4-/2,6-" ON IRIS. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
DINITROTOLUENE, 2,6-		**000606-20-2**								
					IRIS	IRIS		IRIS		005068
	GENERAL COMMENT: LISTED AS "DINITROTOLUENE MIXTURE, 2,4-/2,6-" ON IRIS. ALSO SEE HEAST 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
		RAT	LIVER	TUMORS						
	INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.									
DIRECT BLACK 38		**001937-37-7**								
	ORAL: DIET	93 DAYS			A	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			A	8.1E+0		2.3E-4		010114
		RAT	LIVER	TUMORS						
	ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
DIRECT BROWN 95		**016071-86-6**								
	ORAL: DIET	91 DAYS			A	9.3E+0		2 6E-4		010115
		RAT	LIVER	TUMORS						
	ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
DIRECT SKY BLUE 6B		**002610-05-1**								
					B2					010116
	[EPA GROUP] COMMENT: BASED ON THE FACT THAT 3,3-DIMETHOXYBENZIDINE, A KNOWN EPA GROUP B2 CARCINOGEN, IS A METABOLITE OF DIRECT SKY BLUE 6B.									
EPICHLOROHYDRIN		**000106-89-8**								
					IRIS	IRIS		IRIS		010198
	INHALATION: INTERMITTENT	30 DAYS, OBSERVED LIFETIME			IRIS		4.2E-3		IRIS	010117
		RAT	NASAL CAVITY	TUMORS						
	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).									
ETHYL ACRYLATE		**000140-88-5**								
	ORAL, GAVAGE	104 WEEKS			B2	4 8E-2		1 4E-6		010118
		RAT	FORESTOMACH	TUMORS						

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
ETHYLENE OXIDE		**000075-21-8**								
	ORAL· GAVAGE	LIFETIME, TWICE WEEKLY			B1	1.02E+0		2.9E-5		010421
		RAT	STOMACH	TUMORS						
	ORAL [SLOPE] COMMENT· UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
	INHALATION. INTERMITTENT	2 YEARS			B1		3.5E-1		1E-4	010422
		RAT	BLOOD	LEUKEMIA						
			BRAIN	GLIOMA						
	INHALATION [UNIT RISK] COMMENT· UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
ETHYLENE THIOUREA		**000096-45-7**								
	ORAL. GAVAGE	2 YEARS			B2	1.1E-1		3.4E-6		010947
		MOUSE	LIVER	ADENOMA/ CARCINOMA, COMBINED						
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
FOLPET		**000133-07-3**								
					IRIS	IRIS		IRIS		010120
	GENERAL COMMENT ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
FORMALDEHYDE		**000050-00-0**								
	INHALATION	24 MONTHS			IRIS		4 5E-2		IRIS	010121
		RAT	NASAL CAVITY	TUMORS						
	INHALATION [SLOPE] COMMENT· SEE APPENDIX A-1I: 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 FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
FURAZOLIDONE		**000067-45-8**								
	ORAL· DIET	45 WEEKS			B2	3 8E+0		1E-4		005106
		RAT	MAMMARY	TUMORS						
	GENERAL COMMENT. FORMERLY LISTED UNDER NITROFURANS									
FURIUM		**000531-82-8**								
	ORAL: DIET	28 WEEKS			B2	5 0E+1		1 4E-3		005108
		MOUSE	BLOOD	LEUKEMIA						
	GENERAL COMMENT. FORMERLY LISTED UNDER NITROFURANS									
GLYCIDALDEHYDE		**000765-34-4**								
					IRIS					010122
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
HEPTACHLOR		**000076-44-8**								
	ORAL: DIET				IRIS	IRIS	4.5E+0	IRIS	IRIS	005820
		MOUSE	LIVER	CARCINOMA						
	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).									
HEPTACHLOR EPOXIDE		**001024-57-3**								
	ORAL: DIET	18-24 MONTHS			IRIS	IRIS	9.1E+0	IRIS	IRIS	010424
		MOUSE	LIVER	CARCINOMA						
	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)									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
HEXACHLOROBENZENE		**000118-74-1**								
	ORAL: DIET				IRIS	IRIS	1.6E+0	IRIS	IRIS	010365
		RAT	LIVER	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).

HEXACHLOROBUTADIENE		**000087-68-3**								
	ORAL: DIET	22-24 MONTHS			IRIS	IRIS	7.8E-2	IRIS	IRIS	005088
		RAT	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**								
	ORAL: DIET	24 WEEKS			IRIS	IRIS	6.3E+0	IRIS	IRIS	010123
		MOUSE	LIVER	TUMORS						

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

HEXACHLOROCYCLOHEXANE, BETA-		**000319-85-7**								
	ORAL: DIET	110 WEEKS			IRIS	IRIS	1.8E+0	IRIS	IRIS	010124
		MOUSE	LIVER	TUMORS						

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

HEXACHLOROCYCLOHEXANE, DELTA-		**000319-86-8**								
					IRIS					010125

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

HEXACHLOROCYCLOHEXANE, EPSILON-		**006108-10-7**								
					IRIS					010126

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

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
HEXACHLOROCYCLOHEXANE, GAMMA- 000058-89-9										
	ORAL: DIET	110 WEEKS MOUSE	LIVER	TUMORS	B2-C	1.3E+0		3.7E-5		005098
	ORAL [SLOPE] COMMENT UNDER REVIEW. NUMBER SUBJECT TO CHANGE.									
	GENERAL COMMENT· ALSO SEE HEAST TABLE 1. SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
HEXACHLOROCYCLOHEXANE-TECHNICAL 000608-73-1										
	ORAL: DIET	6-20 MONTHS MOUSE	LIVER	TUMORS	IRIS	IRIS	1.8E+0	IRIS	IRIS	010127
	INHALATION [SLOPE] COMMENT BASED ON ROUTE TO ROUTE EXTRAPOLATION SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST.									
HEXACHLOROETHANE 000067-72-1										
	ORAL. GAVAGE	78 WEEKS MOUSE	LIVER	CARCINOMA	IRIS	IRIS	1 4E-2	IRIS	IRIS	005090
	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).									
HYDRAZINE 000302-01-2										
					IRIS	IRIS		IRIS		010129
	INHALATION: INTERMITTENT	1 YEAR RAT	NASAL CAVITY	TUMORS	IRIS		1 7E+1		IRIS	010128
	INHALATION [SLOPE] COMMENT SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
HYDRAZINE SULFATE		**010034-93-2**								
					IRIS	IRIS		IRIS		010131
	INHALATION: INTERMITTENT	1 YEAR RAT	NASAL CAVITY	TUMORS	IRIS		1 7E+1		IRIS	010130
	INHALATION [SLOPE] COMMENT· SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.									
	GENERAL COMMENT LISTED UNDER "HYDRAZINE" ON IRIS.									
INDENO[1,2,3-CD]PYRENE		**000193-39-5**								
					IRIS					010192
ISOPHORONE		**000078-59-1**								
					IRIS	IRIS		IRIS		005094
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1. SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
LEAD		**007439-92-1**								
					IRIS					005096
	GENERAL COMMENT ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
LINURON		**000330-55-2**								
					IRIS					010383
	GENERAL COMMENT. ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
MERCURIC CHLORIDE		**007487-94-7**								
					IRIS					010971
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
MERCURY, ELEMENTAL		**007439-97-6**								
					IRIS					010973
	GENERAL COMMENT	ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)								
METHOXY-5-NITROANILINE, 2-		**000099-59-2**								
	ORAL. DIET	104 WEEKS			B2	4.6E-2		1 3E-6		010132
		RAT	SKIN	CARCINOMA						
METHYLHYDRAZINE		**000060-34-4**								
	GENERAL COMMENT·	NO EPA GROUP CLASSIFICATION WAS PROVIDED IN THE REFERENCE DOCUMENT. THEREFORE THIS COMPOUND WAS REMOVED FROM TABLE 3								
METHYL-5-NITROANILINE, 2-		**000099-55-8**								
	ORAL· DIET	98 WEEKS			C	3 3E-2		9.4E-7		010140
		MOUSE	LIVER	CARCINOMA						
METHYLANILINE HYDROCHLORIDE, 2-		**000636-21-5**								
	ORAL· DIET	93 WEEKS			B2	1 8E-1		5.1E-6		010134
		RAT	SKIN	FIBROMA						
METHYLANILINE, 2-		**000095-53-4**								
	ORAL· DIET	93 WEEKS			B2	2.4E-1		6 9E-6		010133
		RAT	SKIN	FIBROMA						
	GENERAL COMMENT	THE 1984 HEEP CALLED THIS COMPOUND O-TOLUIDINE. THE 1987 HEEP CALLED IT 2-METHYLANILINE								
METHYLENE CHLORIDE / (DICHLOROMETHANE)		**000075-09-2**								
					IRIS	IRIS		IRIS		005100
					IRIS				IRIS	005904
	GENERAL COMMENT.	ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)								

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)^{1}	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
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 RAT	LUNG	TUMORS	B2	1.3E-1	1 3E-1	3.7E-6	3.7E-5	010425
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).										
METHYLMERCURY 0022967-92-6										
					IRIS					010972
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 RAT	LIVER	ADENOMA	B2					010952
			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 4B.										

CHEMICAL	ROUTE	EXPERIMENT LENGTH SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
NICKEL REFINERY DUST										
	INHALATION:				IRIS		8.4E-1		IRIS	005103
	OCCUPATIONAL	HUMAN	RESPIRATORY SYSTEM	TUMORS						

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.
GENERAL COMMENT: FORMERLY LISTED AS NICKEL. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (UNDER NICKEL, SOLUBLE SALTS).

NICKEL SUBSULFIDE		**012035-72-2**								
	INHALATION:				IRIS		1.7E+0		IRIS	005768
	OCCUPATIONAL	HUMAN	RESPIRATORY SYSTEM	TUMORS						

INHALATION [SLOPE] COMMENT: SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST.
GENERAL COMMENT· FORMERLY LISTED AS NICKEL. ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (UNDER NICKEL).

NITROBENZENE		**000098-95-3**								
	INHALATION	2 YEAR MICE			B2					010969
			LUNG	ADENOMA						
			LUNG	CARCINOMA						
			THYROID	ADENOMA						
			MAMMARY	ADENOCARCINOMA						
	INHALATION	2 YEAR RAT								
			LIVER	ADENOMA						
			LIVER	CARCINOMA						
			KIDNEY	ADENOMA						
			KIDNEY	ADENOCARCINOMA						
			UTERUS	ENDIOMETRIAL POLYPS						

GENERAL COMMENT: ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY.

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
NITROFURAZONE		**000059-87-0**								
	ORAL DIET	46 WEEKS			B2	1.5E+0		4.3E-5		005110
		RAT	MAMMARY	TUMORS						
NITROPROPANE, 2-		**000079-46-9**								
	INHALATION	22 MONTHS			B2		9 4E+0		2 7E-3	010142
	INTERMITTENT	RAT	LIVER	TUMORS						
	INHALATION [SLOPE] COMMENT SEE APPENDIX A-II· DOSE CONVERSIONS ON HEAST.									
NITROSO-DI-N-BUTYLAMINE, N-		**000924-16-3**								
	ORAL· DRINKING	LIFETIME			IRIS	IRIS	5 4E+0	IRIS	IRIS	010143
	WATER	MOUSE	BLADDER	TUMORS						
			ESOPHAGUS	TUMORS						
	INHALATION [SLOPE] COMMENT. BASED ON ROUTE TO ROUTE EXTRAPOLATION SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST									
NITROSO-DI-N-PROPYLAMINE, N-		**000621-64-7**								
					IRIS	IRIS		IRIS		010147
NITROSO-N-ETHYLUREA, N-		**000759-73-9**								
	ORAL: DRINKING	203 DAYS			B2	1.4E+2				010426
	WATER	RAT	INTESTINE	GASTROINTESTINAL TUMORS						
	[EPA GROUP] COMMENT. UNDER REVIEW, CLASSIFICATION SUBJECT TO CHANGE.									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
NITROSO-N-METHYLUREA, N- 000684-93-5										
	ORAL· GAVAGE	308 DAYS GUINEA PIG	PANCREAS	ADENOCARCINOMA	B2					010427
	[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 WATER	6 OR 12 MONTHS RAT	LIVER	TUMORS	IRIS	IRIS	1.5E+2	IRIS	IRIS	010145
	INHALATION [SLOPE] COMMENT· BASED ON ROUTE TO ROUTE EXTRAPOLATION SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST									
NITROSODIMETHYLAMINE, N- 000062-75-9										
	ORAL: DRINKING WATER	RAT	LIVER	TUMORS	IRIS	IRIS	5 1E+1	IRIS	IRIS	010146
	INHALATION [SLOPE] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION SEE APPENDIX A-II, DOSE CONVERSIONS ON HEAST									
NITROSODIPHENYLAMINE, N- 000086-30-6										
					IRIS	IRIS		IRIS		005112
NITROSOMETHYLETHYLAMINE, N- 010595-95-6										
					IRIS	IRIS		IRIS		010148

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)^{1}	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
NITROSOMETHYLVINYLAMINE, N	**004549-40-0**									
	INHALATION	RAT	UPPER RESPIRATORY TRACT	CARCINOMAS	B2					010149
	ORAL: DRINKING WATER		UPPER DIGESTIVE TRACT	CARCINOMAS						
NITROSOPYRROLIDINE, N-	**000930-55-2**									
	ORAL: DIET	LIFETIME RAT	LIVER	TUMORS	IRIS	IRIS	2.1E+0	IRIS	IRIS	010300
	INHALATION [SLOPE] COMMENT BASED ON ROUTE TO ROUTE EXTRAPOLATION. SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST									
PARATHION	**000056-38-2**									
					IRIS					005116
	GENERAL COMMENT· ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
PENTABROMO-6-CHLOROCYCLOHEXANE, 1,2,3,4,5-	**000087-84-3**									
	ORAL DIET	2 YEARS RAT	INTESTINE, LARGE	TUMORS	C	2 3E-2		6.6E-7		010150
	ORAL [SLOPE] COMMENT: BASED ON RESULTS WITH THE ALPHA ISOMER.									
PENTACHLORONITROBENZENE	**000082-68-8**									
	ORAL	72 WEEKS MOUSE	LIVER	TUMORS	C	2.6E-1		7.4E-6		010151
	ORAL [SLOPE] COMMENT: UNDER REVIEW, NUMBER SUBJECT TO CHANGE.									
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
PENTACHLOROPHENOL		**000087-86-5**								
					IRIS	IRIS		IRIS		010381
	GENERAL COMMENT	ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)								
PHENYLENEDIAMINE, O-		**000095-54-5**								
	ORAL· DIET	548 DAYS			B2	4.7E-2		1 3E-6		010152
		RAT	LIVER	TUMORS						
	ORAL [SLOPE] COMMENT. BASED ON LIVER TUMORS IN RATS TREATED WITH O-PHENYLENEDIAMINE DIHYDROCHLORIDE.									
	GENERAL COMMENT· ALSO SEE HEAST TABLE 1. SUBCHRONIC AND CHRONIC TOXICITY.									
PHENYLPHENOL, 2-		**000090-43-7**								
	ORAL. DIET	637 DAYS			C	1.94E-3		5 5E-8		010153
		RAT	URINARY BLADDER	TUMORS						
POLYBROMINATED BIPHENYLS										
	ORAL GAVAGE	25 WEEKS			B2	8.9E+0		2 5E-4		010154
		RAT	LIVER	CARCINOMA						
			LIVER	NEOPLASTIC NODULE						
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)									
POLYCHLORINATED BIPHENYLS		**001336-36-3**								
					IRIS	IRIS		IRIS		005118
	GENERAL COMMENT. CARCINOGENICITY INFORMATION WAS CHANGED ON IRIS									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
PROPYLENE OXIDE		**000075-56-9**								
					IRIS	IRIS		IRIS		010156
	INHALATION. INTERMITTENT	2 YEARS MOUSE	NASAL CAVITY	TUMORS	IRIS		1 3E-2		IRIS	010155
	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).									
QUINOLINE		**000091-22-5**								
	ORAL· DIET	20-40 WEEKS RAT	LIVER	TUMORS	C	1.2E+1		3 5E-4		010158
RDX / (CYCLONITE)		**000121-82-4**								
					IRIS	IRIS		IRIS		010157
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1 SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
SELENIUM SULFIDE		**007446-34-6**								
					IRIS					010194
	ORAL [SLOPE] COMMENT STUDY RESULTS WERE CONSIDERED INCONCLUSIVE FOR QUANTITATIVE RISK ASSESSMENT									
SIMAZINE		**000122-34-9**								
	ORAL. DIET	2 YEARS RAT	MAMMARY	CARCINOMA	C	1 2E-1		3 4E-6		010195
	ORAL [SLOPE] COMMENT: UNDER REVIEW. NUMBER SUBJECT TO CHANGE									
	GENERAL COMMENT: ALSO SEE HEAST TABLE 1. SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
SODIUM DIETHYLDITHIOCARBAMATE		**000148-18-5**								
	ORAL: DIET	77 WEEKS			C	2.7E-1		7.7E-6		005126
		MOUSE	LIVER	TUMORS						
	GENERAL COMMENT	ALSO SEE HEAST TABLE 1: SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY)								
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 (PG/CU M)-1	005128
		RAT	RESPIRATORY SYSTEM	TUMORS						
			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	TUMOR						
	INHALATION [SLOPE] COMMENT	BASED ON ROUTE TO ROUTE EXTRAPOLATION SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST								

