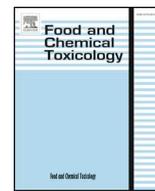




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Short Review

RIFM fragrance ingredient safety assessment, pentyl phenylacetate, CAS Registry Number 5137-52-0



A.M. Api^a, D. Belsito^b, D. Botelho^a, M. Bruze^c, G.A. Burton Jr^d, J. Buschmann^e, M.L. Dagli^f, M. Date^a, W. Dekant^g, C. Deodhar^a, M. Francis^a, A.D. Fryer^h, L. Jones^a, K. Joshi^a, S. La Cava^a, A. Lapczynski^a, D.C. Lieblerⁱ, D. O'Brien^a, A. Patel^a, T.M. Penning^j, G. Ritacco^a, J. Romine^a, N. Sadekar^a, D. Salvito^a, T.W. Schultz^k, I.G. Sipes^l, G. Sullivan^{a,*}, Y. Thakkar^a, Y. Tokura^m, S. Tsang^a

^a Research Institute for Fragrance Materials, Inc., 50 Tice Boulevard, Woodcliff Lake, NJ, 07677, USA

^b Member RIFM Expert Panel, Columbia University Medical Center, Department of Dermatology, 161 Fort Washington Ave., New York, NY, 10032, USA

^c Member RIFM Expert Panel, Malmö University Hospital, Department of Occupational & Environmental Dermatology, Sodra Forstadsgatan 101, Entrance 47, Malmö, SE-20502, Sweden

^d Member RIFM Expert Panel, School of Natural Resources & Environment, University of Michigan, Dana Building G110, 440 Church St., Ann Arbor, MI, 48109, USA

^e Member RIFM Expert Panel, Fraunhofer Institute for Toxicology and Experimental Medicine, Nikolai-Fuchs-Strasse 1, 30625, Hannover, Germany

^f Member RIFM Expert Panel, 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

^g Member RIFM Expert Panel, University of Würzburg, Department of Toxicology, Versbacher Str. 9, 97078, Würzburg, Germany

^h Member RIFM Expert Panel, Oregon Health Science University, 3181 SW Sam Jackson Park Rd., Portland, OR, 97239, USA

ⁱ Member RIFM Expert Panel, Vanderbilt University School of Medicine, Department of Biochemistry, Center in Molecular Toxicology, 638 Robinson Research Building, 2200 Pierce Avenue, Nashville, TN, 37232-0146, USA

^j Member of RIFM Expert Panel, 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

^k Member RIFM Expert Panel, The University of Tennessee, College of Veterinary Medicine, Department of Comparative Medicine, 2407 River Dr., Knoxville, TN, 37996-4500, USA

^l Member RIFM Expert Panel, Department of Pharmacology, University of Arizona, College of Medicine, 1501 North Campbell Avenue, P.O. Box 245050, Tucson, AZ, 85724-5050, USA

^m Member RIFM Expert Panel, The Journal of Dermatological Science (JDS), Editor-in-Chief, Professor and Chairman, Department of Dermatology, Hamamatsu University School of Medicine, 1-20-1 Handayama, Higashi-ku, Hamamatsu, 431-3192, Japan

* Corresponding author.

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

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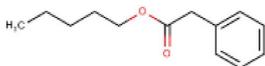
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Version: 050718. This version replaces any previous versions.

Name: Pentyl phenylacetate

CAS Registry Number: 5137-52-0



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., 2015, 2017) compared to a deterministic aggregate approach

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

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

QRA - Quantitative Risk Assessment

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

Pentyl phenylacetate was evaluated for genotoxicity, repeated dose toxicity, reproductive toxicity, local respiratory toxicity, phototoxicity, skin sensitization, and environmental safety. Data on read-across analog methyl phenylacetate (CAS # 101-41-7) show that pentyl phenylacetate is not expected to be genotoxic. The repeated dose, reproductive, and local respiratory toxicity endpoints were completed using the TTC for a Cramer Class I material, and the exposure to pentyl phenylacetate is below the TTC (0.03 mg/kg/day, 0.03 mg/kg/day, and 1.4 mg/day, respectively). Data from read-across analog methyl benzoate (CAS # 93-58-3) show that pentyl phenylacetate does not present a safety concern at the current, declared levels of use for the skin sensitization endpoint. The phototoxicity/photoallergenicity endpoint was completed based on UV spectra; pentyl phenylacetate is not expected to be phototoxic/photoallergenic. The environmental endpoints were evaluated; pentyl phenylacetate was found not to be PBT as per the IFRA Environmental Standards, and its risk quotients, based on its current volume of use in Europe and North America (i.e., PEC/PNEC), are < 1 .

