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IMMEDIATELY DANGEROUS TO LIFE OR HEALTH (IDLH) VALUE PROFILE

FOR

DIKETENE

[CAS No. 674-82-8]

Department of Health and Human Services
Centers for Disease Control and Prevention
National Institute for Occupational Safety and Health

**External Review Draft
March 2015**

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1 Foreword

2 Chemicals are a ubiquitous component of the modern workplace. Occupational exposures to chemicals have the
3 potential to adversely affect the health and lives of workers. Acute or short-term exposures to high concentrations
4 of some airborne chemicals have the ability to quickly overwhelm workers, resulting in a spectrum of undesirable
5 health outcomes that may inhibit the ability to escape from the exposure environment (e.g., irritation of the eyes
6 and respiratory tract or cognitive impairment), cause severe irreversible effects (e.g., damage to the respiratory
7 tract or reproductive toxicity), and in extreme cases, cause death. Airborne concentrations of chemicals capable
8 of causing such adverse health effects or of impeding escape from high-risk conditions may arise from a variety of
9 non-routine workplace situations, including special work procedures (e.g., in confined spaces), industrial
10 accidents (e.g., chemical spills or explosions), and chemical releases into the community (e.g., during
11 transportation incidents or other uncontrolled-release scenarios).

12
13 The “immediately dangerous to life or health air concentration values (IDLH values)” developed by the National
14 Institute for Occupational Safety and Health (NIOSH) characterize these high-risk exposure concentrations and
15 conditions [NIOSH 2013]. IDLH values are based on a 30-minute exposure duration and have traditionally
16 served as a key component of the decision logic for the selection of respiratory protection devices [NIOSH 2004].
17 Occupational health professionals have employed these values beyond their initial purpose as a component of the
18 NIOSH Respirator Selection Logic to assist in developing Risk Management Plans for non-routine work practices
19 governing operations in high-risk environments (e.g., confined spaces) and the development of Emergency
20 Preparedness Plans.

21
22 The approach used to derive IDLH values for high priority chemicals is outlined in the NIOSH Current
23 Intelligence Bulletin (CIB) 66: Derivation of Immediately Dangerous to Life or Health Values [NIOSH 2013].
24 CIB 66 provides 1) an update on the scientific basis and risk assessment methodology used to derive IDLH
25 values, 2) the rationale and derivation process for IDLH values, and 3) a demonstration of the derivation of
26 scientifically credible IDLH values using available data resources.

27
28 The purpose of this technical report is to present the IDLH value for diketene (CAS # 674-82-8). The scientific
29 basis, toxicologic data and risk assessment approach used to derive the IDLH value are summarized to ensure
30 transparency and scientific credibility.

31
32 John Howard, M.D.
33 Director
34 National Institute for Occupational Safety and Health

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1 Abbreviations

2		
3	ACGIH	American Conference of Governmental Industrial Hygienists
4	AEGL	Acute Exposure Guideline Levels
5	AIHA	American Industrial Hygiene Association
6	BMC	benchmark concentration
7	BMCL	benchmark concentration lower confidence limit
8	C	ceiling
9	CAS	chemical abstract service
10	ERPG	Emergency Response Planning Guidelines
11	IDLH	immediately dangerous to life or health
12	kPa	kilopascal
13	LC ₅₀	median lethal concentration
14	LC _{Lo}	lowest concentration of a chemical that caused death in humans or animals
15	LEL	lower explosive limit
16	LOAEL	lowest observed adverse effect level
17	mg/m ³	milligram(s) per cubic meter
18	NAC	National Advisory Committee
19	NAS	National Academy of Sciences
20	NIOSH	National Institute for Occupational Safety and Health
21	NOAEL	no observed adverse effect level
22	OSHA	Occupational Safety and Health Administration
23	PEL	permissible exposure limit
24	ppm	parts per million
25	RD ₅₀	concentration of a chemical in the air that is estimated to cause a 50% decrease in the respiratory rate
26		
27	REL	recommended exposure limit
28	SCP	Standard Completion Program
29	STEL	short term exposure limit
30	TLV	threshold limit value
31	TWA	time weighted average
32	UEL	upper explosive limit
33	WEEL	workplace environmental exposure level

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1 **Glossary**

2
3 **Acute Exposure:** Exposure by the oral, dermal, or inhalation route for 24 hours or less.

