



## Short Review

## RIFM fragrance ingredient safety assessment, 2-(3-phenylpropyl) tetrahydrofuran, CAS Registry Number 3208-40-0



A.M. Api<sup>a</sup>, D. Belsito<sup>b</sup>, S. Biserta<sup>a</sup>, D. Botelho<sup>a</sup>, M. Bruze<sup>c</sup>, G.A. Burton Jr.<sup>d</sup>, J. Buschmann<sup>e</sup>, M. A. Cancellieri<sup>a</sup>, M.L. Dagli<sup>f</sup>, M. Date<sup>a</sup>, W. Dekant<sup>g</sup>, C. Deodhar<sup>a</sup>, A.D. Fryer<sup>h</sup>, S. Gadhia<sup>a</sup>, L. Jones<sup>a</sup>, K. Joshi<sup>a</sup>, M. Kumar<sup>a</sup>, A. Lapczynski<sup>a</sup>, M. Lavelle<sup>a</sup>, I. Lee<sup>a</sup>, D.C. Liebler<sup>i</sup>, H. Moustakas<sup>a</sup>, M. Na<sup>a</sup>, T.M. Penning<sup>j</sup>, G. Ritacco<sup>a</sup>, J. Romine<sup>a</sup>, N. Sadekar<sup>a</sup>, T.W. Schultz<sup>k</sup>, D. Selechnik<sup>a</sup>, F. Siddiqi<sup>a</sup>, I.G. Sipes<sup>l</sup>, G. Sullivan<sup>a,\*</sup>, Y. Thakkar<sup>a</sup>, Y. Tokura<sup>m</sup>

<sup>a</sup> Research Institute for Fragrance Materials, Inc., 50 Tice Boulevard, Woodcliff Lake, NJ, 07677, USA

<sup>b</sup> Columbia University Medical Center, Department of Dermatology, 161 Fort Washington Ave., New York, NY, 10032, USA

<sup>c</sup> Malmo University Hospital, Department of Occupational & Environmental Dermatology, Sodra Forstadsgatan 101, Entrance 47, Malmo, SE, 20502, Sweden

<sup>d</sup> School of Natural Resources & Environment, University of Michigan, Dana Building G110, 440 Church St., Ann Arbor, MI, 48109, USA

<sup>e</sup> Fraunhofer Institute for Toxicology and Experimental Medicine, Nikolai-Fuchs-Strasse 1, 30625, Hannover, Germany

<sup>f</sup> University of Sao Paulo, School of Veterinary Medicine and Animal Science, Department of Pathology, Av. Prof. dr. Orlando Marques de Paiva, 87, Sao Paulo, CEP 05508-900, Brazil

<sup>g</sup> University of Wuerzburg, Department of Toxicology, Versbacher Str. 9, 97078, Würzburg, Germany

<sup>h</sup> Oregon Health Science University, 3181 SW Sam Jackson Park Rd., Portland, OR, 97239, USA

<sup>i</sup> Vanderbilt University School of Medicine, Department of Biochemistry, Center in Molecular Toxicology, 638 Robinson Research Building, 2200 Pierce Avenue, Nashville, TN, 37232-0146, USA

<sup>j</sup> University of Pennsylvania, Perelman School of Medicine, Center of Excellence in Environmental Toxicology, 1316 Biomedical Research Building (BRB) II/III, 421 Curie Boulevard, Philadelphia, PA, 19104-3083, USA

<sup>k</sup> The University of Tennessee, College of Veterinary Medicine, Department of Comparative Medicine, 2407 River Dr., Knoxville, TN, 37996-4500, USA

<sup>l</sup> Department of Pharmacology, University of Arizona, College of Medicine, 1501 North Campbell Avenue, P.O. Box 245050, Tucson, AZ, 85724-5050, USA

<sup>m</sup> The Journal of Dermatological Science (JDS), Department of Dermatology, Hamamatsu University School of Medicine, 1-20-1 Handayama, Higashi-ku, Hamamatsu, 431-3192, Japan

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\* Corresponding author.

E-mail address: [gsullivan@rifm.org](mailto:gsullivan@rifm.org) (G. Sullivan).