Human Health Safety Assessment

Genotoxicity: Not genotoxic. (RIFM, 2001; RIFM, 2015)

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 under the current, declared levels of use. (ECHA REACH Dossier: Methyl benzoate, accessed 6/14/17)

Phototoxicity/Photoallergenicity: Not phototoxic/photoallergenic. (UV Spectra, RIFM DB)

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

Environmental Safety Assessment

Hazard Assessment:

Persistence: Screening-level: 3.1 (BIOWIN) (EPI Suite v4.1; US EPA, 2012a) 3)

Bioaccumulation: Screening-level: 216 L/kg (EPI Suite v4.1; US EPA, 2012a)

Ecotoxicity: Screening-level: Fish LC50: 5.063 mg/L (Salvito et al., 2002)

Conclusion: Not PBT or vPvB as per IFRA Environmental Standards

Risk Assessment:

Screening-level: PEC/PNEC (North America and Europe) < 1 (Salvito et al., 2002)

Critical Ecotoxicity Endpoint: Fish LC50: 5.063 mg/L (Salvito et al., 2002)

RIFM PNEC is: 0.005063 µg/L

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

1. Identification

- Chemical Name:** Pentyl phenylacetate
- CAS Registry Number:** 5137-52-0
- Synonyms:** Amyl phenylacetate; Benzeneacetic acid, pentyl acetate; Pentyl phenylacetate
- Molecular Formula:** C₁₃H₁₈O₂
- Molecular Weight:** 206.29
- RIFM Number:** 6211

2. Physical data

- Boiling Point:** 285.11 °C (EPI Suite)
- Flash Point:** 224.00°F. TCC (106.67 °C)*
- Log K_{ow}:** 4.05 (EPI Suite)
- Melting Point:** 42.34 °C (EPI Suite)
- Water Solubility:** 14.26 mg/L (EPI Suite)
- Specific Gravity:** 0.97700 to 0.98500 @ 25.00 °C.*
- Vapor Pressure:** 0.00164 mm Hg @ 20 °C (EPI Suite), 0.00298 mm Hg @ 25 °C (EPI Suite)

8. **UV Spectra:** No significant absorbance between 290 and 700 nm; molar absorption coefficient is below the benchmark ($1000 \text{ L} \cdot \text{mol}^{-1} \cdot \text{cm}^{-1}$)
9. **Appearance/Organoleptic:** Colorless to pale yellow clear liquid; balsam chocolate animal tobacco at 10.00% in dipropylene glycol*

* <http://www.thegoodscentcompany.com/data/rw1030731.html>.

3. Exposure

- Volume of Use (worldwide band):** < 0.1 metric tons per year (IFRA, 2015)
- 95th Percentile Concentration in Hydroalcohols:** 0.0022% (RIFM, 2017)
- Inhalation Exposure*:** 0.00019 mg/kg/day or 0.0015 mg/day (RIFM, 2017)
- Total Systemic Exposure**:** 0.00038 mg/kg/day (RIFM, 2017)

*95th percentile calculated exposure derived from concentration survey data in the Creme RIFM aggregate exposure model (Comiskey et al., 2015; Safford et al., 2015; Safford et al., 2017; and Comiskey et al., 2017).

**95th percentile calculated exposure; assumes 100% absorption unless modified by dermal absorption data as reported in Section IV. 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 et al., 2015; Safford et al., 2015; Safford et al., 2017; and Comiskey et al., 2017).

