IDLH

IMMEDIATELY DANGEROUS to LIFE or HEALTH

VALUE PROFILE

n-Butyl Acrylate
CAS® No. 141-32-2

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Center for Disease Control and Prevention
National Institute of Occupational Safety and Health

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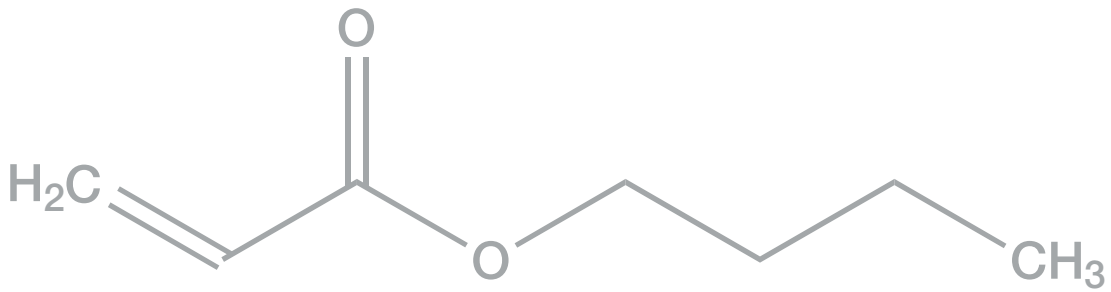
Immediately Dangerous to Life or Health (IDLH) Value Profile

n-Butyl Acrylate

[CAS® No. 141-32-2]



DEPARTMENT OF HEALTH AND HUMAN SERVICES
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September 2016

Foreword

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

The immediately dangerous to life or health (IDLH) air concentration values developed by the National Institute for Occupational Safety and Health (NIOSH) characterize these high-risk exposure concentrations and conditions [NIOSH 2013]. IDLH values are based on a 30-minute exposure duration and have traditionally served as a key component of the decision logic for the selection of respiratory protection devices [NIOSH 2004].

Occupational health professionals have employed these values beyond their initial purpose as a component of the NIOSH Respirator Selection Logic to assist in developing risk management plans for nonroutine work practices governing operations in high-risk environments (e.g., confined spaces) and the development of emergency preparedness plans.

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

The purpose of this technical report is to present the IDLH value for n-butyl acrylate (CAS® #141-32-2). The scientific basis, toxicologic data, and risk assessment approach used to derive the IDLH value are summarized to ensure transparency and scientific credibility.

John Howard, M.D.
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Contents

Abbreviations

ACGIH®	American Conference of Governmental Industrial Hygienists
AEGLs	Acute Exposure Guideline Levels
AIHA®	American Industrial Hygiene Association
BMC	benchmark concentration
BMD	benchmark dose
BMCL	benchmark concentration lower confidence limit
C	ceiling value
°C	degrees Celsius
CAS®	Chemical Abstracts Service, a division of the American Chemical Society
ERPGs™	Emergency Response Planning Guidelines
°F	degrees Fahrenheit
IDLH	immediately dangerous to life or health
LC_{50}	median lethal concentration
LC_{LO}	lowest concentration that caused death in humans or animals
LEL	lower explosive limit
LOAEL	lowest observed adverse effect level
mg/m^3	milligram(s) per cubic meter
min	minutes
mmHg	millimeter(s) of mercury
NAC	National Advisory Committee
NAS	National Academy of Sciences
NIOSH	National Institute for Occupational Safety and Health
NOAEL	no observed adverse effect level
NOEL	no observed effect level
NR	not recommended
OSHA	Occupational Safety and Health Administration
PEL	permissible exposure limit
ppm	parts per million
RD_{50}	concentration of a chemical in the air that is estimated to cause a 50% decrease in the respiratory rate
REL	recommended exposure limit
SCP	Standards Completion Program (joint effort of NIOSH and OSHA)
STEL	short-term exposure limit
TLV®	Threshold Limit Value
TWA	time-weighted average
UEL	upper explosive limit
WEELs®	Workplace Environmental Exposure Levels
µg/kg	microgram(s) per kilogram of body weight

Glossary

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

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

Acute reference concentration (Acute RfC): An estimate (with uncertainty spanning perhaps an order of magnitude) of a continuous inhalation exposure for an acute duration (24 hours or less) of the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime. It can be derived from a NOAEL, LOAEL, or benchmark concentration, with uncertainty factors (UFs) generally applied to reflect limitations of the data used. Generally used in U.S. EPA noncancer health assessments [U.S. EPA 2016].