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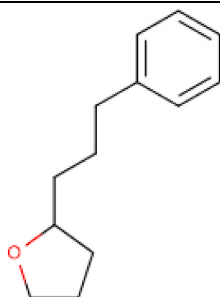
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Name: 2-(3-Phenylpropyl)tetrahydrofuran CAS  
Registry Number: 3208-40-0



#### Abbreviation/Definition List:

**2-Box Model** - A RIFM, Inc. proprietary *in silico* tool used to calculate fragrance air exposure concentration

**AF** - Assessment Factor

**BCF** - Bioconcentration Factor

**Creme RIFM Model** - The Creme RIFM Model uses probabilistic (Monte Carlo) simulations to allow full distributions of data sets, providing a more realistic estimate of aggregate exposure to individuals across a population (Comiskey et al., 2015, 2017; Safford et al., 2015a, 2017) compared to a deterministic aggregate approach

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

**DRF** - Dose Range Finding

**DST** - Dermal Sensitization Threshold

**ECHA** - European Chemicals Agency

**ECOSAR** - Ecological Structure-Activity Relationships Predictive Model

**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

**NOEL** - No Observed Effect Level

**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

**Perfumery** - In this safety assessment, perfumery refers to fragrances made by a perfumer used in consumer products only. The exposures reported in the safety assessment include consumer product use, but do not include occupational exposures.

**QRA** - Quantitative Risk Assessment

**QSAR** - Quantitative Structure-Activity Relationship

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

**RfD** - Reference Dose

**RIFM** - Research Institute for Fragrance Materials

**RQ** - Risk Quotient

**Statistically Significant** - Statistically significant difference in reported results as compared to controls with a  $p < 0.05$  using appropriate statistical test

**TTC** - Threshold of Toxicological Concern

**UV/Vis spectra** - Ultraviolet/Visible spectra

**VCF** - Volatile Compounds in Food

**VoU** - Volume of Use

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

**WoE** - Weight of Evidence

**The Expert Panel for Fragrance Safety\* concludes that this material is safe as described in this safety assessment.**

This safety assessment is based on the RIFM Criteria Document (Api et al., 2015), which should be referred to for clarifications.

Each endpoint discussed in this safety assessment includes 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 2-digit month/day/year), both in the RIFM Database (consisting of publicly available and proprietary data) and through publicly available information sources (e.g., 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

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based on the most conservative endpoint value (e.g., PNEC, NOAEL, LOEL, and NESIL).

\*The Expert Panel for Fragrance Safety 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 with guidance relevant to human health and environmental protection.

**Summary: The existing information supports the use of this material as described in this safety assessment.**

2-(3-Phenylpropyl)tetrahydrofuran was evaluated for genotoxicity, repeated dose toxicity, reproductive toxicity, local respiratory toxicity, phototoxicity/photoallergenicity, skin sensitization, and environmental safety. Data show that 2-(3-phenylpropyl)tetrahydrofuran is not genotoxic. The repeated dose, reproductive, and local respiratory toxicity endpoints were evaluated using the threshold of toxicological concern (TTC) for a Cramer Class III material, and the exposure to 2-(3-phenylpropyl)tetrahydrofuran is below the TTC (0.0015 mg/kg/day, 0.0015 mg/kg/day, and 0.47 mg/day, respectively). The skin sensitization endpoint was completed using the dermal sensitization threshold (DST) for reactive materials (64  $\mu\text{g}/\text{cm}^2$ ); exposure is below the DST. The phototoxicity/photoallergenicity endpoints were evaluated based on ultraviolet (UV) spectra; 2-(3-phenylpropyl)tetrahydrofuran is not expected to be phototoxic/photoallergenic. The environmental endpoints were evaluated; 2-(3-phenylpropyl)tetrahydrofuran was found not to be persistent, bioaccumulative, and toxic (PBT) as per the International Fragrance Association (IFRA) Environmental Standards, and its risk quotients, based on its current volume of use in Europe and North America (i.e., Predicted Environmental Concentration/Predicted No Effect Concentration [PEC/PNEC]), are  $< 1$ .