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
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)									
TETRACHLOROTOLUENE, PARA, ALPHA, ALPHA, ALPHA- 005216-25-1										
	ORAL· GAVAGE	17 5 WEEKS			B2	2.0E+1		5 7E-4		005028
		MOUSE	LUNG	ADENOCARCINOMA						
TETRACHLOROVINPHOS / (STIROFOS) 000961-11-5										
	ORAL: DIET	560 DAYS			C	2.4E-2		6.9E-7		010159
		MOUSE	LIVER	TUMORS						
	[EPA GROUP] COMMENT UNDER REVIEW. CLASSIFICATION SUBJECT TO CHANGE. GENERAL COMMENT: ALSO SEE HEAST TABLE 1· SUBCHRONIC AND CHRONIC TOXICITY (OTHER THAN CARCINOGENICITY).									
TOLUENE-2,4-DIAMINE 000095-80-7										
	ORAL. DIET	84-103 WEEKS			B2	3.2E+0		9.1E-5		010160
		RAT	MAMMARY	TUMORS						
TOLUIDINE, P- 000106-49-0										
	ORAL. DIET	18 MONTHS			C	1 9E-1		5 4E-6		010162
		MOUSE	LIVER	TUMORS						

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day) $^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day) $^{-1}$	[UNIT RISK] ORAL (ug/L) $^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
TOXAPHENE		**008001-35-2**								
	ORAL DIET	18 MONTHS MOUSE	LIVER	TUMORS	IRIS	IRIS	1 1E+0	IRIS	IRIS	005134
	INHALATION [SLOPE] COMMENT SEE APPENDIX A-II. DOSE CONVERSIONS ON HEAST INHALATION [UNIT RISK] COMMENT: BASED ON ROUTE TO ROUTE EXTRAPOLATION.									
TRICHLOROANILINE HYDROCHLORIDE, 2,4,6-				**033663-50-2**						
	ORAL. DIET	18 MONTHS MOUSE	VASCULAR SYSTEM	TUMORS	C	2 9E-2		8.2E-7		005142
TRICHLOROANILINE, 2,4,6-		**000634-93-5**								
	ORAL. DIET	18 MONTHS MOUSE	VASCULAR SYSTEM	TUMORS	C	3 4E-2		1E-6		010487
TRICHLOROETHANE, 1,1,2-		**000079-00-5**								
	ORAL GAVAGE	78 WEEKS MOUSE	LIVER	CARCINOMA	IRIS	IRIS	5.7E-2	IRIS	IRIS	005144
	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									

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
TRICHLOROPHENOL, 2,4,6-	**000088-06-2**									
	ORAL: DIET	107 WEEKS			IRIS	IRIS	1E-2	IRIS	IRIS	010428
		RAT	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		2E-4		010849
		RAT	MULTIPLE SITES	TUMORS, BENIGN/MALIGNANT, COMBINED						

[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		IRIS		010163

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

TRIMETHYL PHOSPHATE	**000512-56-1**									
	ORAL. GAVAGE	103 WEEKS			B2	3.7E-2		1 1E-6		010164
		MOUSE	UTERUS	TUMORS						

TRINITROTOLUENE, 2,4,6-	**000118-96-7**									
					IRIS	IRIS		IRIS		010476

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

CHEMICAL	ROUTE	EXPERIMENT LENGTH / SPECIES	TARGET	CANCER	[EPA GROUP]	[SLOPE FACTOR] ORAL (mg/kg/day)$^{-1}$	[SLOPE FACTOR] INHALATION (mg/kg/day)$^{-1}$	[UNIT RISK] ORAL (ug/L)$^{-1}$	[UNIT RISK] INHALATION (ug/cu m)-1	REFERENCE
VINYL CHLORIDE		**000075-01-4**								
	ORAL: DIET	1001 DAYS			A	1.9E+0		5.4E-5		010368
		RAT	LUNG	TUMORS						
			LIVER	TUMORS						
	INHALATION· INTERMITTENT	1 YEAR			A		3 0E-1		8 4E-5	010367
		RAT	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.
GENERAL COMMENT· THE MOST RECENTLY REVIEWED QUANTITATIVE TOXICITY VALUES LISTED HERE APPEAR IN EPA DOCUMENTS PUBLISHED IN 1984 AND 1985. USE OF THESE VALUES ON AN INTERIM BASIS WAS VALIDATED BY CRAVE (04/05/90). 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 TOXICITY VALUE INCLUDES 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. WASHINGTON. DC 20460

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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

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

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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QUAST, JF, CE WADE, CG HUMISTON ET AL 1980. A 2-YEAR TOXICITY AND ONCOGENICITY STUDY WITH ACRYLONITRILE INCORPORATED IN THE DRINKING WATER OF RATS PREPARED BY THE TOXICOLOGY RESEARCH LABORATORY, HEALTH AND ENVIRONMENTAL RESEARCH, DOW CHEMICAL USA, MIDLAND, MI FOR THE CHEMICAL MANUFACTURERS ASSOCIATION, WASHINGTON, DC.

US EPA 1983 HEALTH ASSESSMENT DOCUMENT FOR ACRYLONITRILE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, RESEARCH TRIANGLE PARK, NC

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US EPA 1987. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

005003 O'BERG, M 1980 EPIDEMIOLOGIC STUDY OF WORKERS EXPOSED TO ACRYLONITRILE J OCCUP MED. 22· 245-252

US EPA 1983 HEALTH ASSESSMENT DOCUMENT FOR ACRYLONITRILE. PREPARED BY THE OFFICE OF HEALTH AND ENVIRONMENTAL ASSESSMENT, ENVIRONMENTAL CRITERIA AND ASSESSMENT OFFICE, RESEARCH TRIANGLE PARK, NC.

US EPA 1987. HEALTH EFFECTS ASSESSMENT FOR ACRYLONITRILE. 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, 1987. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

ALACHLOR 015972-60-8

010180 STOUT, LD, ET AL. 1983 A CHRONIC STUDY OF ALACHLOR ADMINISTERED IN FEED TO LONG-EVANS RATS. EHL #800218, PROJECT NO. ML-80-186, REPORT MSL-3282/3284. VOL. I AND II UNPUBLISHED STUDY RECEIVED FEB 28, 1984 UNDER U S EPA REG NO. 524-316, PREPARED BY MONSANTO ENVIRONMENTAL HEALTH LABORATORY (EHL), SUBMITTED BY MONSANTO CO , WASHINGTON, DC CDL·252496-7.

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US EPA. 1992. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR ALACHLOR 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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ALDRIN 000309-00-2

005006 EPSTEIN, SS. 1975. THE CARCINOGENICITY OF DIELDRIN, PART I. SCI TOTAL ENVIRON 4: 1-52.

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US EPA 1987. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

ALLYL CHLORIDE 000107-05-1

010181 US EPA 1983. HEALTH AND ENVIRONMENTAL EFFECTS PROFILE FOR CHLOROPROPENES. 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 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

ANILINE **000062-53-3**

010088 CIIT (CHEMICAL INDUSTRY INSTITUTE OF TOXICOLOGY) 1982. 104-WEEK CHRONIC TOXICITY STUDY IN RATS. ANILINE HYDROCHLORIDE. FINAL REPORT

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US EPA. 1987. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

ARAMITE **000140-57-8**

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005081 US EPA. 1987. HEALTH AND ENVIRONMENTAL EFFECTS DOCUMENT FOR HALOETHERS. 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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CHLORONITROBENZENE, O- **000088-73-3**

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CHRYSENE **000218-01-9**

010185 REFER TO IRIS

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010191 REFER TO IRIS

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HEXACHLOROCYCLOHEXANE, BETA **000319-85-7**

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US EPA. 1986 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

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US EPA 1986 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

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US EPA 1986 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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US EPA. 1987 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

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INDENO[1,2,3-CD]PYRENE **000193-39-5**

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LEAD **007439-92-1**

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

LINURON **000330-55-2**

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US EPA 1989 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

MERCURIC CHLORIDE **007487-94-7**

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U.S EPA. 1994 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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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

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

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US EPA 1989. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

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US EPA 1987 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

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US EPA 1987 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

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010482 CONTACT THE SUPERFUND HEALTH RISK TECHNICAL SUPPORT CENTER (513) 569-7300

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US EPA 1986 CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP

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US EPA. 1994. CARCINOGEN RISK ASSESSMENT VERIFICATION ENDEAVOR (CRAVE) WORK GROUP.

Table 4

Radionuclide Carcinogencity -- Slope Factors (In Units of Picocuries)

JULY 1997

NOTE: To convert radionuclide slope factors into the International System (SI) activity units of becquerels (Bq), multiply each value in Table 4 by 27.03.

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Actinium (89)	Ac-225	014265-85-1	1 00E+01	D	Y	1.00E-03	1.42E-10	4.16E-09	7.81E-09
	Ac-227	014952-40-0	2.18E+01	Y	Y	1.00E-03	3.52E-10	7.08E-08	2.35E-11
	Ac-227+D	014952-40-0(+D)	2.18E+01	Y	Y	1.00E-03	6.26E-10	7.87E-08	5.97E-07
	Ac-228	014331-83-0	6.13E+00	H	Y	1 00E-03	1 62E-12	3.27E-11	3.28E-06
Americium (95)	Am-241	014596-10-2	4.32E+02	Y	W	1.00E-03	3.28E-10	3.85E-08	4 59E-09
	Am-242	013981-54-9	1.60E+01	H	W	1.00E-03	1.47E-12	1.04E-11	5 76E-09
	Am-242m	013981-54-9(m)	1.52E+02	Y	W	1.00E-03	2.92E-10	3 49E-08	8.76E-11
	Am-243	014993-75-0	7.38E+03	Y	W	1.00E-03	3.27E-10	3.82E-08	2.43E-08
	Am-243+D	014993-75-0(+D)	7.38E+03	Y	W	1.00E-03	3.31E-10	3.82E-08	2.66E-07
Antimony (51)	Sb-122	014374-79-9	2 70E+00	D	W	1 00E-01	8.81E-12	5.46E-12	1.61E-06
	Sb-124	014683-10-4	6.02E+01	D	W	1.00E-01	1.07E-11	1.32E-11	7.35E-06
	Sb-125	014234-35-6	2.77E+00	Y	W	1.00E-01	2.97E-12	5.20E-12	1.34E-06
	Sb-125+D	014234-35-6(+D)	2.77E+00	Y	W	1.00E-01	3.54E-12	5.85E-12	1.34E-06
	Sb-126	015756-32-8	1.24E+01	D	W	1.00E-01	9.73E-12	8.41E-12	1.03E-05
	Sb-126m	015756-32-8(m)	1.90E+01	M	W	1.00E-01	7.28E-14	6.43E-14	5.78E-06
	Sb-127	013968-50-8	3.85E+00	D	W	1.00E-01	8.48E-12	6.05E-12	2.40E-06
	Sb-129	014331-88-5	4 40E+00	H	W	1 00E-01	1.86E-12	8.60E-13	5 56E-06
Argon (18)	Ar-41	014163-25-8	1.83E+00	H	*	---	---	4.71E-16	---

[Table 4 continues on the following page: Refer to Endnotes on the last page]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Astatine (85)	At-217	017239-90-6	3.23E-02	S	D	9.50E-01	8.99E-18	5.14E-16	8.71E-10
Barium (56)	Ba-131	014914-75-1	1.18E+01	D	D	1.00E-01	1.70E-12	4.79E-13	1.27E-06
	Ba-133	013981-41-4	1.05E+01	Y	D	1.00E-01	2.70E-12	4.03E-12	9.15E-07
	Ba-133m	013981-41-4(m)	3.89E+01	H	D	1.00E-01	2.76E-12	5.60E-13	1.00E-07
	Ba-137m	013981-97-0(m)	2.55E+00	M	D	1.00E-01	2.43E-15	1.57E-15	2.21E-06
	Ba-139	014378-25-7	8.31E+01	M	D	1.00E-01	3.04E-13	1.53E-13	8.35E-08
	Ba-140	014798-08-4	1.28E+01	D	D	1.00E-01	1.18E-11	3.17E-12	6.00E-07
Beryllium (4)	Be-7	013966-02-4	5.34E+01	D	Y	5.00E-03	8.64E-14	1.78E-13	1.73E-07
Bismuth (83)	Bi-206	015776-19-9	6.24E+00	D	W	5.00E-02	7 11E-12	5.07E-12	1.20E-05
	Bi-207	013982-38-2	3.34E+01	Y	W	5.00E-02	5.05E-12	9.42E-12	5.49E-06
	Bi-210	014331-79-4	5.01E+00	D	W	5.00E-02	7.29E-12	5 12E-11	0.00E+0
	Bi-211	015229-37-5	2.13E+00	M	W	5.00E-02	1.82E-14	1.74E-12	1.48E-07
	Bi-212	014913-49-6	6 06E+01	M	W	5 00E-02	6.20E-13	3.65E-11	6.67E-07
	Bi-213	015776-20-2	4.57E+01	M	W	5.00E-02	4.40E-13	3.09E-11	4 62E-07
	Bi-214	014733-03-0	1.99E+01	M	W	5.00E-02	1.95E-13	1 46E-11	6 02E-06
Bromine (35)	Br-82	014686-69-2	3.53E+01	H	D	9.50E-01	1.42E-12	7.86E-13	1 01E-05
Cadmium (20)	Cd-109	014109-32-1	4 64E+02	D	Y	5.00E-02	8.01E-12	1.85E-11	5.62E-10
	Cd-115	014336-68-6	5.35E+01	H	Y	5 00E-02	7.29E-12	4.93E-12	7.02E-07

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
	Cd-115m	014336-68-6(m)	4 46E+01	D	Y	5.00E-02	1.42E-11	1.70E-11	8.55E-08
Calcium (20)	Ca-45	013966-05-7	1.63E+02	D	W	3.00E-01	2.02E-12	2.51E-12	3.88E-18
	Ca-47	001439-99-2	4.54E+00	D	W	3.00E-01	6.66E-12	5.22E-12	4.12E-06
Carbon (6)	C-11	014333-33-6	2.05E+01	M	D	9.50E-01	4.49E-14	3.38E-14	3.61E-06
	C-14	014762-75-5	5.73E+03	Y	*	1.00E+00	1.03E-12	6.99E-15	0.00E+0
	C-15	015929-23-4	2.45E+00	S	D	9.50E-01	6.62E-16	8.06E-16	---
Cerium (58)	Ce-141	013967-74-3	3.25E+01	D	Y	3.00E-04	3.91E-12	4.32E-12	1.41E-07
	Ce-143	014119-19-8	3.30E+01	H	Y	3.00E-04	5.91E-12	3.84E-12	7.32E-07
	Ce-144	014762-78-8	2.84E+02	D	Y	3.00E-04	2.96E-11	1.08E-10	2 58E-08
	Ce-144+D	014762-78-8(+D)	2.84E+02	D	Y	3.00E-04	2.97E-11	1.08E-10	1.56E-07
Cesium (55)	Cs-131	014914-76-2	9.69E+00	D	D	9.50E-01	1.80E-13	1 06E-13	2.34E-09
	Cs-134	013967-70-9	2.06E+00	Y	D	9.50E-01	4.73E-11	2.89E-11	5.88E-06
	Cs-134m	013967-70-9(m)	2.90E+00	H	D	9.50E-01	4.54E-14	3.10E-14	1.96E-08
	Cs-135	015726-30-4	2.30E+06	Y	D	9.50E-01	4.53E-12	2.71E-12	0.00E+0
	Cs-136	014234-29-8	1.32E+01	D	D	9 50E-01	7 74E-12	4.65E-12	8.13E-06
	Cs-137	010045-97-3	3.02E+01	Y	D	9.50E-01	3.16E-11	1.91E-11	0.00E+0
	Cs-137+D	010045-97-3(+D)	3.02E+01	Y	D	9.50E-01	3.16E-11	1.91E-11	2.09E-06
	Cs-138	015758-29-9	3.22E+01	M	D	9.50E-01	1.76E-13	1 30E-13	9.45E-06