Acute toxicity: Any poisonous effect produced within a short period of time following an exposure, usually 24 to 96 hours [U.S. EPA 2016].

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

Benchmark dose/concentration (BMD/BMC): A dose or concentration that produces a predetermined change in response rate of an effect (called the benchmark response, or BMR) compared to background [U.S. EPA 2016] (additional information available at http://www.epa.gov/ncea/bmds/).

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

BMCL: A statistical lower confidence limit on the concentration at the BMC [U.S. EPA 2016].

Bolus exposure: A single, relatively large dose.

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

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

Critical study: The study that contributes most significantly to the qualitative and quantitative assessment of risk [U.S. EPA 2016].

Dose: The amount of a substance available for interactions with metabolic processes or biologically significant receptors after crossing the outer boundary of an organism [U.S. EPA 2016].

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

Emergency Response Planning Guidelines (ERPGs™): Maximum airborne concentrations below which nearly all individuals can be exposed without experiencing health effects for 1-hour exposure. ERPGs are presented in a tiered fashion with health effects ranging from mild or transient to serious, irreversible, or life threatening (depending on the tier). ERPGs are developed by the American Industrial Hygiene Association [AIHA 2006].

Endpoint: An observable or measurable biological event or sign of toxicity, ranging from biomarkers of initial response to gross manifestations of clinical toxicity.

Exposure: Contact made between a chemical, physical, or biological agent and the outer boundary of an organism. Exposure is quantified as the amount of an agent available at the exchange boundaries of the organism (e.g., skin, lungs, and gut).

Extrapolation: An estimate of the response at a point outside the range of the experimental data, generally through the use of a mathematical model, although qualitative extrapolation may also be conducted. The model may then be used to extrapolate to response levels that cannot be directly observed.

Hazard: A potential source of harm. Hazard is distinguished from risk, which is the probability of harm under specific exposure conditions.

Immediately dangerous to life or health (IDLH) condition: A condition that poses a threat of exposure to airborne contaminants when that exposure is likely to cause death or immediate or delayed permanent adverse health effects or prevent escape from such an environment [NIOSH 2004, 2013].

IDLH value: A maximum (airborne concentration) level above which only a highly reliable breathing apparatus providing maximum worker protection is permitted [NIOSH 2004, 2013]. IDLH values are based on a 30-minute exposure duration.

LC_{01}: The statistically determined concentration of a substance in the air that is estimated to cause death in 1% of the test animals.

LC_{50}: The statistically determined concentration of a substance in the air that is estimated to cause death in 50% (one half) of the test animals; median lethal concentration.

LC_{LO}: The lowest lethal concentration of a substance in the air reported to cause death, usually for a small percentage of the test animals.

LD_{50}: The statistically determined lethal dose of a substance that is estimated to cause death in 50% (one half) of the test animals; median lethal concentration.

LD_{LO}: The lowest dose of a substance that causes death, usually for a small percentage of the test animals.

LEL: The minimum concentration of a gas or vapor in air, below which propagation of a flame does not occur in the presence of an ignition source.

Lethality: Pertaining to or causing death; fatal; referring to the deaths resulting from acute toxicity studies. May also be used in lethality threshold to describe the point of sufficient substance concentration to begin to cause death.

Lowest observed adverse effect level (LOAEL): The lowest tested dose or concentration of a substance that has been reported to cause harmful (adverse) health effects in people or animals.

Mode of action: The sequence of significant events and processes that describes how a substance causes a toxic outcome. By contrast, the term *mechanism of action* implies a more detailed understanding on a molecular level.

No observed adverse effect level (NOAEL): The highest tested dose or concentration of a substance that has been reported to cause no harmful (adverse) health effects in people or animals.

Occupational exposure limit (OEL): Workplace exposure recommendations developed by governmental agencies and nongovernmental organizations. OELs are intended to represent the maximum airborne concentrations of a chemical substance below which workplace exposures should not cause adverse health effects. OELs may apply to ceiling limits, STELs, or TWA limits.

Peak concentration: Highest concentration of a substance recorded during a certain period of observation.

Permissible exposure limits (PELs): Occupational exposure limits developed by OSHA (29 CFR 1910.1000) or MSHA (30 CFR 57.5001) for allowable occupational airborne exposure concentrations. PELs are legally enforceable and may be designated as ceiling limits, STELs, or TWA limits.