4 **Acute Exposure Guideline Levels (AEGLs):** Threshold exposure limits for the general public applicable to
5 emergency exposure periods ranging from 10 minutes to 8 hours. AEGL-1, AEGL 2, and AEGL-3 are
6 developed for five exposure periods (10 and 30 minutes, 1 hour, 4 hours, and 8 hours) and are distinguished
7 by varying degrees of severity of toxic effects ranging from transient, reversible effects to life-threatening
8 effects [NAS 2001]. AEGLs are intended to be guideline levels used during rare events or single once-in-a-
9 lifetime exposures to airborne concentrations of acutely toxic, high-priority chemicals [NAS 2001]. The
10 threshold exposure limits are designed to protect the general population, including the elderly, children or
11 other potentially sensitive groups that are generally not considered in the development of workplace exposure
12 recommendations (additional information available at <http://www.epa.gov/oppt/aegl/>).

13 **Acute Reference Concentration (RfC):** An estimate (with uncertainty spanning perhaps an order of magnitude)
14 of a continuous inhalation exposure for an acute duration (24 hours or less) of the human population
15 (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a
16 lifetime. It can be derived from a NOAEL, LOAEL, or benchmark concentration, with uncertainty factors
17 (UFs) generally applied to reflect limitations of the data used. Generally used in USEPA noncancer health
18 assessments [USEPA 2014].

19 **Acute Toxicity:** Any poisonous effect produced within a short period of time following an exposure, usually 24
20 to 96 hours.

21 **Adverse Effect:** A substance-related biochemical change, functional impairment, or pathologic lesion that affects
22 the performance of an organ or system or alters the ability to respond to additional environmental challenges.

23 **Benchmark Dose/Concentration (BMD/BMC):** A dose or concentration that produces a predetermined change
24 in response rate of an effect (called the benchmark response, or BMR) compared to background [USEPA
25 2014] (additional information available at <http://www.epa.gov/ncea/bmds/>).

26 **Benchmark Response (BMR):** A predetermined change in response rate of an effect. Common defaults for the
27 BMR are 10% or 5%, reflecting study design, data variability, and sensitivity limits used.

28 **BMCL:** A statistical lower confidence limit on the concentration at the BMC [USEPA 2014].

29 **Bolus Exposure:** A single, relatively large dose.

30 **Ceiling Value ("C"):** U.S. term in occupational exposure indicating the airborne concentration of a potentially
31 toxic substance that should never be exceeded in a worker's breathing zone.

32 **Chronic Exposure:** Repeated exposure for an extended period of time. Typically exposures are more than
33 approximately 10% of life span for humans and >90 days to 2 years for laboratory species.

34 **Critical Study:** The study that contributes most significantly to the qualitative and quantitative assessment of risk
35 [USEPA 2014].

36
37 **Dose:** The amount of a substance available for interactions with metabolic processes or biologically significant
38 receptors after crossing the outer boundary of an organism [USEPA 2014].

39 **EC₅₀:** A combination of the effective concentration of a substance in the air and the exposure duration that is
40 predicted to cause an effect in 50% (one half) of the experimental test subjects.