#### Human Health Safety Assessment

**Genotoxicity:** Not genotoxic. (RIFM, 2019; RIFM, 2020)

**Repeated Dose Toxicity:** No NOAEL available. Exposure is below the TTC.

**Reproductive Toxicity:** No NOAEL available. Exposure is below the TTC.

**Skin Sensitization:** No safety concerns at current, declared use levels; exposure is below the DST.

**Phototoxicity/Photoallergenicity:** Not expected to be phototoxic/photoallergenic. (UV Spectra; RIFM Database)

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

#### Environmental Safety Assessment

##### Hazard Assessment:

**Persistence:** Screening-level: 2.72 (BIOWIN 3) (EPI Suite v4.11; US EPA, 2012a)

**Bioaccumulation:** Screening-level: 216.5 L/kg (EPI Suite v4.11; US EPA, 2012a)

**Ecotoxicity:** Screening-level: Fish LC50: 4.31 mg/L (RIFM Framework; Salvitto et al., 2002)

**Conclusion:** Not PBT or vPvB as per IFRA Environmental Standards

##### Risk Assessment:

**Screening-level:** PEC/PNEC (North America and Europe)  $< 1$  (RIFM Framework; Salvitto et al., 2002)

**Critical Ecotoxicity Endpoint:** Fish LC50: 4.31 mg/L (RIFM Framework; Salvitto et al., 2002)

**RIFM PNEC is:** 0.00431  $\mu\text{g}/\text{L}$

• **Revised PEC/PNECs (2015 IFRA VoU):** North America and Europe: Not applicable; cleared at screening-level

## 1. Identification

- Chemical Name:** 2-(3-Phenylpropyl)tetrahydrofuran
- CAS Registry Number:** 3208-40-0
- Synonyms:** Furan, tetrahydro-2-(3-phenylpropyl)-;  $\alpha$ -(3-Phenylpropyl)tetrahydrofuran; 2-(3-Phenylpropyl)tetrahydrofuran
- Molecular Formula:**  $\text{C}_{13}\text{H}_{18}\text{O}$
- Molecular Weight:** 190.28
- RIFM Number:** 6201
- Stereochemistry:** No isomer specified. One stereocenter and 2 total stereoisomers possible.

## 2. Physical data

- Boiling Point:** 275.08  $^{\circ}\text{C}$  (EPI Suite)
- Flash Point:** Not Available
- Log  $K_{ow}$ :** 4.04 (EPI Suite)

4. **Melting Point:** 36.98 °C (EPI Suite)
5. **Water Solubility:** 17.23 mg/L (EPI Suite)
6. **Specific Gravity:** Not Available
7. **Vapor Pressure:** 0.00316 mm Hg at 20 °C (EPI Suite v4.0), 0.03 mm Hg at 20 °C (Fragrance Materials Association), 0.00566 mm Hg at 25 °C (EPI Suite)
8. **UV Spectra:** No significant absorbance between 290 and 700 nm; molar absorption coefficient is below the benchmark (1000 L mol<sup>-1</sup> · cm<sup>-1</sup>)
9. **Appearance/Organoleptic:** Not Available

### 3. Volume of use (worldwide band)

1. <0.1 metric ton per year (IFRA, 2015)

### 4. Exposure to fragrance ingredient (Creme RIFM Aggregate Exposure Model v1.0)

1. **95th Percentile Concentration in Hydroalcoholics:** 0.00036% (RIFM, 2017)
2. **Inhalation Exposure\*:** 0.0000004 mg/kg/day or 0.000029 mg/day (RIFM, 2017)
3. **Total Systemic Exposure\*\*:** 0.000022 mg/kg/day (RIFM, 2017)

\*95th percentile calculated exposure derived from concentration survey data in the Creme RIFM Aggregate Exposure Model (Comiskey, 2015, 2017; Safford, 2015a, 2017).

\*\*95th percentile calculated exposure; assumes 100% absorption unless modified by dermal absorption data as reported in Section V. It is derived from concentration survey data in the Creme RIFM Aggregate Exposure Model and includes exposure via dermal, oral, and inhalation routes whenever the fragrance ingredient is used in products that include these routes of exposure (Comiskey, 2015, 2017; Safford, 2015a, 2017).

### 5. Derivation of systemic absorption

1. **Dermal:** Assumed 100%
2. **Oral:** Assumed 100%
3. **Inhalation:** Assumed 100%

### 6. Computational toxicology evaluation

#### 1. Cramer Classification: Class III, High

Expert Judgment	Toxtree v 2.6	OECD QSAR Toolbox v 3.2
III	III	III

2. **Analogs Selected:**
  - a. **Genotoxicity:** None
  - b. **Repeated Dose Toxicity:** None
  - c. **Reproductive Toxicity:** None
  - d. **Skin Sensitization:** None
  - e. **Phototoxicity/Photoallergenicity:** None
  - f. **Local Respiratory Toxicity:** None
  - g. **Environmental Toxicity:** None
3. **Read-across Justification:** None

### 7. Metabolism

No relevant data available for inclusion in this safety assessment.