4. Derivation of systemic absorption

- Dermal:** Assumed 100%
- Oral:** Assumed 100%
- Inhalation:** Assumed 100%

5. Computational toxicology evaluation

- Cramer Classification:** Class I, Low

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

- Analogs Selected:
 - Genotoxicity:** Methyl phenylacetate (CAS # 101-41-7)
 - Repeated Dose Toxicity:** None
 - Reproductive Toxicity:** None
 - Skin Sensitization:** methyl benzoate (CAS # 93-58-3)
 - Phototoxicity/Photoallergenicity:** None
 - Local Respiratory Toxicity:** None
 - Environmental Toxicity:** None
- Read-across Justification: See [Appendix](#) below

6. Metabolism

Not considered for this risk assessment.

7. Natural occurrence (discrete chemical) or composition (NCS)

Pentyl phenylacetate is not reported to occur food 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 that contains information on published volatile compounds that

have been found in natural (processed) food products. Includes FEMA GRAS and EU-Flavis data.

8. IFRA standard

None.

9. REACH dossier

Dossier available; accessed 05/07/2018.

10. Summary

10.1. Human health endpoint summaries

10.1.1. Genotoxicity

Based on the current data, pentyl phenylacetate does not present a concern for genotoxicity.

10.1.1.1. Risk assessment. Pentyl phenylacetate was assessed in the BlueScreen assay and found negative for both cytotoxicity and genotoxicity, with and without metabolic activation (RIFM, 2013). There are no studies assessing the mutagenicity of pentyl phenylacetate. The mutagenic activity of read-across analog methyl phenylacetate (CAS # 101-41-7; see Section V) 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 TA97a, TA98, TA100, TA1535, and TA102 were treated with methyl phenylacetate 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 dose in the presence or absence of S9 (RIFM, 2001). Under the conditions of the study, methyl phenylacetate was not mutagenic in the Ames test, and this can be extended to pentyl phenylacetate.

There are no studies assessing the clastogenicity of pentyl phenylacetate. The clastogenic activity of read-across analog methyl phenylacetate 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 methyl phenylacetate in DMSO at concentrations up to 1500 µg/mL in the presence and absence of metabolic activation (S9) for 3 and 24 h. Methyl phenylacetate did not induce binucleated cells with micronuclei when tested up to the cytotoxic doses in either non-activated or S9-activated test systems (RIFM, 2015). Under the conditions of the study, methyl phenylacetate was considered to be non-clastogenic in the *in vitro* micronucleus test, and this can be extended to pentyl phenylacetate.

Based on the available data, pentyl phenylacetate does not present a concern for genotoxic potential.

Additional References: None.

Literature Search and Risk Assessment Completed On: 06/19/2017.

10.1.2. Repeated dose toxicity

There are insufficient repeated dose toxicity data on pentyl phenylacetate or any read-across materials. The total systemic exposure to pentyl phenylacetate is below the TTC for the repeated dose toxicity endpoint of a Cramer Class I material at the current level of use.

10.1.2.1. Risk assessment. There are no repeated dose toxicity data on pentyl phenylacetate or any read-across materials that can be used to support the repeated dose toxicity endpoint. The total systemic exposure to pentyl phenylacetate (0.38 µg/kg/day) is below the TTC (30 µg/kg bw/day; Kroes et al., 2007) for the repeated dose toxicity endpoint of a Cramer Class I material at the current level of use.

Additional References: None.

Literature Search and Risk Assessment Completed On: 06/05/2017.

10.1.3. Reproductive toxicity

There are insufficient reproductive toxicity data on pentyl phenylacetate or any read-across materials. The total systemic exposure to pentyl phenylacetate is below the TTC for the reproductive toxicity endpoint of a Cramer Class I material at the current level of use.

10.1.3.1. Risk assessment. There are no reproductive toxicity data on pentyl phenylacetate or any read-across materials that can be used to support the reproductive toxicity endpoint. The total systemic exposure to pentyl phenylacetate (0.38 µg/kg/day) is below the TTC (30 µg/kg bw/day; Kroes et al., 2007; Laufersweiler et al., 2012) for the reproductive toxicity endpoint of a Cramer Class I material at the current level of use.