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- 1 **Emergency Response Planning Guidelines (ERPGs):** Maximum airborne concentrations below which nearly all
2 individuals can be exposed without experiencing health effects for 1-hour exposure. ERPGs are presented in a
3 tiered fashion with health effects ranging from mild or transient to serious, irreversible, or life threatening
4 (depending on the tier). ERPGs are developed by the American Industrial Hygiene Association [AIHA 2006].
- 5 **Endpoint:** An observable or measurable biological event or sign of toxicity ranging from biomarkers of initial
6 response to gross manifestations of clinical toxicity.
- 7 **Exposure:** Contact made between a chemical, physical, or biological agent and the outer boundary of an
8 organism. Exposure is quantified as the amount of an agent available at the exchange boundaries of the
9 organism (e.g., skin, lungs, gut).
- 10 **Extrapolation:** An estimate of the response at a point outside the range of the experimental data, generally
11 through the use of a mathematical model, although qualitative extrapolation may also be conducted. The
12 model may then be used to extrapolate to response levels that cannot be directly observed.
- 13 **Hazard:** A potential source of harm. Hazard is distinguished from risk, which is the probability of harm under
14 specific exposure conditions.
- 15 **Immediately Dangerous to Life or Health (IDLH) condition:** A situation that poses a threat of exposure to
16 airborne contaminants when that exposure is likely to cause death or immediate or delayed permanent adverse
17 health effects or prevent escape from such an environment [NIOSH 2004, 2013].
- 18 **IDLH value:** A maximum (airborne concentration) level above which only a highly reliable breathing apparatus
19 providing maximum worker protection is permitted [NIOSH 2004, 2013]. IDLH values are based on a 30-
20 minute exposure duration.
- 21 **LC₀₁:** The statistically determined concentration of a substance in the air that is estimated to cause death in 1% of
22 the test animals.
- 23 **LC₅₀:** The statistically determined concentration of a substance in the air that is estimated to cause death in 50%
24 (one half) of the test animals; median lethal concentration.
- 25 **LC_{L0}:** The lowest lethal concentration of a substance in the air reported to cause death, usually for a small
26 percentage of the test animals.
27
- 28 **LD₅₀:** The statistically determined lethal dose of a substance that is estimated to cause death in 50% (one half) of
29 the test animals; median lethal concentration.
- 30 **LD_{L0}:** The lowest dose of a substance that causes death, usually for a small percentage of the test animals.
- 31 **LEL:** The minimum concentration of a gas or vapor in air, below which propagation of a flame does not occur in
32 the presence of an ignition source.
- 33 **Lethality:** Pertaining to or causing death; fatal; referring to the deaths resulting from acute toxicity studies. May
34 also be used in lethality threshold to describe the point of sufficient substance concentration to begin to cause
35 death.
- 36 **Lowest Observed Adverse Effect Level (LOAEL):** The lowest tested dose or concentration of a substance that
37 has been reported to cause harmful (adverse) health effects in people or animals.