**Additional References:** None.

### 8. Natural occurrence (discrete chemical) or composition (NCS)

2-(3-Phenylpropyl)tetrahydrofuran is not reported to occur in foods by the VCF\*.

\*VCF (Volatile Compounds in Food): Database/Nijssen, L.M.; Ingen-Visscher, C.A. van; Donders, J.J.H. (eds). – Version 15.1 – Zeist (The Netherlands): TNO Triskelion, 1963–2014. A continually updated database containing information on published volatile compounds that have been found in natural (processed) food products. Includes FEMA GRAS and EU-Flavis data.

### 9. REACH dossier

Pre-registered for 2010; no dossier available as of 09/06/19.

### 10. Conclusion

The existing information supports the use of this material as described in this safety assessment.

### 11. Summary

#### 11.1. Human health endpoint summaries

##### 11.1.1. Genotoxicity

Based on the current existing data, 2-(3-phenylpropyl)tetrahydrofuran does not present a concern for genotoxicity.

**11.1.1.1. Risk assessment.** The mutagenic activity of 2-(3-phenylpropyl)tetrahydrofuran has been evaluated in a bacterial reverse mutation assay conducted in compliance with GLP regulations and in accordance with OECD TG 471 using the standard plate incorporation method. *Salmonella typhimurium* strains TA98, TA100, TA1535, TA1537, and *Escherichia coli* strain WP2uvrA were treated with 2-(3-phenylpropyl)tetrahydrofuran in dimethyl sulfoxide (DMSO) at concentrations up to 5000 µg/plate. No increases in the mean number of revertant colonies were observed at any tested concentration in the presence or absence of S9 (RIFM, 2019). Under the conditions of the study, 2-(3-phenylpropyl)tetrahydrofuran was not mutagenic in the Ames test.

The clastogenic activity of 2-(3-phenylpropyl)tetrahydrofuran was evaluated in an *in vitro* micronucleus test conducted in compliance with GLP regulations and in accordance with OECD TG 487. Human peripheral blood lymphocytes were treated with 2-(3-phenylpropyl)tetrahydrofuran in dimethyl sulfoxide (DMSO) at concentrations up to 10,000 µM in the dose range finding (DRF) study; micronuclei analysis was conducted at concentrations up to 1317 µM in the presence and absence of metabolic activation. 2-(3-Phenylpropyl)tetrahydrofuran did not induce binucleated cells with micronuclei when tested up to cytotoxic levels concentration in either the presence or absence of an S9 activation system (RIFM, 2020). Under the conditions of the study, 2-(3-phenylpropyl)tetrahydrofuran was considered to be non-clastogenic in the *in vitro* micronucleus test.

Based on the data available, 2-(3-phenylpropyl)tetrahydrofuran does not present a concern for genotoxic potential.

**Additional References:** None.

**Literature Search and Risk Assessment Completed On:** 10/24/19.

##### 11.1.2. Repeated dose toxicity

There are insufficient repeated dose toxicity data on 2-(3-phenylpropyl)tetrahydrofuran or any read-across materials. The total systemic exposure to 2-(3-phenylpropyl)tetrahydrofuran is below the TTC for the repeated dose toxicity endpoint of a Cramer Class III material at the current level of use.

**11.1.2.1. Risk assessment.** There are no repeated dose toxicity data on 2-(3-phenylpropyl)tetrahydrofuran or any read-across materials that can be used to support the repeated dose toxicity endpoint. The total systemic exposure (0.022 µg/kg/day) is below the TTC for 2-(3-phenylpropyl)tetrahydrofuran (1.5 µg/kg/day; Kroes, 2007).

**Additional References:** None.

**Literature Search and Risk Assessment Completed On:** 10/10/19.

### 11.1.3. Reproductive toxicity

There are insufficient reproductive toxicity data on 2-(3-phenylpropyl)tetrahydrofuran or any read-across materials. The total systemic exposure to 2-(3-phenylpropyl)tetrahydrofuran is below the TTC for the reproductive toxicity endpoint of a Cramer Class III material at the current level of use.