Additional References: None.

Literature Search and Risk Assessment Completed On: 06/05/2017.

10.1.4. Skin sensitization

Based on the read-across material methyl benzoate (CAS # 93-58-3), pentyl phenylacetate does not present a safety concern for skin sensitization under the current, declared levels of use.

10.1.4.1. Risk assessment. Insufficient skin sensitization studies are available for pentyl phenylacetate. Based on the read-across analog methyl benzoate (CAS # 93-58-3; see Section V >), pentyl phenylacetate does not present a safety concern for skin sensitization under the current, declared levels of use. The chemical structures of these materials indicate that they would not be expected to react with skin proteins (Toxtree 2.6.13; OECD toolbox v3.4). Read-across analog methyl benzoate does not present a concern for skin sensitization. In a murine local lymph node assay, read-across analog methyl benzoate was found to be negative up to the maximum tested concentration of 100%, which resulted in a Stimulation Index (SI) of 2.98 (ECHA REACH Dossier: methyl benzoate, accessed 6/14/17). In guinea pigs, an open epicutaneous test and Freund's complete adjuvant test with read-across analog methyl benzoate did not present reactions indicative of sensitization (Klecak, 1979, 1985; Hausen et al., 1995). In a human maximization test for read-across analog methyl benzoate, no skin sensitization reactions were observed with 4% or 2760 µg/cm² in petrolatum (RIFM, 1970). Based on weight of evidence from structural analysis and animal and human studies, pentyl phenylacetate does not present a safety concern for skin sensitization under the current, declared levels of use.

Additional References: None.

Literature Search and Risk Assessment Completed On: 6/15/17.

10.1.5. Phototoxicity/photoallergenicity

Based on the available UV/Vis spectra, pentyl phenylacetate would not be expected to present a concern for phototoxicity or photoallergenicity.

10.1.5.1. Risk assessment. There are no phototoxicity studies available for pentyl phenylacetate 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 et al., 2009). Based on lack of absorbance, pentyl phenylacetate does not present a concern for phototoxicity or photoallergenicity.

10.1.5.2. 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⁻¹ · cm⁻¹ (Henry et al., 2009).

Additional References: None.

Literature Search and Risk Assessment Completed On: 05/25/17.

10.1.6. Local Respiratory Toxicity

The margin of exposure could not be calculated due to lack of appropriate data. The material, pentyl phenylacetate, exposure level is below the Cramer Class I TTC value for inhalation exposure local effects.

10.1.6.1. Risk assessment. There are no inhalation data available on pentyl phenylacetate. Based on the Creme RIFM Model, the inhalation exposure is 0.0015 mg/day. This exposure is 933 times lower than the Cramer Class I TTC value of 1.4 mg/day (based on human lung weight of 650 g; Carthew et al., 2009); therefore, the exposure at the current level of use is deemed safe.

Additional References: None.

Literature Search and Risk Assessment Completed On: 6/28/2017.

10.2. Environmental endpoint summary

10.2.1. Screening-level assessment

A screening-level risk assessment of pentyl phenylacetate was performed following the RIFM Environmental Framework (Salvito et al., 2002), which provides 3 tiered levels of screening for aquatic risk. In Tier 1, only the material's regional VoU, its log K_{OW}, 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 biodegradation 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, pentyl phenylacetate 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 v4.1 (US EPA, 2012a) did not identify pentyl phenylacetate as possibly persistent or 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 v4.1). Data on persistence and

bioaccumulation are reported below and summarized in the Environmental Safety Assessment section prior to Section 1.

10.2.2. Risk assessment

Based on the current Volume of Use (2015), pentyl phenylacetate does not present a risk to the aquatic compartment in the screening-level assessment.

Biodegradation: No data available.

Ecotoxicity: No data available.

Other available data: Pentyl phenylacetate has been pre-registered for REACH with no additional data at this time.

10.2.3. Risk assessment refinement

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

Endpoints used to calculate PNEC are underlined.

	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>5.063</u>			1,000,000	0.005063	

Exposure information and PEC calculation (following RIFM Environmental Framework: [Salvito et al., 2002](#)).