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Chlorine (17)	Cl-36	013981-43-6	3.01E+05	Y	D	9.50E-01	2.23E-12	1.30E-12	0.00E+0
	Cl-38	014158-34-0	3.72E+01	M	D	9.50E-01	2.07E-13	1.63E-13	6 47E-06
Chromium (24)	Cr-51	014392-02-0	2.77E+01	D	Y	1.00E-01	1.38E-13	1.74E-13	1.02E-07
Cobalt (27)	Co-57	013981-50-5	2.71E+02	D	Y	3.00E-01	9.71E-13	2.88E-12	2.07E-07
	Co-58	01381-38-9	7.08E+01	D	Y	3.00E-01	2.82E-12	5.17E-12	3.73E-06
	Co-58m	01381-38-9(m)	9.15E+00	H	Y	3.00E-01	9.46E-14	8.90E-14	3.21E-11
	Co-60	010198-40-0	5.27E+00	Y	Y	3.00E-01	1.89E-11	6.88E-11	9.76E-06
Copper (29)	Cu-64	013981-25-4	1.27E+01	H	Y	5.00E-01	5.25E-13	4.18E-13	6.72E-07
Curium (96)	Cm-242	015510-73-3	1.63E+02	D	W	1.00E-03	3.83E-11	3.16E-09	2.34E-11
	Cm-243	015757-87-6	2.85E+01	Y	W	1.00E-03	2.51E-10	2.89E-08	1.71E-07
	Cm-243+D	015757-87-6(+D)	2.85E+01	Y	W	1.00E-03	2.52E-10	2.90E-08	1.72E-07
	Cm-244	013981-15-2	1.81E+01	Y	W	1.00E-03	2.11E-10	2.43E-08	2.07E-11
	Cm-245	015621-76-8	8.50E+03	Y	W	1.00E-03	3.35E-10	3.92E-08	5.51E-08
	Cm-246	015757-90-1	4.75E+03	Y	W	1.00E-03	3.32E-10	3.90E-08	1.81E-11
	Cm-247	015758-32-4	1.56E+07	Y	W	1.00E-03	3 09E-10	3.58E-08	1.03E-06
	Cm-248	015758-33-5	3.39E+05	Y	W	1 00E-03	1 31E-09	1.46E-07	1.47E-11
Dysprosium (66)	Dy-165	013967-64-1	2.33E+00	H	W	3.00E-04	3.26E-13	2.24E-13	6.18E-08
	Dy-166	015840-01-4	8.16E+01	H	W	3.00E-04	9.42E-12	7.82E-12	2 72E-08

[Table 4 continues on the following page· Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Erbium (63)	Er-169	015840-13-8	9.40E+00	D	W	3.00E-04	2.12E-12	1.51E-12	6.52E-12
	Er-171	014391-45-8	7.52E+00	H	W	3.00E-04	1.63E-12	7.50E-13	1.04E-06
Europium (63)	Eu-152	014683-23-9	1.36E+01	Y	W	1.00E-03	5.73E-12	7.91E-11	4.08E-06
	Eu-154	015585-10-1	8.80E+00	Y	W	1.00E-03	9.37E-12	9.15E-11	4.65E-06
	Eu-155	014391-16-3	4.96E+00	Y	W	1.00E-03	1.65E-12	9.60E-12	6.08E-08
	Eu-156	014280-35-4	1 52E+01	D	W	1.00E-03	1.09E-11	9.26E-12	5.40E-06
Fluorine (9)	F-18	013981-56-1	1.10E+02	M	D	9.50E-01	1.09E-13	6.54E-14	3.50E-06
Francium (87)	Fr-221	015756-41-9	4.80E+00	M	D	9.50E-01	1.45E-13	8.02E-12	6.74E-08
	Fr-223	015756-98-6	2.18E+00	M	D	9.50E-01	4.46E-13	5.90E-13	4.17E-08
Gadolinium (64)	Gd-153	014276-65-4	2.42E+02	D	W	3.00E-04	1.32E-12	3.20E-12	7.22E-08
	Gd-159	014041-42-0	1.86E+01	H	W	3.00E-04	2 60E-12	1.24E-12	9.59E-08
Gallium (31)	Ga-67	014119-09-6	3.26E+00	D	W	1.00E-03	8.36E-13	5.14E-13	3.61E-07
	Ga-72	013982-22-4	1.41E+01	H	W	1.00E-03	4.77E-12	2.17E-12	1.12E-05
Germanium (32)	Ge-71	014374-81-3	1.18E+01	D	W	9.50E-01	1 18E-14	5.84E-14	1.56E-11
Gold (79)	Au-196	014914-16-0	6.18E+00	D	Y	1.00E-01	1.30E-12	1.04E-12	1.41E-06
	Au-198	010043-49-0	2.70E+00	D	Y	1.00E-01	5.28E-12	3.64E-12	1.37E-06
Holmium (67)	Ho-166	013967-65-2	2.68E+01	H	W	3.00E-04	7.57E-12	4.06E-12	6 96E-08
Hydrogen (1)	H-3	010028-17-8	1.23E+01	Y	V	1.00E+00	7.15E-14	9.59E-14	0.00E+0

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Indium (49)	In-113m	014885-78-0(m)	1.66E+00	H	W	2.00E-02	8.30E-14	5.77E-14	7.82E-07
	In-114	013981-55-0	7.19E+01	S	W	2.00E-02	4.53E-15	5.81E-15	1.13E-07
	In-114m	013981-55-0(m)	4.95E+01	D	W	2.00E-02	2.06E-11	2.53E-11	2.00E-07
	In-115	014191-71-0	4.60e+15	Y	W	2.00E-02	3.49E-11	2.07E-10	0.00E+0
	In-115m	014191-71-0(m)	4.36E+00	H	W	2.00E-02	3.42E-13	1.75E-13	4.29E-07
Iodine (53)	I-122	018287-75-7	3.62E+00	M	D	9.50E-01	2.16E-14	2.24E-14	3.41E-06
	I-123	015715-08-9	1.31E+01	H	D	9.50E-01	5.42E-13	2.94E-13	2.52E-07
	I-125	014158-31-7	6.01E+01	D	D	9.50E-01	2.58E-11	1.71E-11	2.39E-09
	I-126	014158-32-8	1.29E+01	D	D	9.50E-01	4.82E-11	3.15E-11	1.49E-06
	I-129	015046-84-1	1.57E+07	Y	D	9.50E-01	1.84E-10	1.22E-10	2.69E-09
	I-130	014914-02-4	1.24E+01	H	D	9.50E-01	4.85E-12	2.61E-12	7.93E-06
	I-131	010043-66-0	8.04E+00	D	D	9.50E-01	3.62E-11	2.33E-11	1.25E-06
	I-132	014683-16-0	2.30E+00	H	D	9.50E-01	6.62E-13	3 52E-13	8.75E-06
	I-133	014834-67-4	2.08E+01	H	D	9.50E-01	1.06E-11	6.02E-12	2.20E-06
	I-134	014914-27-3	5.26E+01	M	D	9.50E-01	2 31E-13	1 38E-13	1.02E-05
	I-135	014834-68-5	6.61E+00	H	D	9.50E-01	2.27E-12	1.18E-12	6.23E-06
Iridium (77)	Ir-190	014981-91-0	1.18E+01	D	Y	1.00E-02	4 95E-12	4.49E-12	4.65E-06
	Ir-194	014158-35-1	1.92E+01	H	Y	1.00E-02	7.00E-12	4.18E-12	3.17E-07

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Iron (26)	Fe-55	014681-59-5	2 70E+00	Y	W	1.00E-01	3.51E-13	5 60E-13	0.00E+0
	Fe-59	014596-12-4	4.46E+01	D	W	1.00E-01	5.87E-12	7.08E-12	4.63E-06
Krypton (36)	Kr-83m	013965-98-5(m)	1.83E+00	H	*	---	---	3.48E-17	---
	Kr-85	013983-27-2	1.07E+01	Y	*	---	---	2.87E-16	---
	Kr-85m	013983-27-2(m)	4.48E+00	H	*	---	---	2.75E-16	---
	Kr-87	014809-68-8	7.63E+01	M	*	---	---	1.20E-15	---
	Kr-88	014995-61-0	2.84E+00	H	*	---	---	2.20E-15	---
	Kr-89	016316-03-3	3.16E+00	M	*	---	---	1.61E-15	---
	Kr-90	015741-13-6	3.23E+01	S	*	---	---	1.60E-15	---
Lanthanum (57)	La-140	013981-28-7	4.02E+01	H	W	1.00E-03	9.46E-12	5 10E-12	9.11E-06
Lead (82)	Pb-203	014687-25-3	5.20E+01	H	D	2.00E-01	1.03E-12	3.10E-13	6 40E-07
	Pb-209	014119-30-3	3 25E+00	H	D	2.00E-01	2.09E-13	6.85E-14	0.00E+0
	Pb-210	014255-04-0	2.23E+01	Y	D	2.00E-01	6.75E-10	1.67E-09	1.12E-10
	Pb-210+D	014255-04-0(+D)	2.23E+01	Y	D	2.00E-01	1.01E-09	3.86E-09	1.45E-10
	Pb-211	015816-77-0	3.61E+01	M	D	2.00E-01	3.38E-13	1.03E-11	1.85E-07
	Pb-212	015092-94-1	1.06E+01	H	D	2.00E-01	1.80E-11	3.85E-11	3.00E-07
	Pb-214	015067-28-4	2.68E+01	M	D	2.00E-01	2.94E-13	6.23E-12	7.09E-07
Lutetium (71)	Lu-177	014265-75-9	6.71E+00	D	Y	3.00E-04	2.95E-12	2 20E-12	7.22E-08

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] JULY 1997
(In Units of Picocuries[b])

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Manganese (25)	Mn-52	014092-99-0	5.59E+00	D	W	1.00E-01	6.01E-12	4.40E-12	1.34E-05
	Mn-54	013966-31-9	3.13E+02	D	W	1.00E-01	1.96E-12	3.69E-12	3.26E-06
	Mn-56	014681-52-8	2.58E+00	H	W	1.00E-01	8.57E-13	5.21E-13	6.95E-06
Mercury (80)	Hg-197	013981-51-6	6.41E+01	H	W	2.00E-02	1.18E-12	6.95E-13	5.47E-08
	Hg-203	013982-78-0	4.66E+01	D	W	2.00E-02	2.64E-12	3.03E-12	6.27E-07
Molybdenum (42)	Mo-99	014119-15-4	6.60E+01	H	Y	8.00E-01	2.27E-12	4.48E-12	5.46E-07
Neodymium (60)	Nd-147	014269-74-0	1.10E+01	D	Y	3.00E-04	5.88E-12	4.84E-12	3.22E-07
	Nd-149	015749-81-2	1.73E+00	H	Y	3.00E-04	4.55E-13	4.22E-13	1.08E-06
Neptunium (93)	Np-236	015700-36-4	1.15E+05	Y	W	1.00E-03	9.31E-13	3.87E-12	9.22E-08
	Np-237	013994-20-2	2.14E+06	Y	W	1.00E-03	2.95E-10	3.45E-08	7.56E-09
	Np-237+D	013994-20-2(+D)	2.14E+06	Y	W	1.00E-03	3.00E-10	3.45E-08	4.62E-07
	Np-238	015766-25-3	2.12E+00	D	W	1.00E-03	4.56E-12	4.68E-12	1.95E-06
	Np-239	013968-59-7	2.36E+00	D	W	1.00E-03	4.27E-12	2.41E-12	2.42E-07
	Np-240	015690-84-3	6.50E+01	M	W	1.00E-03	1.77E-13	1.31E-13	3.65E-06
	Np-240m	015690-84-3(m)	7.40E+00	M	W	1.00E-03	2.42E-14	2.83E-14	1.05E-06
Nickel (28)	Ni-59	014336-70-0	7.50E+04	Y	W	5.00E-02	1.85E-13	4.01E-13	0.00E+0
	Ni-63	013981-37-8	1.00E+02	Y	W	5.00E-02	5.50E-13	1.01E-12	0.00E+0
	Ni-65	014833-49-9	2.52E+00	H	W	5.00E-02	5.62E-13	3.59E-13	2.14E-06

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Niobium (41)	Nb-93m	007440-03-1(m)	1.46E+01	Y	Y	1.00E-02	6.64E-13	4.33E-12	3.64E-11
	Nb-94	014681-63-1	2.03E+04	Y	Y	1 00E-02	6.91E-12	8.20E-11	6.08E-06
	Nb-95	013967-76-5	3.51E+01	D	Y	1.00E-02	2.25E-12	3.11E-12	2.94E-06
	Nb-95m	013967-76-5(m)	8.66E+01	H	Y	1.00E-02	3.06E-12	2.25E-12	8.71E-08
	Nb-97	018496-04-3	7.21E+01	M	Y	1.00E-02	1.75E-13	2.13E-13	2.49E-06
	Nb-97m	018496-04-3(m)	6.00E+01	S	Y	1.00E-02	3.27E-15	3.34E-15	2.78E-06
Osmium (76)	Os-185	015766-50-4	9.36E+01	D	Y	1.00E-02	1 80E-12	4.62E-12	2.45E-06
	Os-191	014119-24-5	1.54E+01	D	Y	1.00E-02	3.04E-12	2.70E-12	8.74E-08
	Os-191m	014119-24-5(m)	1.30E+01	H	Y	1.00E-02	4.95E-13	3.32E-13	3.22E-09
	Os-193	016057-77-5	3.00E+01	H	Y	1.00E-02	4.36E-12	2.68E-12	1.82E-07
Palladium (46)	Pd-100	015690-69-4	3.64E+00	D	Y	5.00E-03	3.74E-12	3.55E-12	---
	Pd-101	015749-54-9	8.48E+00	H	Y	5.00E-03	3.74E-13	2.29E-13	---
	Pd-103	014967-68-1	1.70E+01	D	Y	5.00E-03	1.05E-12	1.08E-12	5.38E-10
	Pd-107	017637-99-9	6.50E+06	Y	Y	5.00E-03	2.09E-13	1.46E-12	0.00E+0
	Pd-109	014981-64-7	1.35E+01	H	Y	5.00E-03	3.33E-12	1.99E-12	2 43E-09
Phosphorus (15)	P-32	014596-37-3	1.43E+01	D	D	8.00E-01	6.11E-12	2.93E-12	0.00E+0
	P-33	015749-66-3	2.54E+01	D	D	8.00E-01	7.81E-13	3.96E-13	0 00E+0
Platinum (78)	Pt-191	015706-36-2	2.71E+00	D	D	1.00E-02	1.50E-12	4.13E-13	6.74E-07

[Table 4 continues on the following page· Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
	Pt-193	015735-70-3	5.00E+01	Y	D	1.00E-02	1.62E-13	7.89E-14	0.00E+0
	Pt-193m	015735-70-3(m)	4.33E+00	D	D	1.00E-02	2.51E-12	5.76E-13	7.44E-09
	Pt-197	015735-74-7	1.83E+01	H	D	1 00E-02	2.12E-12	4.54E-13	3.15E-08
	Pt-197m	015735-74-7(m)	9.44E+01	M	D	1.00E-02	3.25E-13	1.00E-13	1.65E-07
Plutonium (94)	Pu-236	015411-92-4	2 85E+00	Y	Y	1.00E-03	7.68E-11	1.34E-08	2 32E-11
	Pu-238	013981-16-3	8 78E+01	Y	Y	1.00E-03	2.95E-10	2.74E-08	1.94E-11
	Pu-239	015117-48-3	2.41E+04	Y	Y	1.00E-03	3.16E-10	2.78E-08	1.26E-11
	Pu-240	014119-33-6	6.57E+03	Y	Y	1.00E-03	3.15E-10	2.78E-08	1.87E-11
	Pu-241	014119-32-5	1.44E+01	Y	Y	1.00E-03	5.20E-12	2.81E-10	0.00E+0
	Pu-241+D	014119-32-5(+D)	1.44E+01	Y	Y	1.00E-03	3.33E-10	3.88E-08	4.59E-09
	Pu-242	013982-10-0	3.76E+05	Y	Y	1.00E-03	3.00E-10	2.64E-08	1.55E-11
	Pu-243	015706-37-3	4.96E+00	H	Y	1 00E-03	3 69E-13	2.67E-13	1.89E-08
	Pu-244	014119-34-7	8.26E+07	Y	Y	1 00E-03	3 13E-10	2.67E-08	1.29E-11
	Pu-244+D	014119-34-7(+D)	8.26E+07	Y	Y	1 00E-03	3.19E-10	2.67E-08	3.65E-06
Polonium (84)	Po-210	013981-52-7	1.38E+02	D	W	1.00E-01	3.26E-10	2.14E-09	3 30E-11
	Po-212	015389-34-1	2.98E-07	S	W	1.00E-01	4.51E-23	5.93E-21	0 00E+0
	Po-213	015756-57-7	4.20E-06	S	W	1.00E-01	6.70E-22	7.80E-20	1.18E-10
	Po-214	015735-67-8	1.64E-04	S	W	1.00E-01	2.12E-20	2.77E-18	3 23E-10