Point of departure (POD): The point on the dose–response curve from which dose extrapolation is initiated. This point can be the lower bound on dose for an estimated incidence or a change in response level from a concentration-response model (BMC), or it can be a NOAEL or LOAEL for an observed effect selected from a dose evaluated in a health effects or toxicology study.

RD_{50}: The statistically determined concentration of a substance in the air that is estimated to cause a 50% (one half) decrease in the respiratory rate.

Recommended exposure limit (REL): Recommended maximum exposure limit to prevent adverse health effects based on human and animal studies and established for occupational (up to 10-hour shift, 40-hour week) inhalation exposure by NIOSH. RELs may be designated as ceiling limits, STELs, or TWA limits.

Short-term exposure limit (STEL): A worker's 15-minute time-weighted average exposure concentration that shall not be exceeded at any time during a work day.

Target organ: Organ in which the toxic injury manifests in terms of dysfunction or overt disease.

Threshold Limit Values (TLVs®): Recommended guidelines for occupational exposure to airborne contaminants, published by the American Conference of Governmental Industrial Hygienists (ACGIH®). TLVs refer to airborne concentrations of chemical substances and represent conditions under which it is believed that nearly all workers may be repeatedly exposed, day after day, over a working lifetime, without adverse effects. TLVs may be designated as ceiling limits, STELs, or 8-hr TWA limits.

Time-weighted average (TWA): A worker's 8-hour (or up to 10-hour) time-weighted average exposure concentration that shall not be exceeded during an 8-hour (or up to 10-hour) work shift of a 40-hour week. The average concentration is weighted to take into account the duration of different exposure concentrations.

Toxicity: The degree to which a substance is able to cause an adverse effect on an exposed organism.

Uncertainty factors (UFs): Mathematical adjustments applied to the POD when developing IDLH values. The UFs for IDLH value derivation are determined by considering the study and effect used for the POD, with further modification based on the overall database.

Workplace Environmental Exposure Levels (WEELs®): Exposure levels developed by the American Industrial Hygiene Association (AIHA®) that provide guidance for protecting most workers from adverse health effects related to occupational chemical exposures, expressed as TWA or ceiling limits.

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1 Introduction

1.1 Overview of the IDLH Value for n-Butyl Acrylate

IDLH value: 113 ppm

Basis for IDLH value: The IDLH value for n-butyl acrylate is based on the 30-minute RD_{50} (concentration estimated to result in a 50% depression in breathing rate) of 340 ppm in mice [Kirkpatrick 2003]. This effect is classified as potentially escape-impairing. Application of an uncertainty factor of 3 to account for extrapolation from a threshold for escape-impairing effects in animals, animal to human differences, and human variability results in an IDLH value of 113 ppm.

1.2 Purpose

This *IDLH Value Profile* presents (1) a brief summary of technical data associated with acute inhalation exposures to n-butyl acrylate and (2) the rationale behind the immediately dangerous to life or health (IDLH) value for n-butyl acrylate. IDLH values are developed on the basis of 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, and LC_{50} values). For n-butyl acrylate, the in-depth literature search was conducted through May 2016.

1.3 General Substance Information

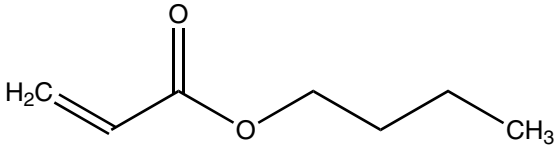
Chemical: n-Butyl Acrylate

CAS No: 141-32-2

Synonyms: Acrylic acid ester, n-butyl ester; Butyl acrylate; n-Butyl ester; n-Butyl propenoate; Butyl 2-propenoate; 2-Propenoic acid, butyl ester*

Chemical category: Carboxylic acid esters†

Structural formula:



References: *NLM [2016]; †IFA [2016]

Table 1 highlights selected physiochemical properties of n-butyl acrylate relevant to IDLH conditions. Table 2 provides alternative exposure guidelines for n-butyl acrylate. Table 3 summarizes the Acute Exposure Guidelines Level (AEGL) values for n-butyl acrylate.