**11.1.3.1. Risk assessment.** There are no reproductive toxicity data on 2-(3-phenylpropyl)tetrahydrofuran or on any read-across materials that can be used to support the reproductive toxicity endpoint. The total systemic exposure to 2-(3-phenylpropyl)tetrahydrofuran (0.022 µg/kg/day) is below the TTC (1.5 µg/kg/day; Kroes, 2007; Laufersweiler, 2012) for the reproductive toxicity endpoint of a Cramer Class III material at the current level of use.

**Additional References:** None.

**Literature Search and Risk Assessment Completed On:** 10/09/19.

### 11.1.4. Skin sensitization

Based on existing data and the application of DST, 2-(3-phenylpropyl)tetrahydrofuran does not present a safety concern for skin sensitization under the current, declared levels of use.

**11.1.4.1. Risk assessment.** The chemical structure of this material indicates that it would not be expected to react with skin proteins directly, but its metabolite will be reactive (Roberts, 2007; Toxtree v3.1.0; OECD Toolbox v4.2). No predictive skin sensitization studies are available for 2-(3-phenylpropyl)tetrahydrofuran. Acting conservatively due to the lack of data, the reported exposure was benchmarked utilizing the reactive DST of 64 µg/cm<sup>2</sup> (Safford, 2008, 2011, 2015b; Roberts, 2015). The current exposure from the 95th percentile concentration is below the DST for reactive materials when evaluated in all QRA categories. Table 1 provides the maximum acceptable concentrations for 2-(3-phenylpropyl)tetrahydrofuran that present no appreciable risk for skin sensitization based on the reactive DST. These levels represent maximum acceptable concentrations based on the DST approach.

**Table 1**

Maximum acceptable concentrations for 2-(3-phenylpropyl)tetrahydrofuran that present no appreciable risk for skin sensitization based on reactive DST.

IFRA Category <sup>a</sup>	Description of Product Type	Maximum Acceptable Concentrations in Finished Products Based on Reactive DST	Reported 95th Percentile Use Concentrations in Finished Products
1	Products applied to the lips	0.0049%	6.1 × 10 <sup>-6</sup>
2	Products applied to the axillae	0.0015%	5.5 × 10 <sup>-4</sup>
3	Products applied to the face using fingertips	0.029%	5.5 × 10 <sup>-5</sup>
4	Fine fragrance products	0.027%	3.6 × 10 <sup>-4</sup>
5	Products applied to the face and body using the hands (palms), primarily leave-on	0.0070%	5.5 × 10 <sup>-4</sup>
6	Products with oral and lip exposure	0.016%	NRU <sup>2</sup>
7	Products applied to the hair with some hand contact	0.056%	1.1 × 10 <sup>-5</sup>
8	Products with significant ano-genital exposure	0.0029%	No Data <sup>c</sup>
9	Products with body and hand exposure, primarily rinse-off	0.054%	2.9 × 10 <sup>-5</sup>
10	Household care products with mostly hand contact	0.19%	3.9 × 10 <sup>-5</sup>
11	Products with intended skin contact but minimal transfer of fragrance to skin from inert substrate	0.11%	No Data <sup>c</sup>
12	Products not intended for direct skin contact, minimal or insignificant transfer to skin	No Restriction	9.4 × 10 <sup>-4</sup>

Note: <sup>a</sup>For a description of the categories, refer to the IFRA/RIFM Information Booklet. <sup>b</sup>No reported use. <sup>c</sup>Fragrance exposure from these products is very low. These products are not currently in the Creme RIFM Aggregate Exposure Model.

However, additional studies may show it could be used at higher levels.

**Additional References:** None.

**Literature Search and Risk Assessment Completed On:** 10/20/19.

### 11.1.5. Phototoxicity/photoallergenicity

Based on the available UV/Vis spectra, 2-(3-phenylpropyl)tetrahydrofuran would not be expected to present a concern for phototoxicity or photoallergenicity.