[Table 4 continues on the following page. Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
	Po-215	015706-52-2	1 78E-03	S	W	1.00E-01	4.99E-19	4.48E-17	5.11E-10
	Po-216	015756-58-8	1.46E-01	S	W	1.00E-01	8.79E-17	2 95E-15	5.62E-11
	Po-218	015422-24-9	3 05E+00	M	W	1.00E-01	5.08E-14	3.69E-12	0 00E+0
Potassium (19)	K-40	013966-00-2	1.28E+09	Y	D	9 50E-01	1.25E-11	7.46E-12	6.11E-07
	K-42	014378-21-3	1.24E+01	H	D	9.50E-01	1.29E-12	7.56E-13	1.09E-06
Praseodymium (59)	Pr-142	014191-64-1	1.91E+01	H	Y	3.00E-04	6.98E-12	4.16E-12	2.34E-07
	Pr-143	014981-79-4	1.36E+01	D	Y	3.00E-04	6.60E-12	5.60E-12	3.41E-14
	Pr-144	014119-05-2	1.73E+01	M	Y	3.00E-04	8.08E-14	1.31E-13	1.33E-07
	Pr-144m	014119-05-2(m)	7.20E+00	M	Y	3.00E-04	3 23E-14	5.61E-14	1.85E-09
Promethium (61)	Pm-147	014380-75-7	2.62E+00	Y	Y	3.00E-04	1.41E-12	7.49E-12	6.35E-12
	Pm-148	014683-19-3	5.37E+00	D	Y	3.00E-04	1.44E-11	1.05E-11	2 21E-06
	Pm-148m	014683-19-3(m)	4.13E+01	D	Y	3.00E-04	9.93E-12	2 95E-11	7.32E-06
	Pm-149	015765-31-8	5.31E+01	H	Y	3.00E-04	5.52E-12	3.57E-12	3.65E-08
Protactinium (91)	Pa-231	014331-85-2	3.73E+04	Y	Y	1.00E-03	1.49E-10	2 42E-08	2.71E-08
	Pa-233	013981-14-1	2.70E+01	D	Y	1.00E-03	4.69E-12	4.92E-12	4.54E-07
	Pa-234	015100-28-4	6.70E+00	H	Y	1.00E-03	2.13E-12	1.30E-12	6.60E-06
	Pa-234m	015100-28-4(m)	1.17E+00	M	Y	1.00E-03	4.77E-15	6.27E-15	4.05E-08
Radium (88)	Ra-223	015623-45-7	1.14E+01	D	W	2.00E-01	2.34E-10	3.60E-09	2.44E-07

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
	Ra-224	013233-32-4	3.62E+00	D	W	2.00E-01	1 49E-10	2.25E-09	2.48E-08
	Ra-225	013981-53-8	1.48E+01	D	W	2.00E-01	1 57E-10	2.38E-09	1.71E-09
	Ra-226	013982-63-3	1.60E+03	Y	W	2.00E-01	2.95E-10	2.72E-09	1.31E-08
	Ra-226+D	013982-63-3(+D)	1.60E+03	Y	W	2.00E-01	2 96E-10	2.75E-09	6.74E-06
	Ra-228	015262-20-1	5.75E+00	Y	W	2.00E-01	2.46E-10	9.61E-10	0.00E+0
	Ra-228+D	015262-20-1(+D)	5.75E+00	Y	W	2.00E-01	2.48E-10	9.94E-10	3.28E-06
Radon (86)	Rn-219	014835-02-0	3.96E+00	S	*	---	---	6.91E-14	1.72E-07
	Rn-220	022481-48-7	5.56E+01	S	*	---	---	1.92E-13	1.88E-09
	Rn-222+D[1]	014859-67-7(+D)	3.82E+00	D	*	---	---	7.57E-12	---
Rhodium (45)	Rh-103m	007440-16-6(m)	5.61E+01	M	Y	5.00E-02	8.19E-15	1.28E-14	5.85E-11
	Rh-105	014913-89-4	3.54E+01	H	Y	5.00E-02	1.93E-12	1.22E-12	2.49E-07
	Rh-105m	014913-89-4(m)	4.50E+01	S	Y	5.00E-02	1.08E-15	9 25E-16	2.27E-08
	Rh-106	014234-34-5	2.99E+01	S	Y	5.00E-02	3.63E-15	4.62E-15	7.57E-07

[1] To derive the inhalation slope factor for Rn-222+D, EPA's Office of Radiation and Indoor Air (ORIA) uses a slightly different risk model and set of exposure assumptions, including an inhalation rate of 2.2E+04 L/day; 50% equilibrium for decay products; and a risk coefficient of 2.36E-4 cases per working level month (WLM). A more detailed description of ORIA's radon risk assessment methodology is provided in the EPA CRAVE Summary Sheet, *Inhaled Rn-222 and its Short Half-Life Decay Products*.

[Table 4 continues on the following page· Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Rubidium (37)	Rb-82	014391-63-0	1.25E+00	M	D	9.50E-01	1.05E-14	1.17E-14	3.89E-06
	Rb-86	014932-53-7	1.87E+01	D	D	9.50E-01	7.12E-12	4.21E-12	3.71E-07
	Rb-87	013982-13-3	4.73e+10	Y	D	9.50E-01	3.68E-12	2.26E-12	0.00E+0
	Rb-88	014928-36-0	1.78E+01	M	D	9.50E-01	1.46E-13	1.36E-13	2.68E-06
	Rb-89	014191-65-2	1.54E+01	M	D	9.50E-01	8.65E-14	6.92E-14	8.47E-06
Ruthenium (44)	Ru-97	015758-35-7	2.90E+00	D	Y	5.00E-02	5.88E-13	4.09E-13	4.52E-07
	Ru-103	013968-53-1	3.94E+01	D	Y	5.00E-02	3.32E-12	4.59E-12	1.70E-06
	Ru-105	014331-95-4	4.44E+00	H	Y	5.00E-02	1.15E-12	8.02E-13	2.88E-06
	Ru-106	013967-48-1	3.68E+02	D	Y	5.00E-02	3.45E-11	1.15E-10	0.00E+0
	Ru-106+D	013967-48-1(+D)	3.68E+02	D	Y	5.00E-02	3.45E-11	1.15E-10	7.57E-07
Samarium (62)	Sm-147	014392-33-7	1.06e+11	Y	W	3.00E-04	2.51E-11	6.93E-09	0.00E+0
	Sm-151	015715-94-3	9.00E+01	Y	W	3.00E-04	4.60E-13	4.63E-12	2.92E-13
	Sm-153	015766-00-4	4.67E+01	H	W	3.00E-04	4.02E-12	2.18E-12	4.65E-08
Scandium (21)	Sc-46	013967-63-0	8.38E+01	D	Y	1.00E-04	5.73E-12	1.31E-11	7.89E-06
	Sc-47	014391-96-9	3.42E+00	D	Y	1.00E-04	2.95E-12	2.01E-12	2.50E-07
	Sc-48	014391-86-7	4.37E+01	H	Y	1.00E-04	6.65E-12	4.20E-12	1.31E-05
Selenium (34)	Se-75	014265-71-5	1.20E+02	D	W	8.00E-01	6.53E-12	4.92E-12	8.89E-07
Silicon (14)	Si-31	014276-49-4	1.57E+02	M	W	1.00E-02	5.04E-13	3.29E-13	3.45E-09

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Silver (47)	Ag-105	014928-14-4	4.13E+01	D	Y	5.00E-02	1.63E-12	2.33E-12	---
	Ag-108	014391-65-2	2.37E+00	M	Y	5.00E-02	6.94E-15	9.43E-15	5.78E-08
	Ag-108m	014391-65-2m	1.27E+02	Y	Y	5.00E-02	6.05E-12	7.02E-11	5.61E-06
	Ag-108m+D	014391-65-2m(+D)	1.27E+02	Y	Y	5.00E-02	6.05E-12	7.02E-11	5.62E-06
	Ag-109m	014378-38-2(m)	3.96E+01	S	Y	5.00E-02	2.71E-16	3.46E-16	1.16E-09
	Ag-110	014391-76-5	2.46E+01	S	Y	5.00E-02	2.44E-15	3.16E-15	1.13E-07
	Ag-110m	014391-76-5(m)	2.50E+02	D	Y	5.00E-02	8.43E-12	3.21E-11	1.05E-05
	Ag-111	157690-04-0	7.46E+00	D	Y	5.00E-02	6.83E-12	5.24E-12	8.51E-08
Sodium (11)	Na-22	013966-32-0	2.60E+00	Y	D	9.50E-01	8.02E-12	4.88E-12	8.18E-06
	Na-24	013982-04-2	1.50E+01	H	D	9.50E-01	1.38E-12	7.51E-13	1.77E-05
Strontium (38)	Sr-82	014809-50-8	2.50E+01	D	D	3.00E-01	2.58E-11	8.87E-12	9.00E-11
	Sr-85	013967-73-2	6.48E+01	D	D	3.00E-01	1.40E-12	1.14E-12	1.54E-06
	Sr-85m	013967-73-2(m)	6.77E+01	M	D	3.00E-01	1.80E-14	7.13E-15	5.24E-07
	Sr-89	014158-27-1	5.06E+01	D	D	3.00E-01	1.03E-11	3.68E-12	5.38E-10
	Sr-90	010098-97-2	2.86E+01	Y	D	3.00E-01	4.09E-11	5.94E-11	0.00E+0
	Sr-90+D	010098-97-2(+D)	2.86E+01	Y	D	3.00E-01	5.59E-11	6.93E-11	0.00E+0
	Sr-91	014331-91-0	9.50E+00	H	D	3.00E-01	2.82E-12	7.79E-13	2.67E-06
	Sr-92	014928-29-1	2.71E+00	H	D	3.00E-01	2.03E-12	4.70E-13	5.20E-06

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Sulfur (16)	S-35	015117-53-0	8.74E+01	D	D	8.00E-01	4.16E-13	1.85E-13	0.00E+0
Tantalum (73)	Ta-182	013982-00-8	1.15E+02	D	Y	1.00E-03	7.03E-12	1.65E-11	4.66E-06
Technetium (43)	Tc-95	014809-56-4	2.00E+01	H	W	8.00E-01	6.81E-14	3.38E-14	2.72E-06
	Tc-95m	014809-56-4(m)	6.10E+01	D	W	8.00E-01	1.24E-12	2.10E-12	2.08E-06
	Tc-96	014808-44-7	4.28E+00	D	W	8.00E-01	2.28E-12	1.94E-12	9.36E-06
	Tc-96m	014808-44-7(m)	5.15E+01	M	W	8.00E-01	2.61E-14	2.26E-14	7.72E-08
	Tc-97	015759-35-0	2.60E+06	Y	W	8.00E-01	1.58E-13	3.44E-13	2.49E-10
	Tc-97m	015759-35-0(m)	8.90E+01	D	W	8.00E-01	1.20E-12	1.96E-12	2.67E-10
	Tc-99	014133-76-7	2.13E+05	Y	W	8.00E-01	1.40E-12	2.89E-12	6.19E-13
	Tc-99m	014133-76-7(m)	6.02E+00	H	W	8.00E-01	5.58E-14	3.49E-14	2.51E-07
Tellurium (52)	Te-125m	014390-73-9(m)	5.80E+01	D	W	2.00E-01	2.51E-12	2.85E-12	2.16E-09
	Te-127	013981-49-2	9.35E+00	H	W	2.00E-01	8.55E-13	4.32E-13	1.62E-08
	Te-127m	013981-49-2(m)	1.09E+02	D	W	2.00E-01	6.01E-12	1.31E-11	7.10E-10
	Te-129	014269-71-7	6.96E+01	M	W	2.00E-01	1.48E-13	1.46E-13	1.46E-07
	Te-129m	014269-71-7(m)	3.36E+01	D	W	2.00E-01	1.17E-11	1.33E-11	6.92E-08
	Te-131	014683-12-6	2.50E+01	M	W	2.00E-01	3.90E-13	2.48E-13	1.35E-06
	Te-131m	014683-12-6(m)	3.00E+01	H	W	2.00E-01	8.81E-12	8.40E-12	5.31E-06
	Te-132	014234-28-7	7.82E+01	H	W	2.00E-01	1.22E-11	8.38E-12	4.31E-07

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Terbium (65)	Tb-158	015759-55-4	1.50E+02	Y	W	3.00E-04	4.20E-12	7.04E-11	---
	Tb-160	013981-29-8	7.23E+01	D	W	3.00E-04	7.62E-12	1.14E-11	4.03E-06
Thallium (81)	Tl-202	015720-57-7	1.22E+01	D	D	9 50E-01	1.01E-12	6.07E-13	1.42E-06
	Tl-204	013968-51-9	3.78E+00	Y	D	9.50E-01	1.97E-12	1.15E-12	8 72E-10
	Tl-208	014913-50-9	3.05E+00	M	D	9.50E-01	1.75E-14	1.36E-14	1.45E-05
	Tl-209	015690-73-0	2.20E+00	M	D	9.50E-01	1.40E-14	1.12E-14	7.83E-06
Thorium (90)	Th-227	015623-47-9	1.87E+01	D	Y	2 00E-04	4.04E-11	4.31E-09	1.74E-07
	Th-228	014274-82-9	1.91E+00	Y	Y	2 00E-04	6 29E-11	9.45E-08	5.28E-10
	Th-228+D	014274-82-9(+D)	1.91E+00	Y	Y	2.00E-04	2 31E-10	9.68E-08	9.94E-07
	Th-229	015594-54-4	7.34E+03	Y	Y	2.00E-04	5.65E-11	7.60E-08	5.94E-08
	Th-229+D	015594-54-4(+D)	7.34E+03	Y	Y	2.00E-04	3.56E-10	8.26E-08	5.99E-07
	Th-230	014269-63-7	7.70E+04	Y	Y	2.00E-04	3.75E-11	1.72E-08	4 40E-11
	Th-231	014932-40-2	2.55E+01	H	Y	2 00E-04	1.79E-12	1.10E-12	2.09E-09
	Th-232	007440-29-1	1 41E+10	Y	Y	2.00E-04	3.28E-11	1.93E-08	1.97E-11
	Th-234	015065-10-8	2 41E+01	D	Y	2.00E-04	1.93E-11	1.90E-11	3.50E-09
Thulium (69)	Tm-170	013981-30-1	1.29E+02	D	W	3.00E-04	7.50E-12	1.10E-11	3 84E-09
	Tm-171	014333-45-0	1.92E+00	Y	W	3.00E-04	5.86E-13	1.84E-12	3 15E-10
Tin (50)	Sn-113	013966-06-8	1 15E+02	D	W	2.00E-02	3.72E-12	6.61E-12	2.96E-09

[Table 4 continues on the following page· Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
	Sn-121	014683-06-8	2.71E+01	H	W	2.00E-02	1.22E-12	6.13E-13	---
	Sn-121m	014683-06-8(m)	5.55E+01	Y	W	2.00E-02	2.00E-12	7.46E-12	---
	Sn-125	014683-08-0	9.64E+00	D	W	2 00E-02	1.68E-11	1 19E-11	1.21E-06
	Sn-126	015832-50-5	1.00E+05	Y	W	2.00E-02	2.12E-11	4.26E-11	3.32E-08
Tungsten (74)	W-181	015749-46-9	1.21E+02	D	D	3.00E-01	2.72E-13	8.02E-14	2.11E-08
	W-185	014932-41-3	7.51E+01	D	D	3 00E-01	2.04E-12	4.26E-13	5 03E-11
	W-187	014983-48-3	2.38E+01	H	D	3.00E-01	2.46E-12	5.29E-13	1 63E-06
Uranium (92)	U-232	014158-29-3	7.20E+01	Y	Y	5 00E-02	8.12E-11	5.29E-08	3.42E-11
	U-233	013968-55-3	1.59E+05	Y	Y	5.00E-02	4.48E-11	1.41E-08	3.52E-11
	U-234	013966-29-5	2.45E+05	Y	Y	5.00E-02	4.44E-11	1.40E-08	2.14E-11
	U-235	015117-96-1	7.04E+08	Y	Y	5.00E-02	4 52E-11	1 30E-08	2.63E-07
	U-235+D	015117-96-1(+D)	7 04E+08	Y	Y	5.00E-02	4.70E-11	1.30E-08	2 65E-07
	U-236	013982-70-2	2 34E+07	Y	Y	5.00E-02	4.21E-11	1.32E-08	1.72E-11
	U-237	014269-75-1	6.75E+00	D	Y	5.00E-02	3 98E-12	3.12E-12	1.37E-07
	U-238	007440-61-1	4.47E+09	Y	Y	5.00E-02	4.27E-11	1.24E-08	1.50E-11
	U-238+D	007440-61-1(+D)	4.47E+09	Y	Y	5.00E-02	6.20E-11	1.24E-08	5.25E-08
	U-240	015687-53-3	1.41E+01	H	Y	5 00E-02	5.47E-12	3 35E-12	1.09E-10
Vanadium (23)	V-48	014331-97-6	1.60E+01	D	W	1.00E-02	7.56E-12	6.84E-12	1 12E-05

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
Xenon (54)	Xe-122	015151-09-4	2.01E+01	H	*	---	---	3.08E-15	---
	Xe-123	015700-10-4	2.14E+00	H	*	---	---	8 92E-16	---
	Xe-125	013994-18-8	1.68E+01	H	*	---	---	1 20E-15	---
	Xe-127	013994-19-9	3.64E+01	D	*	---	---	4.09E-16	---
	Xe-129m	013965-99-6(m)	8.89E+00	D	*	---	---	5 74E-16	---
	Xe-131m	014683-11-5(m)	1.18E+01	D	*	---	---	4.13E-16	---
	Xe-133	014932-42-4	5.25E+00	D	*	---	---	4.14E-16	---
	Xe-133m	014932-42-4(m)	2.19E+00	D	*	---	---	5.12E-16	---
	Xe-135	014995-62-1	9.11E+00	H	*	---	---	7.45E-16	---
	Xe-135m	014995-62-1(m)	1.54E+01	M	*	---	---	1.88E-16	---
	Xe-137	014835-21-3	3.83E+00	M	*	---	---	1.39E-15	---
	Xe-138	015751-81-2	1.41E+01	M	*	---	---	2.06E-15	---
Yttrium (39)	Y-90	010098-91-6	6.41E+01	H	Y	1.00E-04	1.50E-11	9.90E-12	0 00E+0
	Y-91	014234-24-3	5 85E+01	D	Y	1.00E-04	1.35E-11	1.85E-11	1 41E-08
	Y-91m	014234-24-3(m)	4.97E+01	M	Y	1 00E-04	3.69E-14	2.99E-14	1 90E-06
	Y-92	015751-59-4	3.54E+00	H	Y	1.00E-04	1.95E-12	1.61E-12	9.80E-07
	Y-93	014981-70-5	1.01E+01	H	Y	1.00E-04	5.74E-12	3.48E-12	3.41E-07
Zinc (30)	Zn-65	013982-39-3	2.44E+02	D	Y	5.00E-01	9.93E-12	9.98E-12	2.27E-06

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

Element (Atomic Number)	Isotope[c]	CASRN[d]	Radioactive Half-life[e]		ICRP Lung Class[f]	GI Absorption Factor (f_1)[g]	Slope Factor Lifetime Excess Total Cancer Risk Per Unit Intake or Exposure: Ingestion (Risk/pCi)	Inhalation (Risk/pCi)	External Exposure (Risk/yr per pCi/g soil)
	Zn-69	013982-23-5	5 56E+01	M	Y	5.00E-01	6 19E-14	1.04E-13	2.03E-11
	Zn-69m	013982-23-5(m)	1.38E+01	H	Y	5.00E-01	1.52E-12	1.17E-12	1.43E-06
Zirconium (40)	Zr-93	015751-77-6	1.53E+06	Y	W	2.00E-03	5 21E-13	5.26E-12	0.00E+0
	Zr-95	013967-71-0	6.40E+01	D	W	2.00E-03	3.92E-12	6.48E-12	2.81E-06
	Zr-97	014928-30-4	1.69E+01	H	W	2.00E-03	1.04E-11	4.73E-12	6.85E-07

[Table 4 continues on the following page: Refer to Endnotes on the last page.]