Exposure	Europe (EU)	North America (NA)
Log K_{ow} Used	3.12	3.12
Biodegradation Factor Used	0	0
Dilution Factor	3	3
Regional Volume of Use Tonnage Band	< 1	< 1
Risk Characterization: PEC/PNEC	< 1	< 1

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

The RIFM PNEC is 0.005063 µg/L. The revised PEC/PNECs for EU and NA: not applicable; cleared at the screening-level and therefore does not present a risk to the aquatic environment at the current reported volumes of use.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.fct.2018.11.031>.

Appendix

Read-across Justification

Methods

The read-across analogs were identified following the strategy for structuring and reporting a read-across prediction of toxicity described in [Schultz et al. \(2015\)](#). The strategy is also consistent with the guidance provided by OECD within Integrated Approaches for Testing and Assessment (OECD, 2015) and the European Chemical Agency read-across assessment framework ([ECHA, 2016](#)).

- First, materials were clustered based on their structural similarity. Second, data availability and data quality on the selected cluster were examined. Third, appropriate read-across analogs from the cluster were confirmed by expert judgment.
- Tanimoto structure similarity scores were calculated using FCFC4 fingerprints ([Rogers and Hahn, 2010](#)).

Literature Search and Risk Assessment Completed On: 6/16/17.

11. Literature Search*

- **RIFM Database:** Target, Fragrance Structure Activity Group materials, other references, JECFA, CIR, SIDS
- **ECHA:** <http://echa.europa.eu/>
- **NTP:** <https://ntp.niehs.nih.gov/>
- **OECD Toolbox**
- **SciFinder:** <https://scifinder.cas.org/scifinder/view/scifinder/scifinderExplore.jsf>
- **PubMed:** <http://www.ncbi.nlm.nih.gov/pubmed>
- **TOXNET:** <http://toxnet.nlm.nih.gov/>
- **IARC:** <http://monographs.iarc.fr>
- **OECD SIDS:** <http://webnet.oecd.org/hpv/ui/Default.aspx>
- **EPA ACToR:** <https://actor.epa.gov/actor/home.xhtml>

- **US EPA HPVIS:** 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:** <http://www.safe.nite.go.jp/english/db.html>
- **Japan Existing Chemical Data Base (JECDB):** http://dra4.nihs.go.jp/mhlw_data/jsp/SearchPageENG.jsp
- **Google:** <https://www.google.com>
- **ChemIDplus:** <https://chem.nlm.nih.gov/chemidplus/>

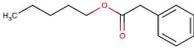
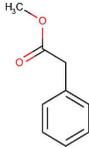
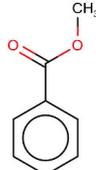
Search keywords: CAS number and/or material names.

*Information sources outside of RIFM's database are noted as appropriate in the safety assessment. This is not an exhaustive list.

Conflicts of interest

The authors declare that they have no conflicts of interest.

- The physical–chemical properties of the target substance and the read-across analogs were calculated using EPI Suite (US EPA, 2012a).
- J_{\max} values were calculated using RIFM's skin absorption model (SAM). The parameters were calculated using the consensus model (Shen et al., 2014).
- DNA binding, mutagenicity, genotoxicity alerts, and oncologic classification predictions were generated using OECD QSAR Toolbox v3.4 (OECD, 2012).
- ER binding and repeat dose categorization were generated using OECD QSAR Toolbox v3.4 (OECD, 2012).
- Developmental toxicity was predicted using CAESAR v2.1.7 (Cassano et al., 2010) and skin sensitization was predicted using Toxtree 2.6.13.
- Protein binding was predicted using OECD QSAR Toolbox v3.4 (OECD, 2012).
- The major metabolites for the target and read-across analogs were determined and evaluated using OECD QSAR Toolbox v3.4 (OECD, 2012).