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- 1 **Mode of Action:** The sequence of significant events and processes that describes how a substance causes a toxic
2 outcome. Mode of action is distinguished from the more detailed mechanism of action, which implies a more
3 detailed understanding on a molecular level.
- 4 **No Observed Adverse Effect Level (NOAEL):** The highest tested dose or concentration of a substance that has
5 been reported to cause no harmful (adverse) health effects in people or animals.
- 6 **Occupational Exposure Limit (OEL):** Workplace exposure recommendations developed by governmental
7 agencies and non-governmental organizations. OELs are intended to represent the maximum airborne
8 concentrations of a chemical substance below which workplace exposures should not cause adverse health
9 effects. OELs may apply to ceiling, short-term (STELs), or time-weighted average (TWA) limits.
- 10 **Peak Concentration:** Highest concentration of a substance recorded during a certain period of observation.
- 11 **Permissible Exposure Limit (PEL):** Occupational exposure limits developed by OSHA (29 CFR 1910.1000) or
12 MSHA (30 CFR 57.5001) for allowable occupational airborne exposure concentrations. PELs are legally
13 enforceable and may be designated as ceiling, STEL, or TWA limits.
- 14
- 15 **Point of Departure (POD):** The point on the dose–response curve from which dose extrapolation is initiated.
16 This point can be the lower bound on dose for an estimated incidence or a change in response level from a
17 concentration-response model (BMC), or it can be a NOAEL or LOAEL for an observed effect selected from
18 a dose evaluated in a health effects or toxicology study.
- 19 **RD₅₀:** The statistically determined concentration of a substance in the air that is estimated to cause a 50% (one
20 half) decrease in the respiratory rate.
- 21 **Recommended Exposure Limit (REL):** Recommended maximum exposure limit to prevent adverse health
22 effects based on human and animal studies and established for occupational (up to 10-hour shift, 40-hour
23 week) inhalation exposure by NIOSH. RELs may be designated as ceiling, STEL, or TWA limits.
- 24 **Short-Term Exposure Limit (STEL):** A worker’s 15-minute time-weighted average exposure concentration that
25 shall not be exceeded at any time during a work day.
- 26 **Target Organ:** Organ in which the toxic injury manifests in terms of dysfunction or overt disease.
- 27 **Threshold Limit Values (TLVs®):** Recommended guidelines for occupational exposure to airborne
28 contaminants, published by the American Conference of Governmental Industrial Hygienists (ACGIH). TLVs
29 refer to airborne concentrations of chemical substances and represent conditions under which it is believed
30 that nearly all workers may be repeatedly exposed, day after day, over a working lifetime, without adverse
31 effects. TLVs may be designated as ceiling, short-term (STELs), or 8-hr TWA limits.
- 32 **Time-Weighted Average (TWA):** A worker’s 8-hour (or up to 10-hour) time-weighted average exposure
33 concentration that shall not be exceeded during an 8-hour (or up to 10-hour) work shift of a 40-hour week.
34 The average concentration is weighted to take into account the duration of different exposure concentrations.
- 35 **Toxicity:** The degree to which a substance is able to cause an adverse effect on an exposed organism.
- 36
- 37 **Uncertainty Factors (UFs):** Mathematical adjustments applied to the POD when developing IDLH values. The
38 UFs for IDLH value derivation are determined by considering the study and effect used for the POD, with
39 further modification based on the overall database.

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- 1 **Workplace Environmental Exposure Levels (WEELs):** Exposure levels developed by the American Industrial
- 2 Hygiene Association (AIHA) that provide guidance for protecting most workers from adverse health
- 3 effects related to occupational chemical exposures expressed as a TWA or ceiling limit.

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2

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24 University

25

1.0 Introduction

1.1 Overview of the IDLH Value for Diketene

IDLH Value: 7.6 ppm

Basis for IDLH Value: The IDLH value for diketene is based on the rat BMCL₀₅ of 181 ppm for lethality in a 1-hour exposure [Katz 1987]. The duration adjusted BMCL₀₅ for a 30-minute exposure is 228 ppm. A composite uncertainty factor of 30 was applied to account for extrapolation from a lethal concentration threshold in animals, animal to human differences, human variability, and database uncertainties, resulting in an IDLH value of **7.6 ppm**.

1.2 Purpose

This *IDLH Value Profile* presents (1) a brief summary of technical data associated with acute inhalation exposures to diketene and (2) the rationale behind the Immediately Dangerous to Life or Health (IDLH) value for diketene. IDLH values are developed based on the scientific rationale and logic outlined in the NIOSH Current Intelligence Bulletin (CIB) 66: Derivation of Immediately Dangerous to Life or Health (IDLH) values [NIOSH 2013]. As described in CIB 66, NIOSH performs in-depth literature searches to ensure that all relevant data from human and animal studies with acute exposures to the substance are identified. Information included in CIB 66 on the literature search includes pertinent databases, key terms, and guides for evaluating data quality and relevance for the establishment of an IDLH value. The information that is identified in the in-depth literature search is evaluated with general considerations that include description of studies (i.e., species, study protocol, exposure concentration and duration), health endpoint evaluated, and critical effect levels (e.g., NOAELs, LOAELs, LC₅₀ values). For diketene, the in-depth literature search was conducted through February 2014.