Table 4: Radionuclide Carcinogenicity -- Slope Factors[a] (In Units of Picocuries[b])

JULY 1997

ENDNOTES:

[a] 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 central estimates in a linear model of the age-averaged, lifetime attributable radiation cancer incidence (fatal and nonfatal cancer) risk per unit of activity inhaled or ingested, expressed as risk/picocurie (pCi). External exposure slope factors are central estimates of the lifetime attributable radiation 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. If required, slope factors in Table 4 can be converted into the International System (SI) units of becquerels (1 Bq = 1 nuclear transformation per second) by dividing each inhalation, ingestion, or external exposure value by 27.03. 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. 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 3.7×10^{10} nuclear transformations per second. 1 picocurie (pCi) = 10^{12} Ci.

[c] 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 associated short-lived radioactive decay products (i.e., those decay products with radioactive half-lives less than or equal to 6 months) are also included. 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 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 NUMBER 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 and the IRIS Pilot:

Risk assessment values for chemicals currently being considered by EPA, but not yet on IRIS, are included in HEAST. In the past the EPA Reference Dose/Reference Concentration (RfD/RfC) and the Carcinogen Risk Assessment Verification Endeavor (CRAVE) Work Groups used to validate Agency systemic toxicity and carcinogen risk assessments, respectively. These Work Groups are now replaced by the IRIS Pilot which will be responsible for resolving any conflicts regarding toxicity values developed by various Program Offices. The IRIS Pilot peer review panels represent different EPA offices and external scientists experienced in issues related to both the qualitative and quantitative risk assessment of carcinogenic and toxic agents. Values verified by this system must undergo extensive peer review and represent an Agency consensus. Verified risk assessment values or changes are entered into the IRIS data base monthly.

* Questions concerning the IRIS Pilot: Call Amy Mills, NCEA Washington at (202) 260-0569.

3. Office of Research and Development (ORD/National Center for Environmental Assessment (NCEA) OSWER-OAQPS (Office of Solid Waste and Emergency Response-Office of Air Quality Planning and Standards) Documents:

A listing of most ORD/NCEA 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 National Center for Environmental Assessment (NCEA). 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
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Health Effects Assessments (HEAs): This document series was prepared by the National Center for Environmental Assessment (NCEA-CIN) 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 National Center for Environmental Assessment (NCEA-CIN) 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 National Center for Environmental Assessment (NCEA-CIN) 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 National Center for Environmental Assessment (NCEA-RTP) 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 National Center for Environmental Assessment (NCEA-RTP and NCEA-CIN) 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:

Drinking Water Criteria Documents (DWCDs): The National Center for Environmental Assessment (NCEA-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 National Center for 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 4 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

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/m^3), 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_3 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 <u>given appropriate circumstances and Superfund program objectives.</u>

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

III. 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. Chemicals may be searched either alphabetically by compound name or numerically by the CASRN.

The list has been updated to only include chemicals that are currently documented in this issue of the HEAST.

CHEMICAL NAME AND CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER CROSS REFERENCE
(LISTED BY NAME)

July 1997

ACENAPHTHENE	000083-32-9
ACENAPTHYLENE	000208-96-8
ACEPHATE	030560-19-1
ACETONE	000067-64-1
ACETONE CYANOHYDRIN	000075-86-5
ACETONITRILE	000075-05-8
ACETOPHENONE	000098-86-2
ACROLEIN	000107-02-8
ACRYLAMIDE	000079-06-1
ACRYLIC ACID	000079-10-7
ACRYLONITRILE	000107-13-1
ADIPONITRILE	000111-69-3
ALACHLOR	015972-60-8
ALDICARB	000116-06-3
ALDRIN	000309-00-2
ALLIDOCHLOR	000093-71-0
ALLYL ALCOHOL	000107-18-6
ALLYL CHLORIDE	000107-05-1
ALUMINUM	007429-90-5
ALUMINUM PHOSPHIDE	020859-73-8
AMETRYN	000834-12-8
AMINO-2-NAPHTHOL, 1-	002834-92-6
AMINO-2-NAPHTOL HYDROCHLORIDE, 1-	001198-27-2
AMINOPHENOL, M-	000591-27-5
AMINOPHENOL, O-	000095-55-6
AMINOPHENOL, P-	000123-30-8
AMINOPYRIDINE, 4-	000504-24-5
AMMONIA	007664-41-7
ANILINE	000062-53-3
ANTHRACENE	000120-12-7
ANTIMONY, METALLIC	007440-36-0
ANTIMONY PENTOXIDE	001314-60-9
ANTIMONY POTASSIUM TARTRATE	000304-61-0
ANTIMONY TETROXIDE	001332-81-6
ANTIMONY TRIOXIDE	001309-64-4
ARAMITE	000140-57-8
AROCLOR 1248	012672-29-6
AROCLOR 1254	011097-69-1
ARSENIC, INORGANIC	007440-38-2
ASBESTOS	001332-21-4
ATRAZINE	001912-24-9
AZOBENZENE	000103-33-3
BARIUM	007440-39-3
BARIUM CYANIDE	000542-62-1
BENEFIN	001861-40-1
BENZAL CHLORIDE	000098-87-3
BENZALDEHYDE	000100-52-7
BENZALDEHYDE CYANOHYDRIN	000532-28-5
BENZENE	000071-43-2
BENZENETHIOL / (THIOPHENOL)	000108-98-5
BENZIDINE	000092-87-5
BENZOIC ACID	000065-85-0
BENZOTRICHLORIDE	000098-07-7
BENZO[A]ANTHRACENE	000056-55-3
BENZO[A]PYRENE	000050-32-8
BENZO[K]FLUORANTHENE	000207-08-9
BENZYL ALCOHOL	000100-51-6
BENZYL CHLORIDE	000100-44-7
BERYLLIUM	007440-41-7
BIPHENYL, 1,1'	000092-52-4
BIS(2-CHLOROETHYL) ETHER	000111-44-4
BIS(2-CHLOROISOPROPYL) ETHER	039638-32-9
BIS(2-CHLORO-1-METHYLETHYL) ETHER	000108-60-1
BIS(2-ETHYLHEXYL) PHTHALATE / (DEHP)	000117-81-7
BIS(CHLOROMETHYL) ETHER	000542-88-1
BISPHENOL A	000080-05-7
BORON, ELEMENTAL	007440-42-8
BORON TRIFLUORIDE	007637-07-2
BROMINATED DIBENZO-P-DIOXINS	NO CASRN
BROMINATED DIBENZOFURANS	NO CASRN
BROMOACETONE	000598-31-2
BROMOCHLOROETHANES	NO CASRN
BROMODICHLOROMETHANE	000075-27-4
BROMOETHENE / (VINYL BROMIDE)	000593-60-2
BROMOFORM	000075-25-2
BROMOMETHANE	000074-83-9
BROMOPHENYL PHENYL ETHER, 4-	000101-55-3
BROMOPHOS	002104-96-3
BROMOXYNIL	001689-84-5
BROMOXYNIL OCTANOATE	001689-99-2
BUSAN 77	031512-74-0
BUSAN 90	002491-38-5
BUTADIENE, 1,3-	000106-99-0
BUTANOL, 1-	000071-36-3
BUTYL BENZYL PHTHALATE, N-	000085-68-7
BUTYLATE	002008-41-5
BUTYLCHLORIDE, T-	000507-20-0
BUTYROLACTONE, GAMMA-	000096-48-0
CACODYLIC ACID	000075-60-5
CADMIUM	007440-43-9
CALCIUM CYANIDE	000592-01-8
CAPROLACTAM	000105-60-2
CAPTAFOL	002425-06-1
CAPTAN	000133-06-2
CARBARYL	000063-25-2
CARBAZOLE	000086-74-8
CARBOFURAN	001563-66-2
CARBON DISULFIDE	000075-15-0
CARBON MONOXIDE	000630-05-0
CARBON TETRACHLORIDE	000056-23-5
CHLORAL	000075-87-6
CHLORANIL	000118-75-2
CHLORDANE	000057-74-9
CHLORINE CYANIDE	000506-77-4
CHLORO-1,3-BUTADIENE, 2- / (CHLOROPRENE)	000126-99-8
CHLORO-2-METHYLANILINE, 4-	000095-69-2
CHLORO-2-METHYLANILINE HYDROCHLORIDE, 4-	003165-93-3
CHLORO-M-CRESOL, P-	000059-50-7
CHLOROACETALDEHYDE	000107-20-0
CHLOROACETIC ACID	000079-11-8
CHLOROANILINE, 2-	000095-51-2
CHLOROANILINE, 3-	000108-42-9
CHLOROANILINE, 4-	000106-47-8
CHLOROBENZENE	000108-90-7
CHLOROBENZILATE	000510-15-6
CHLOROBENZOIC ACID, P-	000074-11-3
CHLOROBENZOTRIFLUORIDE, 4-	000098-56-6
CHLOROBUTANE, 1-	000109-69-3
CHLOROBUTANE, 2-	000078-86-4
CHLOROCYCLOPENTADIENE	041851-50-7
CHLOROFORM	000067-66-3
CHLOROMETHANE / (METHYL CHLORIDE)	000074-87-3
CHLOROMETHYL METHYL ETHER	000107-30-2
CHLORONITROBENZENE, M-	000121-73-3
CHLORONITROBENZENE, O-	000088-73-3
CHLORONITROBENZENE, P-	000100-00-5
CHLOROPHENOL, 2-	000095-57-8
CHLOROPHENOL, 3-	000108-43-0
CHLOROPHENOL, 4-	000106-48-9
CHLOROPRENE	000126-99-8
CHLOROPROPANE, 2-	000075-29-6
CHLOROTHALONIL	001897-45-6
CHLOROTOLUENE, M-	000108-41-8
CHLOROTOLUENE, O-	000095-49-8
CHLOROTOLUENE, P-	000106-43-4
CHLORPYRIFOS	002921-88-2
CHLORPYRIFOS METHYL	005598-13-0
CHLORTHIOPHOS	060238-56-4
CHROMIUM(III)	016065-83-1
CHROMIUM(VI)	018540-29-9
CHRYSENE	000218-01-9
COKE OVEN EMISSIONS	008007-45-2
COPPER	007440-50-8
COPPER CYANIDE	000544-92-3
CREOSOTE, COAL TAR	008001-58-9
CRESOL, M- / (3-METHYLPHENOL)	000108-39-4
CRESOL, O- / (2-METHYLPHENOL)	000095-48-7
CRESOL, P- / (4-METHYLPHENOL)	000106-44-5

CROTONALDEHYDE	000123-73-9
CUMENE	000098-82-8
CYANAZINE	021725-46-2
CYANIDE	000057-12-5
CYANOGEN	000460-19-5
CYANOGEN BROMIDE	000506-68-3
CYCLOATE	001134-23-2
CYCLOHEXANOL	000108-93-0
CYCLOHEXYLAMINE	000108-91-8
CYCLONITE	000121-82-4
CYCLOPENTADIENE	000542-92-7
DACTHAL	001861-32-1
DALAPON	000075-99-0
2,4-DB	000094-82-6
DDD	000072-54-8
DDE	000072-55-9
DDT	000050-29-3
DECABROMODIPHENYL ETHER	001163-19-5
DEHP	000117-81-7
DI-N-OCTYL PHTHALATE	000117-84-0
DIALLATE	002303-16-4
DIAZINON	000333-41-5
DIBENZOFURAN	000132-64-9
DIBENZO[A,H]ANTHRACENE	000053-70-3
DIBROMO-3-CHLOROPROPANE, 1,2	000096-12-8
DIBROMOBENZENE, 1,4-	000106-37-6
DIBROMOCHLOROMETHANE	000124-48-1
DIBROMOETHANE, 1,2-	000106-93-4
DIBUTYL PHTHALATE	000084-74-2
DICAMBA	001918-00-9
DICHLORO-2-BUTENE, 1,4-	000764-41-0
DICHLOROBENZENE, 1,2-	000095-50-1
DICHLOROBENZENE, 1,3-	000541-73-1
DICHLOROBENZENE, 1,4-	000106-46-7
DICHLOROBENZIDINE, 3,3'-	000091-94-1
DICHLOROBUTENES	NO CASRN
DICHLORODIFLUOROMETHANE	000075-71-8
DICHLOROETHANE, 1,1-	000075-34-3
DICHLOROETHYLENE, 1,1-	000075-35-4
DICHLOROETHYLENE, 1,2- (MIXED ISOMERS)	000540-59-0
DICHLOROETHYLENE, 1,2-C-	000156-59-2
DICHLOROETHYLENE, 1,2-T-	000156-60-5
DICHLOROMETHANE	000075-09-2
DICHLOROPHENOL, 2,3-	000576-24-9
DICHLOROPHENOL, 2,4-	000120-83-2
DICHLOROPHENOL, 2,5-	000583-78-8
DICHLOROPHENOL, 2,6-	000087-65-0
DICHLOROPHENOL, 3,4-	000095-77-2
DICHLOROPHENOL, 3,5-	000591-35-5
DICHLOROPHENOXY ACETIC ACID, 2,4-	000094-75-7
DICHLOROPHENOXY) BUTYRIC ACID, 4-(2,4- / (2,4-DB)	000094-82-6
DICHLOROPROPANE, 1,1-	000078-99-9
DICHLOROPROPANE, 1,2-	000078-87-5
DICHLOROPROPANE, 1,3-	000142-28-9
DICHLOROPROPANE, 2,2-	000594-20-7
DICHLOROPROPENE, 1,3- / (TELONE II)	000542-75-6
DICHLORPROP	000120-36-5
DICYCLOPENTADIENE	000077-73-6
DIELDRIN	000060-57-1
DIETHYL-P-NITROPHENYL PHOSPHATE	000311-45-5
DIETHYL PHTHALATE	000084-66-2
DIETHYLANILINE, N,N-	000091-66-7
DIETHYLENE GLYCOL MONOBUTYL ETHER	000112-34-5
DIETHYLENE GLYCOL MONOETHYL ETHER	000111-90-0
DIETHYLFORMAMIDE	000617-84-5
DIETHYLHYDRAZINE, 1,2-	001615-80-1
DIETHYLSTILBESTROL	000056-53-1
DIMETHOATE	000060-51-5
DIMETHOXYBENZIDINE, 3,3'-	000119-90-4
DIMETHYLANILINE, 2,4-	000095-68-1
DIMETHYLANILINE HYDROCHLORIDE, 2,4-	021436-96-4
DIMETHYLANILINE, N,N-	000121-69-7
DIMETHYLBENZIDINE, 3,3'-	000119-93-7
DIMETHYLFORMAMIDE, N,N-	000068-12-2
DIMETHYLHYDRAZINE, 1,1-	000057-14-7
DIMETHYLHYDRAZINE, 1,2-	000540-73-8
DIMETHYLPHENOL, 2,3-	000526-75-0
DIMETHYLPHENOL, 2,4-	000105-67-9
DIMETHYLPHENOL, 2,5-	000095-87-4
DIMETHYLPHENOL, 2,6-	000576-26-1
DIMETHYLPHENOL, 3,4-	000095-65-8
DIMETHYLPHTHALATE	000131-11-3
DIMETHYLSULFATE	000077-78-1
DIMETHYLTEREPHTHALATE	000120-61-6
DIMETHYLUREA, N,N-	000598-94-7
DINITRO-O-CRESOL, 4,6-	000534-52-1
DINITRO-P-CRESOL, 2,6-	000609-93-8
DINITROBENZENE, 1,2-	000528-29-0
DINITROBENZENE, 1,3-	000099-65-0
DINITROBENZENE, 1,4-	000100-25-4
DINITROPHENOL, 2,3-	000066-56-8
DINITROPHENOL, 2,4-	000051-28-5
DINITROPHENOL, 2,5-	000329-71-5
DINITROPHENOL, 2,6-	000573-56-8
DINITROPHENOL, 3,5-	000586-11-8
DINITROTOLUENE, 2,3-	000602-01-7
DINITROTOLUENE, 2,4	000121-14-2
DINITROTOLUENE, 2,5-	000619-15-8
DINITROTOLUENE, 2,6-	000606-20-2
DINITROTOLUENE, 3,4-	000610-39-9
DINOSEB	000088-85-7
DIOXANE, 1,4-	000123-91-1
DIPHENYLAMINE, N,N-	000122-39-4
DIPHENYLHYDRAZINE, 1,2-	000122-66-7
DIPHENYLMETHANE DIISOCYANATE	000101-68-8
DIRECT BLACK 38	001937-37-7
DIRECT BLUE 6	002602-46-2
DIRECT BROWN 95	016071-86-6
DIRECT LIGHTFAST BLUE	004399-55-7
DIRECT SKY BLUE 6B	002610-05-1
DISULFOTON	000298-04-4
ENDOSULFAN	000115-29-7
ENDOTHALL	000145-73-3
ENDRIN	000072-20-8
EPICHLOROHYDRIN	000106-89-8
EPTC	000759-94-4
ETHOPROP	013194-48-4
ETHOXYETHANOL, 2-	000110-80-5
ETHOXYETHANOL ACETATE, 2-	000111-15-9
ETHOXYETHANOL ACRYLATE, 2-	000106-74-1
ETHOXYETHANOL DODECANOATE, 2-	000106-13-8
ETHOXYETHANOL PHOSPHATE, 2-	068554-00-7
ETHOXYETHYL METHACRYLATE, 2-	002370-63-0
ETHYL ACETATE	000141-78-6
ETHYL ACRYLATE	000140-88-5
ETHYL BENZENE	000100-41-4
ETHYL CHLORIDE	000075-00-3
ETHYL ETHER	000060-29-7
ETHYL METHACRYLATE	000097-63-2
ETHYL-O-XYLENE, 4-	000934-80-5
ETHYLANILINE, N-	000103-69-5
ETHYLENE CYANOHYDRIN	000109-78-4
ETHYLENE DIAMINE	000107-15-3
ETHYLENE GLYCOL	000107-21-1
ETHYLENE GLYCOL MONOBUTYL ETHER	000111-76-2
ETHYLENE OXIDE	000075-21-8
ETHYLENE THIOUREA	000096-45-7
ETHYLTOLUENE, M-	000620-14-4
ETHYLTOLUENE, O-	000611-14-3
ETHYLTOLUENE, P-	000622-96-8
FLUORANTHENE	000206-44-0
FLUORENE	000086-73-7
FLUORINE / (SOLUBLE FLUORIDE)	007782-41-4
FLURIDONE	059756-60-4
FOLPET	000133-07-3

