



## Short review

## RIFM fragrance ingredient safety assessment, 2-methyl-3-buten-2-ol, CAS Registry Number 115-18-4



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## ABSTRACT

The use of this material under current use conditions is supported by the existing information.

This material was evaluated for genotoxicity, repeated dose toxicity, developmental toxicity, reproductive toxicity, local respiratory toxicity, phototoxicity, skin sensitization potential as well as environmental safety. Repeated dose, developmental, and reproductive toxicities were determined to have the most conservative systemic exposure derived NO[A]EL of 50 mg/kg/day, based on OECD gavage toxicity studies in rats, that resulted in a MOE of 4545455 after considering 100% absorption from skin contact and inhalation. A MOE of >100 is deemed acceptable.

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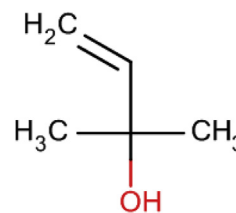
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**Version: 050515. This version replaces any previous versions.**

**Name:** 2-Methyl-3-buten-2-ol

**CAS Registry Number:** 115-18-4



**Abbreviation list:** **2-Box Model** – a RIFM, Inc. proprietary *in silico* tool used to calculate fragrance air exposure concentration

**97.5th percentile**- The concentration of the fragrance ingredient is obtained from examination of several thousand commercial fine fragrance formulations. The upper 97.5th percentile concentration is calculated from these data and is then used to estimate the dermal systemic exposure in ten types of the most frequently used personal care and cosmetic products. The dermal route is the major route in assessing the safety of fragrance ingredients. Further explanation of how the data were obtained and of how exposures were determined has been previously reported by [Cadby et al. \(2002\)](#) and [Ford et al. \(2000\)](#).

**AF**-Assessment Factor

**BCF**-Bioconcentration factor

**DEREK**-Derek nexus is an *in silico* tool used to identify structural alerts

**DST**-Dermal Sensitization Threshold

**ECHA**-European Chemicals Agency

**EU**- Europe/European Union

**GLP**-Good Laboratory Practice

**IFRA**-The International Fragrance Association

**LOEL**-Lowest Observable Effect Level

**MOE**-Margin of Exposure

**MPPD**-Multiple-Path Particle Dosimetry. An *in silico* model for inhaled vapors used to simulate fragrance lung deposition

**NA**-North America

**NESIL**-No Expected Sensitization Induction Level

**NOAEC**-No Observed Adverse Effect Concentration

**NOAEL**-No Observed Adverse Effect Level

**NOEC**-No Observed Effect Concentration

**OECD**-Organisation for Economic Co-operation and Development

**OECD TG**-Organisation for Economic Co-operation and Development Testing Guidelines

**PBT**-Persistent, Bioaccumulative, and Toxic

**PEC/PNEC**- Predicted Environmental Concentration/Predicted No Effect Concentration

**QRA**-quantitative risk assessment

**REACH**-Registration, Evaluation, Authorisation, and Restriction of Chemicals

**RIFM**-Research Institute for Fragrance Materials

**RQ**-Risk Quotient

**TTC**-Threshold of Toxicological Concern

**UV/Vis Spectra**-Ultra Violet/Visible spectra

**VCF**-Volatile Compounds in Food

**VoU**-Volume of Use

**vPvB**-(very) Persistent, (very) Bioaccumulative

**WOE**-Weight of Evidence

**RIFM's Expert Panel\*** concludes that this material is safe under the limits described in this safety assessment.

This safety assessment is based on RIFM's Criteria Document ([Api et al., 2015](#)) and should be referred to for clarifications.

Each endpoint discussed in this safety assessment reviews the relevant data that were available at the time of writing (version number in the top box is indicative of the date of approval based on a two digit month/day/year), both in the RIFM database (consisting of publicly available and proprietary data) and through publicly available information sources (i.e., SciFinder and PubMed). Studies selected for this safety assessment were based on appropriate test criteria such as, acceptable guidelines, sample size, study duration, route of exposure, relevant animal species, most relevant testing endpoints, etc. A key study for each endpoint was selected based on the most conservative end-point value (e.g., PNEC, NOAEL, LOEL, and NESIL).

\*RIFM's Expert Panel is an independent body that selects its own members and establishes its own operating procedures. The Expert Panel is comprised of internationally known scientists that provide RIFM guidance relevant to human health and environmental protection.

Summary: The use of this material under current use conditions is supported by the existing information.

This material was evaluated for genotoxicity, repeated dose toxicity, developmental toxicity, reproductive toxicity, local respiratory toxicity, phototoxicity, skin sensitization potential as well as environmental safety. Repeated dose, developmental, and reproductive toxicities were determined to have the most conservative systemic exposure derived NO[A]EL of 50 mg/kg/day, based on OECD gavage toxicity studies in rats, that resulted in a MOE of 4545455 after considering 100% absorption from skin contact and inhalation. A MOE of >100 is deemed acceptable.

#### Human Health Safety Assessment

**Genotoxicity:** Not Genotoxic

**Repeated Dose Toxicity:** NOAEL = 50 mg/kg/day

**Developmental and Reproductive Toxicity:** NOAEL = 50 mg/kg/day

**Skin Sensitization:** Not sensitizing

**Phototoxicity/Photoallergenicity:** Not phototoxic/photoallergenic

**Local Respiratory Toxicity:** No NOAEC available. Exposure is below TTC.

#### Environmental Safety Assessment

##### **Hazard Assessment:**

**Persistence:** Critical Measured Value: 67%

(RIFM, 1989a; RIFM, 1992a)  
(OECD SIDS, 1995: 3-Buten-2-ol, 2-methyl-)  
(OECD SIDS, 1995: 3-Buten-2-ol, 2-methyl-)  
(RIFM, 1991)  
(UV spectra, RIFM DB)

(REACH dossier accessed 14 May 2013; see Section 9)

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