1.3 General Substance Information

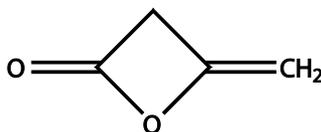
Chemical: Diketene

CAS No: 674-82-8

Synonyms: Acetyl ketene; 4-Methylene-2-oxetanone; But-3-en-3-olide; Dimer ketene *

Chemical category: Lactones[†]

1
2 **Structural formula:**



7 Table 1 highlights selected physiochemical properties of diketene relevant to IDLH conditions. Table 2 provides
8 alternative exposure guidelines for diketene. Table 3 summarizes the Acute Exposure Guidelines Level (AEGL)
9 values for diketene.

10
11 **Table 1: Physiochemical Properties of Diketene**
12

Property	Value
Molecular weight	84.04 [‡]
Chemical formula	C ₄ H ₄ O ₂
Description	Colorless to light-colored liquid
Odor	Pungent
Odor Threshold	Not available
UEL	11.7% [†]
LEL	2% [†]
Vapor pressure	1.07 kPa at 20°C (68°F) [‡]
Flash point	34°C (93.2°F) [‡]
Ignition temperature	275°C (527°F) [‡]
Solubility	Hydrolysis [‡]

13 **Abbreviation:** °C – Celsius; °F – Fahrenheit; kPa – kilopascal; LEL – lower explosive limit; UEL – upper explosive limit

14 * NLM [2014]

15 † IFA [2014]

16 ‡ HSDB [2014]

17
18 **Table 2: Alternative Exposure Guidelines for Diketene**
19

Organization	Value
Original (SCP) IDLH value	None
NIOSH REL	Not available
OSHA PEL [2014]	Not available
ACGIH TLV [2014]	Not available
AIHA ERPG [2010]	ERPG-1: 1 ppm; ERPG-2: 5 ppm; ERPG-3: 20 ppm
AIHA WEEL [2010]	Not available

20 **Abbreviation:** ACGIH – American Conference of Governmental Industrial Hygienists; AIHA – American Industrial Hygiene
21 Association; ERPG – Emergency Response Preparedness Guidelines; IDLH – immediately dangerous to life or health; NIOSH – National
22 Institute for Occupational Safety and Health; OSHA – Occupational Safety and Health Administration; PEL – permissible exposure limit;
23 REL – recommended exposure limit; SCP – Standard Completion Program; WEEL – workplace environmental exposure level

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1 **Table 3: AEGL Values for Diketene**
2

Classification	10-min	30-min	1-hour	4-hour	8-hour	Endpoint [reference]
AEGL-1	NR	NR	NR	NR	NR	Insufficient data
AEGL-2	11 ppm 38.0 mg/m ³	7.7 ppm 26.0 mg/m ³	6.0 ppm 21.0 mg/m ³	1.5 ppm 5.2 mg/m ³	0.77 ppm 2.6 mg/m ³	AEGL-3 reduced by factor of 3
AEGL-3	33.0 ppm 110.0 mg/m ³	23.0 ppm 79.0 mg/m ³	18.0 ppm 62.0 mg/m ³	4.5 ppm 15.0 mg/m ³	2.3 ppm 7.9 mg/m ³	BMCL ₀₅ for lethality [Katz 1987]

3 **Abbreviation:** AEGL – acute exposure guideline levels; mg/m³ – milligrams per cubic meter; min – minute; NR – not recommended due to insufficient data; ppm – parts per million

4 ***References:** NAS [2008]

2.0 Animal Toxicity Data

The available acute lethality database for diketene is limited to one rat study [Katz 1987] and one multi-species study [Wooster et al. 1947]. Katz [1987] exposed rats to 0, 250, 500, or 750 ppm diketene for one hour with a 14-day post-exposure period. Clinical signs providing evidence of severe irritation included excessive tearing and gasping at all exposure levels, and porphyrin discharge from the nose at the two highest concentrations. No gross lesions were identified during necropsy. Mortality occurred at all concentrations except 250 ppm. Modeling of the mortality responses yielded an LC_{50} of 612 ppm. The lethality thresholds were estimated using benchmark concentration modeling, resulting in a $BMCL_{05}$ (lower bound estimate of the concentration associated with a benchmark response of 5%) of 181 ppm for lethality. The 30-minute duration adjusted equivalent value of the $BMCL_{05}$ is 228 ppm.