FORMALDEHYDE	000050-00-0
FORMALDEHYDE CYANOHYDRIN	000107-16-4
FORMIC ACID	000064-18-6
FURAN	000110-00-9
FURAZOLIDONE	000067-45-8
FURFURAL	000098-01-1
FURIUM	000531-82-8
GLYCIDALDEHYDE	000765-34-4
HEPTACHLOR	000076-44-8
HEPTACHLOR EPOXIDE	001024-57-3
HEPTANE, N-	000142-82-5
HEXABROMOBENZENE	000087-82-1
HEXACHLOROBENZENE	000118-74-1
HEXACHLOROBUTADIENE	000087-68-3
HEXACHLOROCYCLOHEXANE, ALPHA-	000319-84-6
HEXACHLOROCYCLOHEXANE, BETA-	000319-85-7
HEXACHLOROCYCLOHEXANE, DELTA-	000319-86-8
HEXACHLOROCYCLOHEXANE, EPSILON-	006108-10-7
HEXACHLOROCYCLOHEXANE, GAMMA-	000058-89-9
HEXACHLOROCYCLOHEXANE-TECHNICAL	000608-73-1
HEXACHLOROCYCLOPENTADIENE	000077-47-4
HEXACHLOROETHANE	000067-72-1
HEXACHLOROPHENE	000070-30-4
HEXAMETHYLENE DIAMINE	000124-09-4
HEXANE, N-	000110-54-3
HEXANONE, 2-	000591-78-6
HYDRAZINE	000302-01-2
HYDRAZINE SULFATE	010034-93-2
HYDROGEN SULFIDE	007783-06-4
HYDROQUINONE	000123-31-9
INDENO[1,2,3-CD]PYRENE	000193-39-5
IRON	007439-89-6
ISOBUTYL ALCOHOL	000078-83-1
ISOPHORONE	000078-59-1
ISOPROPALIN	033820-53-0
LACTONITRILE	000078-97-7
LEAD	007439-92-1
LEAD ALKYLS	NO CASRN
LINURON	000330-55-2
MALATHION	000121-75-5
MALEIC ANHYDRIDE	000108-31-6
MALEIC HYDRAZIDE	000123-33-1
MALONONITRILE	000109-77-3
MANCOZEB	008018-01-7
MANEB	012427-38-2
MANGANESE	007439-96-5
MEPHOSFOLAN	000950-10-7
MERCURIC CHLORIDE	007487-94-7
MERCURY, ELEMENTAL	007439-97-6
MERPHOS	000150-50-5
MERPHOS OXIDE	000078-48-8
METHACRYLONITRILE	000126-98-7
METHANOL	000067-56-1
METHOMYL	016752-77-5
METHOXY-5-NITROANILINE, 2-	000099-59-2
METHOXYCHLOR	000072-43-5
METHOXYETHANOL, 2-	000109-86-4
METHOXYETHANOL ACETATE, 2-	000110-49-6
METHYL-4-CHLOROPHENOXY) BUTYRIC ACID, 4-(2-	000094-81-5
METHYL-4-CHLOROPHENOXY) PROPIONIC ACID, 2-(2-	000093-65-2
METHYL-4-CHLOROPHENOXY ACETIC ACID, 2-	000094-74-6
METHYL-5-NITROANILINE, 2-	000099-55-8
METHYL ACETATE	000079-20-9
METHYL ACRYLATE	000096-33-3
METHYL CHLORIDE	000074-87-3
METHYL CHLOROCARBONATE	000079-22-1
METHYL ETHYL KETONE	000078-93-3
METHYL ETHYL KETONE PEROXIDE	001338-23-4
METHYL HYDRAZINE	000060-34-4
METHYL ISOBUTYL KETONE	000108-10-1
METHYL ISOCYANATE	000624-83-9
METHYL MERCURY	022967-92-6
METHYL METHACRYLATE	000080-62-6
METHYL PARATHION	000298-00-0
METHYL STYRENE (MIXED ISOMERS)	025013-15-4
METHYL STYRENE, ALPHA	000098-83-9
METHYLANILINE, 2-	000095-53-4
METHYLANILINE HYDROCHLORIDE, 2-	000636-21-5
METHYLCYCLOHEXANE	000108-87-2
METHYLENE-BIS(2-CHLOROANILINE), 4,4'-	000101-14-4
METHYLENE-BIS(N,N'-DIMETHYL)ANILINE, 4,4'-	000101-61-1
METHYLENE BROMIDE	000074-95-3
METHYLENE CHLORIDE / (DICHLOROMETHANE)	000075-09-2
METHYLENE-BIS(BENZENEAMINE), 4,4- / (METHYLENE DIANILINE, 4,4-)	000101-77-9
METHYLENEDIPHENYL ISOCYANATE, 4,4- / (DIPHENYLMETHANE DIISOCYANATE)	000101-68-8
2-METHYLLACTONITRILE	000075-86-5
2-METHYLPHENOL	000095-48-7
3-METHYLPHENOL	000108-39-4
4-METHYLPHENOL	000106-44-5
METOLACHLOR	051218-45-2
METRIBUZIN	021087-64-9
MIREX	002385-85-5
MOLINATE	002212-67-1
MOLYBDENUM	007439-98-7
MONOCHLORAMINE	010599-90-3
NAPHTHALENE	000091-20-3
NAPHTHOQUINONE, 1,4-	000130-15-4
NIAGARA BLUE 4B	002429-74-5
NICKEL CYANIDE	000557-19-7
NICKEL, REFINERY DUST	NO CASRN
NICKEL, SOLUBLE SALTS	Various
NICKEL SUBSULFIDE	012035-72-2
NICOTINONITRILE	000100-54-9
NITRIC OXIDE	010102-43-9
NITRITE	014797-65-0
NITROANILINE, 2-	000088-74-4
NITROANILINE, M-	000099-09-2
NITROANILINE, P-	000100-01-6
NITROBENZENE	000098-95-3
NITROFURANTOIN	000067-20-9
NITROFURAZONE	000059-87-0
NITROGEN DIOXIDE	010102-44-0
NITROGEN OXIDES	NO CASRN
NITROMETHANE	000075-52-5
NITROPHENOLS	NO CASRN
NITROPROPANE, 2-	000079-46-9
NITROSO-DI-N-BUTYLAMINE, N-	000924-16-3
NITROSO-DI-N-PROPYLAMINE, N-	000621-64-7
NITROSO-N-ETHYLUREA, N-	000759-73-9
NITROSO-N-METHYLUREA, N-	000684-93-5
NITROSODIETHANOLAMINE, N-	001116-54-7
NITROSODIETHYLAMINE, N-	000055-18-5
NITROSODIMETHYLAMINE, N-	000062-75-9
NITROSODIPHENYLAMINE, P-	000156-10-5
NITROSODIPHENYLAMINE, N-	000086-30-6
NITROSOMETHYLETHYLAMINE, N-	010595-95-6
NITROSOMETHYLVINYLAMINE, N	004549-40-0
NITROSOPYRROLIDINE, N-	000930-55-2
NITROTOLUENE, M-	000099-08-1
NITROTOLUENE, O-	000088-72-2
NITROTOLUENE, P-	000099-99-0
OCTABROMODIPHENYL ETHER	032536-52-0
OCTAMETHYLPYROPHOSPHORAMIDE	000152-16-9
OSMIUM TETROXIDE	020816-12-0
OZONE	010028-15-6
PARALDEHYDE	000123-63-7
PARATHION	000056-38-2
PARTICULATE MATTER	NO CASRN
PEBULATE	001114-71-2

PENDIMETHALIN	040487-42-1
PENTABROMO-6-CHLOROCYCLOHEXANE, 1,2,3,4,5-	000087-84-3
PENTABROMODIPHENYL ETHER	032534-81-9
PENTACHLOROBENZENE	000608-93-5
PENTACHLOROCYCLOPENTADIENE	025329-35-5
PENTACHLORONITROBENZENE	000082-68-8
PENTACHLOROPHENOL	000087-86-5
PENTACHLOROPROPENE, 1,1,2,3,3,-	001600-37-9
PENTANE, N-	000109-66-0
PHENANTHRENE	000085-01-8
PHENOL	000108-95-2
PHENYLENEDIAMINE, M-	000108-45-2
PHENYLENEDIAMINE, O-	000095-54-5
PHENYLENEDIAMINE, P-	000106-50-3
PHENYLMERCURIC ACETATE	000062-38-4
PHENYLPHENOL, 2-	000090-43-7
PHORATE	000298-02-2
PHOSGENE	000075-44-5
PHOSPHINE	007803-51-2
PHOSPHORUS, WHITE	007723-14-0
PHOTOCHEMICAL OXIDANTS	NO CASRN
PHTHALIC ACID, M-	000121-91-5
PHTHALIC ACID, O-	000088-99-3
PHTHALIC ACID, P-	000100-21-0
PHTHALIC ANHYDRIDE	000085-44-9
POLYBROMINATED BIPHENYLS	NO CASRN
POLYCHLORINATED BIPHENYLS	001336-36-3
POTASSIUM CYANIDE	000151-50-8
POTASSIUM SILVER CYANIDE	000506-61-6
PROFLURALIN	026399-36-0
PRONAMIDE	023950-58-5
PROPACHLOR	001918-16-7
PROPAZINE	000139-40-2
PROPIONITRILE	000107-12-0
PROPYL ALCOHOL, N-	000071-23-8
PROPYLENE GLYCOL	000057-55-6
PROPYLENE GLYCOL MONOETHYL ETHER	001569-02-4
PROPYLENE GLYCOL MONOMETHYL ETHER	000107-98-2
PROPYLENE OXIDE	000075-56-9
PYRENE	000129-00-0
PYRIDINE	000110-86-1
QUINOLINE	000091-22-5
RDX / (CYCLONITE)	000121-82-4
RONNEL	000299-84-3
SELENIOUS ACID	007783-00-8
SELENIUM	007782-49-2
SELENIUM SULFIDE	007446-34-6
SELENOUREA	000630-10-4
SILVER	007440-22-4
SILVER CYANIDE	000506-64-9
SIMAZINE	000122-34-9
SODIUM CYANIDE	000143-33-9
SODIUM DIETHYLDITHIOCARBAMATE	000148-18-5
SODIUM METAVANADATE	013718-26-8
STIROPHOS	000961-11-5
STRONTIUM, STABLE	007440-24-6
STRYCHNINE	000057-24-9
STYRENE	000100-42-5
SUCCINONITRILE	000110-61-2
SULFUR DIOXIDE	007446-09-5
SULFUR OXIDES	NO CASRN
SULFURIC ACID	007664-93-9
TCDD, 2,3,7,8-	001746-01-6
TELONE II	000542-75-6
TEMEPHOS	003383-96-8
TERBUFOS	013071-79-9
TEREPHTHALIC ACID	000100-21-0
TETRACHLOROAZOXYBENZENE	021232-47-3
TETRACHLOROBENZENE, 1,2,4,5-	000095-94-3
TETRACHLOROCYCLOPENTADIENE	000695-77-2
TETRACHLOROETHANE, 1,1,1,2-	000630-20-6
TETRACHLOROETHANE, 1,1,2,2-	000079-34-5
TETRACHLOROETHYLENE	000127-18-4
TETRACHLOROHYDRAZOBENZENE	071753-42-9
TETRACHLOROPHENOL, 2,3,4,5-	004901-51-3
TETRACHLOROPHENOL, 2,3,4,6-	000058-90-2
TETRACHLOROPHENOL, 2,3,5,6-	000935-95-5
TETRACHLOROPROPENE, 1,1,2,3-	010436-39-2
TETRACHLOROTOLUENE, PARA, ALPHA, ALPHA, ALPHA-	005216-25-1
TETRACHLOROVINPHOS / (STIROPHOS)	000961-11-5
TETRAETHYL DITHIOPYROPHOSPHATE	003689-24-5
THALLIC OXIDE	001314-32-5
THALLIUM (I) ACETATE	000563-68-8
THALLIUM (I) CARBONATE	006533-73-9
THALLIUM (I) CHLORIDE	007791-12-0
THALLIUM, INSOLUBLE SALTS	NO CASRN
THALLIUM (I) NITRATE	010102-45-1
THALLIUM SELENITE	012039-52-0
THALLIUM (I) SULFATE	007446-18-6
THIOCYANOMETHYLTHIO)BENZOTHIAZOLE, 2-(	021564-17-0
THIOFANOX	013196-18-4
THIOPHENOL	000108-98-5
THIRAM	000137-26-8
TIN AND COMPOUNDS	NO CASRN
TOLUENE	000108-88-3
TOLUENE-2,4-DIAMINE	000095-80-7
TOLUENE-2,5-DIAMINE	000095-70-5
TOLUENE-2,6-DIAMINE	000823-40-5
TOLUENEDIAMINE, 2,3-	002687-25-4
TOLUENEDIAMINE, 3,4-	000496-72-0
TOLUIDINE, M-	000108-44-1
TOLUIDINE, P-	000106-49-0
TOXAPHENE	008001-35-2
TRIALLATE	002303-17-5
TRIBROMOBENZENE, 1,2,4-	000615-54-3
TRICHLORO-1,2,2-TRIFLUOROETHANE, 1,1,2-	000076-13-1
TRICHLORO-2'-HYDROXYDIPHENYLETHER, 2,2,4'-	003380-34-5
TRICHLOROANILINE, 2,4,6-	000634-93-5
TRICHLOROANILINE HYDROCHLORIDE, 2,4,6-	033663-50-2
TRICHLOROBENZENE, 1,2,4-	000120-82-1
TRICHLOROCYCLOPENTADIENE	077323-84-3
TRICHLOROETHANE, 1,1,1-	000071-55-6
TRICHLOROETHANE, 1,1,2-	000079-00-5
TRICHLOROETHYLENE	000079-01-6
TRICHLOROFLUOROMETHANE	000075-69-4
TRICHLOROPHENOL, 2,3,4-	015950-66-0
TRICHLOROPHENOL, 2,3,5-	000933-78-8
TRICHLOROPHENOL, 2,3,6-	000933-75-5
TRICHLOROPHENOL, 2,4,5-	000095-95-4
TRICHLOROPHENOL, 2,4,6-	000088-06-2
TRICHLOROPHENOL, 3,4,5-	000609-19-8
TRICHLOROPHENOXY) PROPIONIC ACID, 2(2,4,5-	000093-72-1
TRICHLOROPHENOXY ACETIC ACID, 2,4,5-	000093-76-5
TRICHLOROPROPANE, 1,1,1-	007789-89-1
TRICHLOROPROPANE, 1,1,2-	000598-77-6
TRICHLOROPROPANE, 1,2,2-	003175-23-3
TRICHLOROPROPANE, 1,2,3-	000096-18-4
TRICHLOROPROPENE, 1,2,3-	000096-19-5
TRICHLOROTOLUENE, 2,3,6-	002077-46-5
TRICHLOROTOLUENE, ALPHA,2,6-	002014-83-7
TRIFLURALIN	001582-09-8
TRIMETHYL PHOSPHATE	000512-56-1
TRIMETHYLBENZENES	NO CASRN
TRINITROBENZENE, 1,3,5-	000099-35-4
TRINITROPHENOLS	NO CASRN
TRINITROPHENYLMETHYLNITRAMINE	000479-45-8