**11.1.5.1. Risk assessment.** There are no phototoxicity studies available for 2-(3-phenylpropyl)tetrahydrofuran in experimental models. UV/Vis absorption spectra indicate no significant absorption between 290 and 700 nm. The corresponding molar absorption coefficient is well below the benchmark of concern for phototoxicity and photoallergenicity (Henry, 2009). Based on the lack of absorbance, 2-(3-phenylpropyl)tetrahydrofuran does not present a concern for phototoxicity or photoallergenicity.

**11.1.5.2. UV spectra analysis.** UV/Vis absorption spectra (OECD TG 101) were obtained. The spectra indicate no significant absorbance in the range of 290–700 nm. The molar absorption coefficient is below the benchmark of concern for phototoxic effects, 1000 L mol<sup>-1</sup> · cm<sup>-1</sup> (Henry, 2009).

**Additional References:** None.

**Literature Search and Risk Assessment Completed On:** 09/30/19.

### 11.1.6. Local Respiratory Toxicity

The margin of exposure could not be calculated due to a lack of appropriate data. The exposure level for 2-(3-phenylpropyl)tetrahydrofuran is below the Cramer Class III TTC value for inhalation exposure local effects.

**11.1.6.1. Risk assessment.** There are no inhalation data available on 2-(3-phenylpropyl)tetrahydrofuran. Based on the Creme RIFM Model, the inhalation exposure is 0.000029 mg/day. This exposure is 16,207 times lower than the Cramer Class III TTC value of 0.47 mg/day (based on human lung weight of 650 g; Carthew, 2009); therefore, the exposure at the current level of use is deemed safe.

**Additional References:** None.

**Literature Search and Risk Assessment Completed On:** 10/08/19.

## 11.2. Environmental endpoint summary

### 11.2.1. Screening-level assessment

A screening-level risk assessment of 2-(3-phenylpropyl)tetrahydrofuran was performed following the RIFM Environmental Framework (Salvito, 2002), which provides 3 tiered levels of screening for aquatic risk. In Tier 1, only the material's regional VoU, its log K<sub>ow</sub>, and its molecular weight are needed to estimate a conservative risk quotient (RQ), expressed as the ratio Predicted Environmental Concentration/Predicted No Effect Concentration (PEC/PNEC). A general QSAR with a high uncertainty factor applied is used to predict fish toxicity, as discussed in Salvito et al. (2002). In Tier 2, the RQ is refined by applying a lower uncertainty factor to the PNEC using the ECOSAR model (US EPA, 2012b), which provides chemical class-specific ecotoxicity estimates. Finally, if necessary, Tier 3 is conducted using measured biodeg-

	LC50 (Fish) (mg/L)	EC50 ( <i>Daphnia</i> ) (mg/L)	EC50 (Algae) (mg/L)	AF	PNEC (µg/L)	Chemical Class
RIFM Framework Screening-level (Tier 1)	<u>4.31</u>			1000000	0.00431	

radation and ecotoxicity data to refine the RQ, thus allowing for lower PNEC uncertainty factors. The data for calculating the PEC and PNEC for this safety assessment are provided in the table below. For the PEC, the range from the most recent IFRA Volume of Use Survey is reviewed. The PEC is then calculated using the actual regional tonnage, not the extremes of the range. Following the RIFM Environmental Framework, 2-(3-phenylpropyl)tetrahydrofuran was identified as a fragrance material with no potential to present a possible risk to the aquatic environment (i.e., its screening-level PEC/PNEC <1).

A screening-level hazard assessment using EPI Suite v4.11 (US EPA, 2012a) identified 2-(3-phenylpropyl)tetrahydrofuran as possibly persistent but not bioaccumulative based on its structure and physical-chemical properties. This screening-level hazard assessment considers the potential for a material to be persistent and bioaccumulative and toxic, or very persistent and very bioaccumulative as defined in the Criteria Document (Api et al., 2015). As noted in the Criteria Document, the screening criteria applied are the same as those used in the EU for REACH (ECHA, 2012). For persistence, if the EPI Suite model BIOWIN 3 predicts a value < 2.2 and either BIOWIN 2 or BIOWIN 6 predicts a value < 0.5, then the material is considered potentially persistent. A material would be considered potentially bioaccumulative if the EPI Suite model BCFBAF predicts a fish BCF ≥2000 L/kg. Ecotoxicity is determined in the above screening-level risk assessment. If, based on these model outputs (Step 1), additional assessment is required, a WoE-based review is then performed (Step 2). This review considers available data on the material's physical-chemical properties, environmental fate (e.g., OECD Guideline biodegradation studies or die-away studies), fish bioaccumulation, and higher-tier model outputs (e.g., US EPA's BIOWIN and BCFBAF found in EPI Suite v4.11).