Table 1: Physiochemical properties of n-butyl acrylate

Property	Value
Molecular weight	128.17[*]
Chemical formula	$C_7H_{12}O_2$
Description	Colorless liquid
Odor	Strong, fruity; Rancid; Plastic
Odor threshold	0.00096-0.10 ppm[§]
UEL	8%[†]
LEL	1.2%[†]
Vapor pressure	4.3 torr at 20°C (68°F)[*]
Flash point	36.5°C (97.7°F), closed cup[*]
Ignition temperature	292.78°C (559°F)[‡]
Solubility	Miscible with most common organic solvents[*]

References: [*]ACGIH [2015]; [§]NAS [2007]; [†]IFA [2016]; [‡]HSDB [2016]

Table 2: Alternative exposure guidelines for n-butyl acrylate

Organization	Value
Original (SCP) IDLH value[*]	None
NIOSH REL[†]	10 ppm (55 mg/m^3), TWA
OSHA PEL[§]	None
ACGIH TLV®[‡]	2 ppm, TWA
AIHA ERPGs™[¶]	ERPG-1: 0.05 ppm; ERPG-2: 25 ppm; ERPG-3: 250 ppm
AIHA WEELs®[¶]	Not available

References: [*]NIOSH [1994]; [§]OSHA [2016]; [†]NIOSH [2016]; [‡]ACGIH [2015]; [¶]AIHA [2014]

Table 3: AEGL values for n-butyl acrylate

Classification	10-min	30-min	1-hour	4-hour	8-hour	Endpoint [reference]
AEGL-1	8.3 ppm 44.0 mg/m^3	8.3 ppm 44.0 mg/m^3	8.3 ppm 44.0 mg/m^3	8.3 ppm 44.0 mg/m^3	8.3 ppm 44.0 mg/m^3	No clinical signs with repeated exposures [Rohm and Haas Co. 1992; Merkle and Klimisch 1983]
AEGL-2	160.0 ppm 850.0 mg/m^3	160.0 ppm 850.0 mg/m^3	130.0 ppm 690.0 mg/m^3	81.0 ppm 430.0 mg/m^3	53.0 ppm 280.0 mg/m^3	Clinical signs and histopathology with repeated exposure [Klimisch et al. 1978]
AEGL-3	820.0 ppm 4,400.0 mg/m^3	820.0 ppm 4,400.0 mg/m^3	480.0 ppm 2,600.0 mg/m^3	170.0 ppm 906.0 mg/m^3	97.0 ppm 520.0 mg/m^3	Calculated $BMCL_{05}$ from LC_{50} data [Oberly and Tansy 1985]

Reference: NAS [2007]

2 Animal Toxicity Data

The available data on n-butyl acrylate lethality and toxicity are limited. LC_{50} values were not substantially different among hamsters, rats, and mice. Engelhardt and Klimisch [1983] exposed Chinese hamsters and Sprague-Dawley rats to n-butyl acrylate for 6 hours for 3 days, followed by a single 5-hour exposure on day 4, at concentrations of 817 and 820 ppm, respectively. Clinical signs, dyspnea, disequilibrium, and bloody discharge from the eyes and noses were observed. No rats died, but four male hamsters died during the exposure period.

The most reliable animal lethality data came from the Oberly and Tansy [1985] 4-hour LC_{50} study in rats. Male Sprague-Dawley rats underwent whole-body exposure for 4 hours to n-butyl acrylate (1,990, 2,035, 2,500, 2,828, or 3,041 ppm), followed by a 14-day observation period. Vapor concentration was determined by gas chromatography. The animals had normal behavior during the first few minutes of the exposures but then exhibited labored breathing and irritation of the eyes, nose, and respiratory tract. All deaths occurred within 24 hours and were attributed to cardiopulmonary collapse. The number of deaths at each concentration was 0, 1, 3, 5, and 7, respectively. The 4-hour LC_{50} was calculated as 2,730 ppm. The NAS [2007] calculated a 4-hour lethality $BMCL_{05}$ value (a statistical lower confidence limit on the dose that produces a predetermined change in response rate of an adverse effect) by log-probit analysis, using U.S. EPA Benchmark Dose Software version 1.3.2. The resulting 4-hour $BMCL_{05}$ of 1,652 ppm was adjusted to a 30-minute exposure duration concentration of 8,179 ppm. In a 4-hour LC_{50} study in male and female Sprague-Dawley rats, BASF [1979b,c; 1980] noted severe irritation of the mucous membranes in the rats after they breathed n-butyl acrylate at concentrations above 677 ppm. Deaths occurred at concentrations above 1,278 ppm. Kirkpatrick [2003] measured respiratory depression in Swiss Webster mice. Test groups (n = 8) were exposed to n-butyl acrylate concentrations of 30, 100, 200, 350, 650, or 900 ppm. Kirkpatrick [2003] reported respiratory rates decreased by 5% to 60% in a concentration-response manner. The no observed effect level (NOEL) was 30 ppm, whereas the lowest effect level for respiratory depression was 100 ppm. Kirkpatrick reported that the estimated chemical concentration that caused a 50% decrease in the respiratory rate, or RD_{50} value, was 340 ppm. No deaths were reported to have occurred during this study.