TRINITROTOLUENE, 2,4,6-	000118-96-7
VANADIUM	007440-62-2
VANADIUM PENTOXIDE	001314-62-1
VANADIUM SULFATE	036907-42-3
VERNAM / (VERNOLATE)	001929-77-7
VERNOLATE	001929-77-7
VINYL-1-CYCLOHEXENE, 4-	000100-40-3
VINYL ACETATE	000108-05-4
VINYL BROMIDE	000593-60-2
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

CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER AND CHEMICAL NAME CROSS REFERENCE
(LISTED BY CHEMICAL ABSTRACTS SERVICE REGISTRY NUMBER)

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000050-00-0 FORMALDEHYDE
000050-29-3 DDT
000050-32-8 BENZO[A]PYRENE
000051-28-5 DINITROPHENOL, 2,4-
000053-70-3 DIBENZO[A,H]ANTHRACENE
000055-18-5 NITROSODIETHYLAMINE, N-
000056-23-5 CARBON TETRACHLORIDE
000056-38-2 PARATHION
000056-53-1 DIETHYLSTILBESTROL
000056-55-3 BENZO[A]ANTHRACENE
000057-12-5 CYANIDE
000057-14-7 DIMETHYLHYDRAZINE, 1,1-
000057-24-9 STRYCHNINE
000057-55-6 PROPYLENE GLYCOL
000057-74-9 CHLORDANE
000058-89-9 HEXACHLOROCYCLOHEXANE, GAMMA-
000058-90-2 TETRACHLOROPHENOL, 2,3,4,6-
000059-50-7 CHLORO-M-CRESOL, P-
000059-87-0 NITROFURAZONE
000060-29-7 ETHYL ETHER
000060-34-4 METHYL HYDRAZINE
000060-51-5 DIMETHOATE
000060-57-1 DIELDRIN
000062-38-4 PHENYLMERCURIC ACETATE
000062-53-3 ANILINE
000062-75-9 NITROSODIMETHYLAMINE, N-
000063-25-2 CARBARYL
000064-18-6 FORMIC ACID
000065-85-0 BENZOIC ACID
000066-56-8 DINITROPHENOL, 2,3-
000067-20-9 NITROFURANTOIN
000067-45-8 FURAZOLIDONE
000067-56-1 METHANOL
000067-64-1 ACETONE
000067-66-3 CHLOROFORM
000067-72-1 HEXACHLOROETHANE
000068-12-2 DIMETHYLFORMAMIDE, N,N-
000070-30-4 HEXACHLOROPHENE
000071-23-8 PROPYL ALCOHOL, N-
000071-36-3 BUTANOL, 1-
000071-43-2 BENZENE
000071-55-6 TRICHLOROETHANE, 1,1,1-
000072-20-8 ENDRIN
000072-43-5 METHOXYCHLOR
000072-54-8 DDD
000072-55-9 DDE
000074-11-3 CHLOROBENZOIC ACID, P-
000074-83-9 BROMOMETHANE
000074-87-3 METHYL CHLORIDE
000074-87-3 CHLOROMETHANE
000074-95-3 METHYLENE BROMIDE
000075-00-3 ETHYL CHLORIDE
000075-01-4 VINYL CHLORIDE
000075-05-8 ACETONITRILE
000075-09-2 METHYLENE CHLORIDE
000075-09-2 DICHLOROMETHANE
000075-15-0 CARBON DISULFIDE
000075-21-8 ETHYLENE OXIDE
000075-25-2 BROMOFORM
000075-27-4 BROMODICHLOROMETHANE
000075-29-6 CHLOROPROPANE, 2-
000075-34-3 DICHLOROETHANE, 1,1-
000075-35-4 DICHLOROETHYLENE, 1,1-
000075-44-5 PHOSGENE
000075-52-5 NITROMETHANE
000075-56-9 PROPYLENE OXIDE
000075-60-5 CACODYLIC ACID
000075-69-4 TRICHLOROFLUOROMETHANE
000075-71-8 DICHLORODIFLUOROMETHANE
000075-86-5 2-METHYLLACTONITRILE
000075-86-5 ACETONE CYANOHYDRIN
000075-87-6 CHLORAL
000075-99-0 DALAPON
000076-13-1 TRICHLORO-1,2,2-TRIFLUOROETHANE, 1,1,2-
000076-44-8 HEPTACHLOR
000077-47-4 HEXACHLOROCYCLOPENTADIENE
000077-73-6 DICYCLOPENTADIENE
000077-78-1 DIMETHYLSULFATE
000078-48-8 MERPHOS OXIDE
000078-59-1 ISOPHORONE
000078-83-1 ISOBUTYL ALCOHOL
000078-86-4 CHLOROBUTANE, 2-
000078-87-5 DICHLOROPROPANE, 1,2-
000078-93-3 METHYL ETHYL KETONE
000078-97-7 LACTONITRILE
000078-99-9 DICHLOROPROPANE, 1,1-
000079-00-5 TRICHLOROETHANE, 1,1,2-
000079-01-6 TRICHLOROETHYLENE
000079-06-1 ACRYLAMIDE
000079-10-7 ACRYLIC ACID
000079-11-8 CHLOROACETIC ACID
000079-20-9 METHYL ACETATE
000079-22-1 METHYL CHLOROCARBONATE
000079-34-5 TETRACHLOROETHANE, 1,1,2,2-
000079-46-9 NITROPROPANE, 2-
000080-05-7 BISPHENOL A
000080-62-6 METHYL METHACRYLATE
000081-81-2 WARFARIN
000082-68-8 PENTACHLORONITROBENZENE
000083-32-9 ACENAPHTHENE
000084-66-2 DIETHYL PHTHALATE
000084-74-2 DIBUTYL PHTHALATE
000085-01-8 PHENANTHRENE
000085-44-9 PHTHALIC ANHYDRIDE
000085-68-7 BUTYL BENZYL PHTHALATE, N-
000086-30-6 NITROSODIPHENYLAMINE, N-
000086-73-7 FLUORENE
000086-74-8 CARBAZOLE
000087-65-0 DICHLOROPHENOL, 2,6-
000087-68-3 HEXACHLOROBUTADIENE
000087-82-1 HEXABROMOBENZENE
000087-84-3 PENTABROMO-6-CHLOROCYCLOHEXANE, 1,2,3,4,5-
000087-86-5 PENTACHLOROPHENOL
000088-06-2 TRICHLOROPHENOL, 2,4,6-
000088-72-2 NITROTOLUENE, O-
000088-73-3 CHLORONITROBENZENE, O-
000088-74-4 NITROANILINE, 2-
000088-85-7 DINOSEB
000088-99-3 PHTHALIC ACID, O-
000090-43-7 PHENYLPHENOL, 2-
000091-20-3 NAPHTHALENE
000091-22-5 QUINOLINE
000091-66-7 DIETHYLANILINE, N,N-
000091-94-1 DICHLOROBENZIDINE, 3,3'-
000092-52-4 BIPHENYL, 1,1'
000092-87-5 BENZIDINE
000093-65-2 METHYL-4-CHLOROPHENOXY) PROPIONIC ACID, 2-(2-
000093-71-0 ALLIDOCHLOR
000093-72-1 TRICHLOROPHENOXY) PROPIONIC ACID, 2(2,4,5-
000093-76-5 TRICHLOROPHENOXY ACETIC ACID, 2,4,5-
000094-74-6 METHYL-4-CHLOROPHENOXY ACETIC ACID, 2-
000094-75-7 DICHLOROPHENOXY ACETIC ACID, 2,4-
000094-81-5 METHYL-4-CHLOROPHENOXY) BUTYRIC ACID, 4-(2-
000094-82-6 2,4-DB
000094-82-6 DICHLOROPHENOXY) BUTYRIC ACID, 4-(2,4-
000095-47-6 XYLENE, O-
000095-48-7 CRESOL, O-
000095-48-7 2-METHYLPHENOL
000095-49-8 CHLOROTOLUENE, O-
000095-50-1 DICHLOROBENZENE, 1,2-
000095-51-2 CHLOROANILINE, 2-
000095-53-4 METHYLANILINE, 2-

000095-54-5	PHENYLENEDIAMINE, O-
000095-55-6	AMINOPHENOL, O-
000095-57-8	CHLOROPHENOL, 2-
000095-65-8	DIMETHYLPHENOL, 3,4-
000095-68-1	DIMETHYLANILINE, 2,4-
000095-69-2	CHLORO-2-METHYLANILINE, 4-
000095-70-5	TOLUENE-2,5-DIAMINE
000095-77-2	DICHLOROPHENOL, 3,4-
000095-80-7	TOLUENE-2,4-DIAMINE
000095-87-4	DIMETHYLPHENOL, 2,5-
000095-94-3	TETRACHLOROBENZENE, 1,2,4,5-
000095-95-4	TRICHLOROPHENOL, 2,4,5-
000096-12-8	DIBROMO-3-CHLOROPROPANE, 1,2
000096-18-4	TRICHLOROPROPANE, 1,2,3-
000096-19-5	TRICHLOROPROPENE, 1,2,3-
000096-33-3	METHYL ACRYLATE
000096-45-7	ETHYLENE THIOUREA
000096-48-0	BUTYROLACTONE, GAMMA-
000097-63-2	ETHYL METHACRYLATE
000098-01-1	FURFURAL
000098-07-7	BENZOTRICHLORIDE
000098-56-6	CHLOROBENZOTRIFLUORIDE, 4-
000098-82-8	CUMENE
000098-83-9	METHYL STYRENE, ALPHA
000098-86-2	ACETOPHENONE
000098-87-3	BENZAL CHLORIDE
000098-95-3	NITROBENZENE
000099-08-1	NITROTOLUENE, M-
000099-09-2	NITROANILINE, M-
000099-35-4	TRINITROBENZENE, 1,3,5-
000099-55-8	METHYL-5-NITROANILINE, 2-
000099-59-2	METHOXY-5-NITROANILINE, 2-
000099-65-0	DINITROBENZENE, 1,3-
000099-99-0	NITROTOLUENE, P-
000100-00-5	CHLORONITROBENZENE, P-
000100-01-6	NITROANILINE, P-
000100-21-0	PHTHALIC ACID, P-
000100-21-0	TEREPHTHALIC ACID
000100-25-4	DINITROBENZENE, 1,4-
000100-40-3	VINYL-1-CYCLOHEXENE, 4-
000100-41-4	ETHYL BENZENE
000100-42-5	STYRENE
000100-44-7	BENZYL CHLORIDE
000100-51-6	BENZYL ALCOHOL
000100-52-7	BENZALDEHYDE
000100-54-9	NICOTINONITRILE
000101-14-4	METHYLENE-BIS(2-CHLOROANILINE), 4,4'-
000101-55-3	BROMOPHENYL PHENYL ETHER, 4-
000101-61-1	METHYLENE-BIS(N,N'-DIMETHYL)ANILINE, 4,4'-
000101-68-8	DIPHENYLMETHANE DIISOCYANATE
000101-68-8	METHYLENEDIPHENYL ISOCYANATE, 4,4-
000101-77-9	METHYLENE-BIS(BENZENEAMINE), 4,4-
000103-33-3	AZOBENZENE
000103-69-5	ETHYLANILINE, N-
000105-60-2	CAPROLACTAM
000105-67-9	DIMETHYLPHENOL, 2,4-
000106-13-8	ETHOXYETHANOL DODECANOATE, 2-
000106-37-6	DIBROMOBENZENE, 1,4-
000106-42-3	XYLENE, P-
000106-43-4	CHLOROTOLUENE, P-
000106-44-5	4-METHYLPHENOL
000106-44-5	CRESOL, P-
000106-46-7	DICHLOROBENZENE, 1,4-
000106-47-8	CHLOROANILINE, 4-
000106-48-9	CHLOROPHENOL, 4-
000106-49-0	TOLUIDINE, P-
000106-50-3	PHENYLENEDIAMINE, P-
000106-74-1	ETHOXYETHANOL ACRYLATE, 2-
000106-89-8	EPICHLOROHYDRIN
000106-93-4	DIBROMOETHANE, 1,2-
000106-99-0	BUTADIENE, 1,3-
000107-02-8	ACROLEIN
000107-05-1	ALLYL CHLORIDE
000107-12-0	PROPIONITRILE
000107-13-1	ACRYLONITRILE
000107-15-3	ETHYLENE DIAMINE
000107-16-4	FORMALDEHYDE CYANOHYDRIN
000107-18-6	ALLYL ALCOHOL
000107-20-0	CHLOROACETALDEHYDE
000107-21-1	ETHYLENE GLYCOL
000107-30-2	CHLOROMETHYL METHYL ETHER
000107-98-2	PROPYLENE GLYCOL MONOMETHYL ETHER
000108-05-4	VINYL ACETATE
000108-10-1	METHYL ISOBUTYL KETONE
000108-31-6	MALEIC ANHYDRIDE
000108-38-3	XYLENE, M-
000108-39-4	3-METHYLPHENOL
000108-39-4	CRESOL, M-
000108-41-8	CHLOROTOLUENE, M-
000108-42-9	CHLOROANILINE, 3-
000108-43-0	CHLOROPHENOL, 3-
000108-44-1	TOLUIDINE, M-
000108-45-2	PHENYLENEDIAMINE, M-
000108-60-1	BIS(2-CHLORO-1-METHYLETHYL) ETHER
000108-87-2	METHYLCYCLOHEXANE
000108-88-3	TOLUENE
000108-90-7	CHLOROBENZENE
000108-91-8	CYCLOHEXYLAMINE
000108-93-0	CYCLOHEXANOL
000108-95-2	PHENOL
000108-98-5	THIOPHENOL
000108-98-5	BENZENETHIOL
000109-66-0	PENTANE, N-
000109-69-3	CHLOROBUTANE, 1-
000109-77-3	MALONONITRILE
000109-78-4	ETHYLENE CYANOHYDRIN
000109-86-4	METHOXYETHANOL, 2-
000110-00-9	FURAN
000110-49-6	METHOXYETHANOL ACETATE, 2-
000110-54-3	HEXANE, N-
000110-61-2	SUCCINONITRILE
000110-80-5	ETHOXYETHANOL, 2-
000110-86-1	PYRIDINE
000111-15-9	ETHOXYETHANOL ACETATE, 2-
000111-44-4	BIS(2-CHLOROETHYL) ETHER
000111-69-3	ADIPONITRILE
000111-76-2	ETHYLENE GLYCOL MONOBUTYL ETHER
000111-90-0	DIETHYLENE GLYCOL MONOETHYL ETHER
000112-34-5	DIETHYLENE GLYCOL MONOBUTYL ETHER
000115-29-7	ENDOSULFAN
000116-06-3	ALDICARB
000117-81-7	BIS(2-ETHYLHEXYL) PHTHALATE
000117-81-7	DEHP
000117-84-0	DI-N-OCTYL PHTHALATE
000118-74-1	HEXACHLOROBENZENE
000118-75-2	CHLORANIL
000118-96-7	TRINITROTOLUENE, 2,4,6-
000119-90-4	DIMETHOXYBENZIDINE, 3,3'-
000119-93-7	DIMETHYLBENZIDINE, 3,3'-
000120-12-7	ANTHRACENE
000120-36-5	DICHLORPROP
000120-61-6	DIMETHYLTEREPHTHALATE
000120-82-1	TRICHLOROBENZENE, 1,2,4-
000120-83-2	DICHLOROPHENOL, 2,4-
000121-14-2	DINITROTOLUENE, 2,4
000121-69-7	DIMETHYLANILINE, N,N-
000121-73-3	CHLORONITROBENZENE, M-
000121-75-5	MALATHION
000121-82-4	CYCLONITE
000121-82-4	RDX
000121-91-5	PHTHALIC ACID, M-
000122-34-9	SIMAZINE
000122-39-4	DIPHENYLAMINE, N,N-
000122-66-7	DIPHENYLHYDRAZINE, 1,2-