Wooster et al. [1947] exposed mice to 194, 580, or 870 ppm diketene in an acetone solvent for 10 minutes with a 15-day post-exposure period. Clinical signs were not reported. Pathological examination revealed proteinaceous edematous fluid in the alveoli and in the perivascular connective tissue of the bronchi. Only the highest concentration (870 ppm) resulted in mortality, one mouse died. Wooster et al. [1947] also exposed guinea pigs to 194 ppm diketene for 10 minutes with a 15-day post-exposure observation period. No clinical signs were reported. The authors implied that the gross pathology findings indicated pulmonary edema as the cause of death. All guinea pigs in 194 ppm group died. Wooster et al. [1947] also exposed rats and rabbits to 194 ppm diketene for 10 minutes. No lethality, clinical signs, or gross pathology findings were noted.

Table 4 summarizes the LC data identified in animal studies and provides 30-minute equivalent derived values for diketene. Information in these tables includes species of test animals, toxicological metrics (i.e., LC, NOAEL, LOAEL), adjusted 30-minute concentration, and the justification for the composite uncertainty factors applied to calculate the derived values.

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1 **Table 4: Lethal Concentration Data for Diketene**

2

Reference	Species (sex)	LC ₅₀ (ppm)	Other lethality concentration (ppm)	Time (min)	Adjusted 30-min Concentration* (ppm)	Composite Uncertainty Factor	Derived Value (ppm) [†]
Katz [1987]	Rats (male and female)	612	--	60	771	30 [±]	25.7
Katz [1987]	Rats		370 [^]	60	466	30 ⁺	15.5
Katz [1987]	Rats		181[‡]	60	228	30⁺	7.6

3
4 **Abbreviation:** LC – lethal concentration; LC₅₀ – median lethal concentration; LC_{Lo} – lowest concentration of a chemical that caused death in humans or animals; min – minute; ppm – parts per million

5
6
7 * For exposures other than 30 minutes the ten Berge et al. [1986] relationship is used for duration adjustment ($C^n \times t = k$); no empirically estimated n values were available, therefore the default values were used, n = 3 for exposures greater than 30 minutes and n = 1 for exposures less than 30 minutes.

8
9 [†] The derived value is the result of the adjusted 30-minute concentration divided by the composite uncertainty factor.

10 [±]Composite uncertainty factor to account for adjustment of LC₅₀ values to LC₀₁ values, use of lethal concentration threshold in animals, interspecies differences and human variability.

11
12 [^]LC₁₀

13 ⁺Composite uncertainty factor to account for adjustment from a lethal concentration threshold in animals, interspecies differences, human variability and database uncertainties.

14
15 [‡] Lower bound estimate of the concentration associated with a benchmark response of 5% for lethality.

1 **3.0 Human Data**

2 Only very limited information on human effects and effect levels was available. Occupational exposure to
3 diketene was reported to cause mild irritation to the conjunctiva and mucosa of the nose and throat at a
4 concentration of 0.58 ppm for one minute [Fel'dman 1967]. These studies were not appropriate for an IDLH
5 value derivation due to the very short duration and limited details available.

6 **4.0 Summary**

7 The IDLH value for diketene is based on the rat BMCL₀₅ of 181 ppm for lethality in a 1-hour exposure [Katz
8 1987]. The duration adjusted BMCL₀₅ for a 30-minute exposure is 228 ppm. Application of composite
9 uncertainty factor of 30 to account for extrapolation from a lethal concentration threshold in animals, animal to
10 human differences, human variability, and database uncertainties, including uncertainty regarding the threshold
11 for severe irritation, yielded an IDLH value of **7.6 ppm**.

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