Table 4 summarizes the LC data identified in animal studies and provides 30-minute-equivalent derived values for n-butyl acrylate. Table 5 provides nonlethal concentration data reported from animal studies with 30-minute-equivalent derived values. Information in this table includes species of test animals, toxicological metrics (i.e., LC, NOAEL, and LOAEL), adjusted 30-minute concentration, and the justification for the composite uncertainty factors applied to calculate the derived values.

Table 4: Lethal concentration data for n-butyl acrylate

Reference	Species	LC_{50} (ppm)	LC_{10} (ppm)	Time (min)	Adjusted 30-min concentration*	Composite uncertainty factor	30-min equivalent derived value (ppm)†	Final value (ppm)‡
BASF [1979a]	Hamster	1,201	—	240	5,946	30§	198.8	199
BASF [1979a]	Mouse	1,278	—	240	6,327	30§	210.9	211
Oberly and Tansy [1985]	Rat	2,730	—	240	13,516	30§	450.5	451
Oberly and Tansy [1985]	Rat	—	1,652	240	8,179	10¶	817.9	818
BASF [1979b,c; 1980]	Rat	1,936	—	240	9,585	30§	319.5	320

*NAS [2007] empirically estimated n = 1.3 by combining 1- and 4- hour LC_{50} data sets from ethyl acrylate in a 3-dimensional probit analysis. The n = 1.3 was used during all duration adjustments for n-butyl acrylate. Additional information on the calculation of duration-adjusted concentrations can be located in NIOSH [2013].

†The derived value is the result of the adjusted 30-minute LC value divided by the composite uncertainty factor.

‡Values rounded to the appropriate significant figure.

§Composite uncertainty factor to account for adjustment of LC_{50} values to LC_{01} values, use of lethal concentration threshold in animals, interspecies differences, and human variability.

¶Composite uncertainty factor to account for lethal concentration threshold in animals, interspecies differences, human variability, and extrapolation from a threshold for escape-impairing effects in animals.

Table 5: Nonlethal concentration data for n-butyl acrylate

Reference	Species	Critical nonlethal effect	RD_{50} (ppm)	Time (min)	Adjusted 30-min concentration*	Composite uncertainty factor	30-min equivalent derived value (ppm)†	Final value (ppm)‡
Kirkpatrick [2003]§	Mouse	Respiratory depression	340¶	30	340	3**	113.3	113

*NAS [2007] empirically estimated n = 1.3 by combining 1- and 4- hour LC50 data sets from ethyl acrylate in a 3-dimensional probit analysis. The n = 1.3 was used during all duration adjustments for n-butyl acrylate. Additional information on the calculation of duration-adjusted concentrations can be located in NIOSH [2013].

†The derived value is the result of the adjusted 30-min value divided by the composite uncertainty factor. The composite uncertainty factor used varies for each study on the basis of the nature and severity of the endpoint observed.

‡Values rounded to the appropriate significant figure.

§Identified study is the primary basis of the IDLH value for n-butyl acrylate.

¶This concentration was determined to be an RD_{50} (concentration estimated to result in a 50% depression in breathing rate).

**Composite uncertainty factor to account for extrapolation from a threshold for escape-impairing effects in animals, animal to human differences, and human variability.

3 Human Data

No reports of human fatalities from exposure to n-butyl acrylate were found. Very little information is available concerning human exposure to n-butyl acrylate. Symptoms of irritation in chemical plant workers were occasionally reported [Tucek et al. 2002]. Dermal sensitization has been reported [BIBRA 1991], but not respiratory sensitization.

4 Summary

In the absence of adequate human data, the IDLH value is based on animal data. Although lethality data are available (see Table 4), the preferred data that serve as the basis of the IDLH value indicate respiratory depression, which is categorized as an escape-impairing effect. Kirkpatrick [2003] reported a 30-minute RD_{50} value in mice exposed to 340 ppm. This effect is classified as potentially escape-impairing. No duration adjustment was needed. Application of a composite uncertainty factor of 3, to account for extrapolation from a threshold for escape-impairing effects in animals, animal to human differences, and human variability, yielded an IDLH value of 113 ppm.

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