000123-30-8	AMINOPHENOL, P-
000123-31-9	HYDROQUINONE
000123-33-1	MALEIC HYDRAZIDE
000123-63-7	PARALDEHYDE
000123-73-9	CROTONALDEHYDE
000123-91-1	DIOXANE, 1,4-
000124-09-4	HEXAMETHYLENE DIAMINE
000124-48-1	DIBROMOCHLOROMETHANE
000126-98-7	METHACRYLONITRILE
000126-99-8	CHLORO-1,3-BUTADIENE, 2-
000126-99-8	CHLOROPRENE
000127-18-4	TETRACHLOROETHYLENE
000129-00-0	PYRENE
000130-15-4	NAPHTHOQUINONE, 1,4-
000131-11-3	DIMETHYLPHTHALATE
000132-64-9	DIBENZOFURAN
000133-06-2	CAPTAN
000133-07-3	FOLPET
000137-26-8	THIRAM
000139-40-2	PROPAZINE
000140-57-8	ARAMITE
000140-88-5	ETHYL ACRYLATE
000141-78-6	ETHYL ACETATE
000142-28-9	DICHLOROPROPANE, 1,3-
000142-82-5	HEPTANE, N-
000143-33-9	SODIUM CYANIDE
000145-73-3	ENDOTHALL
000148-18-5	SODIUM DIETHYLDITHIOCARBAMATE
000150-50-5	MERPHOS
000151-50-8	POTASSIUM CYANIDE
000152-16-9	OCTAMETHYLPYROPHOSPHORAMIDE
000156-10-5	NITROSODIPHENYLAMINE, P-
000156-59-2	DICHLOROETHYLENE, 1,2-C-
000156-60-5	DICHLOROETHYLENE, 1,2-T-
000193-39-5	INDENO[1,2,3-CD]PYRENE
000206-44-0	FLUORANTHENE
000207-08-9	BENZO[K]FLUORANTHENE
000208-96-8	ACENAPTHYLENE
000218-01-9	CHRYSENE
000298-00-0	METHYL PARATHION
000298-02-2	PHORATE
000298-04-4	DISULFOTON
000299-84-3	RONNEL
000302-01-2	HYDRAZINE
000304-61-0	ANTIMONY POTASSIUM TARTRATE
000309-00-2	ALDRIN
000311-45-5	DIETHYL-P-NITROPHENYL PHOSPHATE
000319-84-6	HEXACHLOROCYCLOHEXANE, ALPHA-
000319-85-7	HEXACHLOROCYCLOHEXANE, BETA-
000319-86-8	HEXACHLOROCYCLOHEXANE, DELTA-
000329-71-5	DINITROPHENOL, 2,5-
000330-55-2	LINURON
000333-41-5	DIAZINON
000460-19-5	CYANOGEN
000479-45-8	TRINITROPHENYLMETHYLNITRAMINE
000496-72-0	TOLUENEDIAMINE, 3,4-
000504-24-5	AMINOPYRIDINE, 4-
000506-61-6	POTASSIUM SILVER CYANIDE
000506-64-9	SILVER CYANIDE
000506-68-3	CYANOGEN BROMIDE
000506-77-4	CHLORINE CYANIDE
000507-20-0	BUTYLCHLORIDE, T-
000510-15-6	CHLOROBENZILATE
000512-56-1	TRIMETHYL PHOSPHATE
000526-75-0	DIMETHYLPHENOL, 2,3-
000528-29-0	DINITROBENZENE, 1,2-
000531-82-8	FURIUM
000532-28-5	BENZALDEHYDE CYANOHYDRIN
000534-52-1	DINITRO-O-CRESOL, 4,6-
000540-59-0	DICHLOROETHYLENE, 1,2- (MIXED ISOMERS)
000540-73-8	DIMETHYLHYDRAZINE, 1,2-
000541-73-1	DICHLOROBENZENE, 1,3-
000542-62-1	BARIUM CYANIDE
000542-75-6	TELONE II
000542-75-6	DICHLOROPROPENE, 1,3-
000542-88-1	BIS(CHLOROMETHYL) ETHER
000542-92-7	CYCLOPENTADIENE
000544-92-3	COPPER CYANIDE
000557-19-7	NICKEL CYANIDE
000557-21-1	ZINC CYANIDE
000563-68-8	THALLIUM (I) ACETATE
000573-56-8	DINITROPHENOL, 2,6-
000576-24-9	DICHLOROPHENOL, 2,3-
000576-26-1	DIMETHYLPHENOL, 2,6-
000583-78-8	DICHLOROPHENOL, 2,5-
000586-11-8	DINITROPHENOL, 3,5-
000591-27-5	AMINOPHENOL, M-
000591-35-5	DICHLOROPHENOL, 3,5-
000591-78-6	HEXANONE, 2-
000592-01-8	CALCIUM CYANIDE
000593-60-2	VINYL BROMIDE
000593-60-2	BROMOETHENE
000594-20-7	DICHLOROPROPANE, 2,2-
000598-31-2	BROMOACETONE
000598-77-6	TRICHLOROPROPANE, 1,1,2-
000598-94-7	DIMETHYLUREA, N,N-
000602-01-7	DINITROTOLUENE, 2,3-
000606-20-2	DINITROTOLUENE, 2,6-
000608-73-1	HEXACHLOROCYCLOHEXANE-TECHNICAL
000608-93-5	PENTACHLOROBENZENE
000609-19-8	TRICHLOROPHENOL, 3,4,5-
000609-93-8	DINITRO-P-CRESOL, 2,6-
000610-39-9	DINITROTOLUENE, 3,4-
000611-14-3	ETHYLTOLUENE, O-
000615-54-3	TRIBROMOBENZENE, 1,2,4-
000617-84-5	DIETHYLFORMAMIDE
000619-15-8	DINITROTOLUENE, 2,5-
000620-14-4	ETHYLTOLUENE, M-
000621-64-7	NITROSO-DI-N-PROPYLAMINE, N-
000622-96-8	ETHYLTOLUENE, P-
000624-83-9	METHYL ISOCYANATE
000630-05-0	CARBON MONOXIDE
000630-10-4	SELENOUREA
000630-20-6	TETRACHLOROETHANE, 1,1,1,2-
000634-93-5	TRICHLOROANILINE, 2,4,6-
000636-21-5	METHYLANILINE HYDROCHLORIDE, 2-
000684-93-5	NITROSO-N-METHYLUREA, N-
000695-77-2	TETRACHLOROCYCLOPENTADIENE
000759-73-9	NITROSO-N-ETHYLUREA, N-
000759-94-4	EPTC
000764-41-0	DICHLORO-2-BUTENE, 1,4-
000765-34-4	GLYCIDALDEHYDE
000823-40-5	TOLUENE-2,6-DIAMINE
000834-12-8	AMETRYN
000924-16-3	NITROSO-DI-N-BUTYLAMINE, N-
000930-55-2	NITROSOPYRROLIDINE, N-
000933-75-5	TRICHLOROPHENOL, 2,3,6-
000933-78-8	TRICHLOROPHENOL, 2,3,5-
000934-80-5	ETHYL-O-XYLENE, 4-
000935-95-5	TETRACHLOROPHENOL, 2,3,5,6-
000950-10-7	MEPHOSFOLAN
000961-11-5	TETRACHLOROVINPHOS
000961-11-5	STIROPHOS
001024-57-3	HEPTACHLOR EPOXIDE
001114-71-2	PEBULATE
001116-54-7	NITROSODIETHANOLAMINE, N-
001134-23-2	CYCLOATE
001163-19-5	DECABROMODIPHENYL ETHER
001198-27-2	AMINO-2-NAPHTOL HYDROCHLORIDE, 1-
001309-64-4	ANTIMONY TRIOXIDE
001314-32-5	THALLIC OXIDE
001314-60-9	ANTIMONY PENTOXIDE
001314-62-1	VANADIUM PENTOXIDE
001314-84-7	ZINC PHOSPHIDE
001330-20-7	XYLENE, MIXTURE
001332-21-4	ASBESTOS
001332-81-6	ANTIMONY TETROXIDE

001336-36-3 POLYCHLORINATED BIPHENYLS
001338-23-4 METHYL ETHYL KETONE PEROXIDE
001563-66-2 CARBOFURAN
001569-02-4 PROPYLENE GLYCOL MONOETHYL ETHER
001582-09-8 TRIFLURALIN
001600-37-9 PENTACHLOROPROPENE, 1,1,2,3,3,-
001615-80-1 DIETHYLHYDRAZINE, 1,2-
001689-84-5 001689-99-2BROMOXYNIL OCTANOATE
001746-01-6 TCDD, 2,3,7,8-
001861-32-1 DACTHAL
001861-40-1 BENEFIN
001897-45-6 CHLOROTHALONIL
001912-24-9 ATRAZINE
001918-00-9 DICAMBA
001918-16-7 PROPACHLOR
001929-77-7 VERNOLATE
001929-77-7 VERNAM
001937-37-7 DIRECT BLACK 38
002008-41-5 BUTYLATE
002014-83-7 TRICHLOROTOLUENE, ALPHA,2,6-
002077-46-5 TRICHLOROTOLUENE, 2,3,6-
002104-96-3 BROMOPHOS
002212-67-1 MOLINATE
002303-16-4 DIALLATE
002303-17-5 TRIALLATE
002370-63-0 ETHOXYETHYL METHACRYLATE, 2-
002385-85-5 MIREX
002425-06-1 CAPTAFOL
002429-74-5 NIAGARA BLUE 4B
002491-38-5 BUSAN 90
002602-46-2 DIRECT BLUE 6
002610-05-1 DIRECT SKY BLUE 6B
002687-25-4 TOLUENEDIAMINE, 2,3-
002834-92-6 AMINO-2-NAPHTHOL, 1-
002921-88-2 CHLORPYRIFOS
003165-93-3
CHLORO-2-METHYLANILINE HYDROCHLORIDE, 4-
003175-23-3 TRICHLOROPROPANE, 1,2,2-
003380-34-5
TRICHLORO-2'-HYDROXYDIPHENYLETHER, 2,2,4'-
003383-96-8 TEMEPHOS
003689-24-5 TETRAETHYL DITHIOPYROPHOSPHATE
004399-55-7 DIRECT LIGHTFAST BLUE
004549-40-0 NITROSOMETHYLVINYLAMINE, N
004901-51-3 TETRACHLOROPHENOL, 2,3,4,5-
005216-25-1
TETRACHLOROTOLUENE, PARA, ALPHA, ALPHA, ALPHA-
005598-13-0 CHLORPYRIFOS METHYL
006108-10-7 HEXACHLOROCYCLOHEXANE, EPSILON-
006533-73-9 THALLIUM (I) CARBONATE
007429-90-5 ALUMINUM
007439-89-6 IRON
007439-92-1 LEAD
007439-96-5 MANGANESE
007439-97-6 MERCURY, ELEMENTAL
007439-98-7 MOLYBDENUM
007440-22-4 SILVER
007440-24-6 STRONTIUM, STABLE
007440-36-0 ANTIMONY, METALLIC
007440-38-2 ARSENIC, INORGANIC
007440-39-3 BARIUM
007440-41-7 BERYLLIUM
007440-42-8 BORON, ELEMENTAL
007440-43-9 CADMIUM
007440-50-8 COPPER
007440-62-2 VANADIUM
007440-66-6 ZINC (METALLIC)
007446-09-5 SULFUR DIOXIDE
007446-18-6 THALLIUM (I) SULFATE
007446-34-6 SELENIUM SULFIDE
007487-94-7 MERCURIC CHLORIDE
007637-07-2 BORON TRIFLUORIDE
007664-41-7 AMMONIA
007664-93-9 SULFURIC ACID
007723-14-0 PHOSPHORUS, WHITE
007782-41-4 FLUORINE / (SOLUBLE FLUORIDE)
007782-49-2 SELENIUM
007783-00-8 SELENIOUS ACID
007783-06-4 HYDROGEN SULFIDE
007789-89-1 TRICHLOROPROPANE, 1,1,1-
007791-12-0 THALLIUM (I) CHLORIDE
007803-51-2 PHOSPHINE
008001-35-2 TOXAPHENE
008001-58-9 CREOSOTE, COAL TAR
008007-45-2 COKE OVEN EMISSIONS
008018-01-7 MANCOZEB
010028-15-6 OZONE
010034-93-2 HYDRAZINE SULFATE
010102-43-9 NITRIC OXIDE
010102-44-0 NITROGEN DIOXIDE
010102-45-1 THALLIUM (I) NITRATE
010436-39-2 TETRACHLOROPROPENE, 1,1,2,3-
010595-95-6 NITROSOMETHYLETHYLAMINE, N-
010599-90-3 MONOCHLORAMINE
011097-69-1 AROCLOR 1254
012035-72-2 NICKEL SUBSULFIDE
012039-52-0 THALLIUM SELENITE
012122-67-7 ZINEB
012427-38-2 MANEB
012672-29-6 AROCLOR 1248
013071-79-9 TERBUFOS
013194-48-4 ETHOPROP
013196-18-4 THIOFANOX
013718-26-8 SODIUM METAVANADATE
014797-65-0 NITRITE
015950-66-0 TRICHLOROPHENOL, 2,3,4-
015972-60-8 ALACHLOR
016065-83-1 CHROMIUM(III)
016071-86-6 DIRECT BROWN 95
016752-77-5 METHOMYL
018540-29-9 CHROMIUM(VI)
020816-12-0 OSMIUM TETROXIDE
020859-73-8 ALUMINUM PHOSPHIDE
021087-64-9 METRIBUZIN
021232-47-3 TETRACHLOROAZOXYBENZENE
021436-96-4 DIMETHYLANILINE HYDROCHLORIDE, 2,4-
021564-17-0
THIOCYANOMETHYLTHIO)BENZOTHIAZOLE, 2-(
021725-46-2 CYANAZINE
022967-92-6 METHYL MERCURY
023950-58-5 PRONAMIDE
025013-15-4 METHYL STYRENE (MIXED ISOMERS)
025329-35-5 PENTACHLOROCYCLOPENTADIENE
026399-36-0 PROFLURALIN
030560-19-1 ACEPHATE
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
VARIOUS	BROMINATED DIBENZOFURANS
NO CASRN	BROMINATED DIBENZO-P-DIOXINS
NO CASRN	BROMOCHLOROETHANES
NO CASRN	DICHLOROBUTENES
NO CASRN	LEAD ALKYLS
NO CASRN	NICKEL, REFINERY DUST
VARIOUS	NICKEL, SOLUBLE SALTS
NO CASRN	NITROGEN OXIDES
NO CASRN	NITROPHENOLS
NO CASRN	PARTICULATE MATTER
NO CASRN	PHOTOCHEMICAL OXIDANTS
NO CASRN	POLYBROMINATED BIPHENYLS
NO CASRN	SULFUR OXIDES
NO CASRN	THALLIUM, INSOLUBLE SALTS
NO CASRN	TIN AND COMPOUNDS
NO CASRN	TRIMETHYLBENZENES
NO CASRN	TRINITROPHENOLS

APPENDIX A-IV

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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. National Center for Environmental Assessment, Cincinnati, OH.

APPENDIX A-V

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 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_2), Particulate Matter, less than 10 µm in size, (PM_{10}), Ozone (O_3) and Sulfur Dioxide (SO_2). 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[a]
(as of December 2, 1991)

Pollutant	Primary Standards[b]	Averaging Time	Secondary Standards[b]
Carbon monoxide (CO)	9 ppm (10 mg/m^3) 35 ppm (40 mg/m^3)	8 hour[c] 1 hour[c]	None
Lead (Pb) (and Lead compounds	1.5 µg/m^3	Quarterly	Same as primary
Nitrogen dioxide (NO_2) (Nitrogen oxide) (Nitric oxide)	0.053 ppm (100 µg/m^3)	Annual	Same as primary
Particulate Matter (PM_{10})	50 µg/m^3 150 µg/m^3	Annual[d] 24 hours[e]	Same as primary
Ozone (O_3)	0.12 ppm (235 µg/m^3)	1 hour[f]	Same as primary
Sulfur dioxide (SO_2) (Sulfur oxide)	0.03 ppm (80 µg/m^3)	Annual	---
	0.14 ppm (365 µg/m^3)	24 hours[c]	---
	---	3 hours[c]	0.5 ppm (1300 µg/m^3)

[a]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.

[b]Primary standards are designed to protect public health; Secondary standards are designed to protect public welfare.

[c]Not to be exceeded more than once per year.

[d]The standard is attained when the expected annual arithmetic mean concentration is less than or equal to 50 µg/m^3.

[e]The standard is attained when the expected number of days per calendar year with a 24-hour average concentration above 150 µg/m^3 is equal to or less than 1.

[f]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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