### 11.2.2. Risk assessment

Based on the current Volume of Use (2015), 2-(3-phenylpropyl)tetrahydrofuran presents no risk to the aquatic compartment in the screening-level assessment.

### 11.2.3. Key studies

11.2.3.1. Biodegradation. No data available.

11.2.3.2. Ecotoxicity. No data available.

11.2.3.3. Other available data. 2-(3-Phenylpropyl)tetrahydrofuran has been pre-registered for REACH with no additional data available at this time.

Ecotoxicological data and PNEC derivation (all endpoints reported in mg/L; PNECs in µg/L).

Endpoints used to calculate PNEC are underlined.

Exposure information and PEC calculation (following RIFM Framework: Salvito, 2002).

Exposure	Europe (EU)	North America (NA)
Log K <sub>ow</sub> Used	4.04	4.04
Biodegradation Factor Used	0	0
Dilution Factor	3	3
Regional Volume of Use Tonnage Band	<1	<1
<b>Risk Characterization: PEC/PNEC</b>	<b>&lt;1</b>	<b>&lt;1</b>

Based on available data, the RQ for this material is < 1. No additional assessment is necessary.

The RIFM PNEC is 0.00431 µg/L. The revised PEC/PNECs for EU and NA are not applicable. The material was cleared at the screening-level; therefore, it does not present a risk to the aquatic environment at the current reported volumes of use.

**Literature Search and Risk Assessment Completed On:** 09/25/19.

## 12. Literature Search\*

- **RIFM Database:** Target, Fragrance Structure-Activity Group materials, other references, JECFA, CIR, SIDS
- **ECHA:** <https://echa.europa.eu/>
- **NTP:** <https://ntp.niehs.nih.gov/>
- **OECD Toolbox:** <https://www.oecd.org/chemicalsafety/risk-assessment/oecd-qsar-toolbox.htm>
- **SciFinder:** <https://scifinder.cas.org/scifinder/view/scifinder/scifinderExplore.jsf>
- **PubMed:** <https://www.ncbi.nlm.nih.gov/pubmed>
- **National Library of Medicine's Toxicology Information Services:** <https://toxnet.nlm.nih.gov/>
- **IARC:** <https://monographs.iarc.fr>
- **OECD SIDS:** <https://hpvchemicals.oecd.org/ui/Default.aspx>
- **EPA ACToR:** <https://actor.epa.gov/actor/home.xhtml>

- **US EPA HPVVIS:** [https://ofmpub.epa.gov/opthpv/public\\_search\\_publicdetails?submission\\_id=24959241&ShowComments=Yes&sqlstr=null&recordcount=0&User\\_title=DetailQuery%20Results&EndPointRpt=Y#submission](https://ofmpub.epa.gov/opthpv/public_search_publicdetails?submission_id=24959241&ShowComments=Yes&sqlstr=null&recordcount=0&User_title=DetailQuery%20Results&EndPointRpt=Y#submission)
- **Japanese NITE:** [https://www.nite.go.jp/en/chem/chrip/chrip\\_search/systemTop](https://www.nite.go.jp/en/chem/chrip/chrip_search/systemTop)
- **Japan Existing Chemical Data Base (JECDB):** [http://dra4.nihs.go.jp/mhlw\\_data/jsp/SearchPageENG.jsp](http://dra4.nihs.go.jp/mhlw_data/jsp/SearchPageENG.jsp)
- **Google:** <https://www.google.com>
- **ChemIDplus:** <https://chem.nlm.nih.gov/chemidplus/>

Search keywords: CAS number and/or material names.

\*Information sources outside of RIFM's database are noted as appropriate in the safety assessment. This is not an exhaustive list. The links listed above were active as of 01/31/20.

#### Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper. We wish to confirm that there are no known conflicts of interest associated with this publication and there has been no significant financial support for this work that could have influenced its outcome. RIFM staff are employees of the Research Institute for Fragrance Materials, Inc. (RIFM). The Expert Panel receives a small honorarium for time spent reviewing the subject work.

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