	Target material	Read-across material	
Principal Name	Pentyl phenylacetate	Methyl phenylacetate	Methyl benzoate
CAS No.	5137-52-0	101-41-7	93-58-3
Structure			
Similarity (Tanimoto score)		0.74	0.61
Read-across endpoint		• Genotoxicity	• Skin Sensitization
Molecular Formula	$C_{13}H_{18}O_2$	$C_9H_{10}O_2$	$C_8H_8O_2$
Molecular Weight	206.29	150.18	136.15
Melting Point (°C, EPI Suite)	42.34	−0.50	−11.87
Boiling Point (°C, EPI Suite)	285.11	215.57	195.93
Vapor Pressure (Pa @ 25 °C, EPI Suite)	0.398	20.9	5.07E+001
Log Kow (KOWWIN v1.68 in EPI Suite)	4.05	1.83	2.12
Water Solubility (mg/L, @ 25 °C, WSKOW v1.42 in EPI Suite)	14.26	2072	2100
J_{\max} (mg/cm ² /h, SAM)	12.045	78.176	77.618
Henry's Law (Pa·m ³ /mol, Bond Method, EPI Suite)	4.45E+000	1.43E+000	3.52E+000
Genotoxicity			
DNA binding (OASIS v 1.4 QSAR Toolbox 3.4)	• No alert found	• No alert found	
DNA binding by OECD QSAR Toolbox (3.4)	• Michael addition	• Michael addition	
Carcinogenicity (genotox and non-genotox) alerts (ISS)			
	• Non-carcinogen (low reliability)	• Non-carcinogen (moderate reliability)	
DNA alerts for Ames, MN, CA by OASIS v 1.1	• No alert found	• No alert found	
<i>In vitro</i> Mutagenicity (Ames test) alerts by ISS	• No alert found	• No alert found	
<i>In vivo</i> mutagenicity (Micronucleus) alerts by ISS	• No alert found	• No alert found	
Oncologic Classification	• Not classified	• Not classified	
Skin Sensitization			
Protein binding by OASIS v1.1	• No alert found		• Acylation
Protein binding by OECD	• No alert found		• No alert found
Protein binding potency	• Not possible to classify		• Not possible to classify
Protein binding alerts for skin sensitization by OASIS v1.1	• No alert found		• No alert found
Skin Sensitization reactivity domains (ToxTree v2.6.13)	• No alert found		• No alert found
Metabolism			
OECD QSAR Toolbox (3.4)	See Supplemental Data 1	See Supplemental Data 2	See Supplemental Data 3
Rat liver S9 metabolism simulator and structural alerts for metabolites			

Summary

There are insufficient toxicity data on the target material pentyl phenylacetate (CAS # 5137-52-0). Hence, *in silico* evaluation was conducted by determining a read-across analog for this material. Based on structural similarity, reactivity, metabolism data, physical–chemical properties, and expert judgment, analogs methyl phenylacetate (CAS # 101-41-7) and methyl benzoate (CAS # 93-58-3) were identified as read-across materials with sufficient data for toxicological evaluation.

Conclusion

- Methyl phenylacetate (CAS # 101-41-7) was used as a read-across analog for the target material pentyl phenylacetate (CAS # 5137-52-0) for the genotoxicity endpoint.
 - The target substance and the read-across analog are structurally similar and belong to the structural class of aromatic esters.
 - The target substance and the read-across analog share an aromatic acid portion and a straight chain saturated alcohol portion.
 - The key difference between the target substance and the read-across analog is that the target substance has a C5 alcohol portion while the read-across analog has a C1 (methyl alcohol) alcohol portion.
 - Similarity between the target substance and the read-across analog is indicated by the Tanimoto score in the table above. Differences between the structures that affect the Tanimoto score do not affect consideration of the toxicity endpoint.
 - The physical–chemical properties of the target substance and the read-across analog are sufficiently similar to enable comparison of their

- toxicological properties.
- According to the QSAR OECD Toolbox (v3.4), structural alerts for the toxicological endpoint are consistent between the target substance and the read-across analog.
 - The target substance and the read-across analog have a DNA binding alert by OECD. This shows that the read-across analog and the target substance have comparable reactivity. The data described in the genotoxicity section show that the read-across analog does not pose a concern for this endpoint. Therefore, the alert will be superseded by the available data.
 - The target substance and the read-across analog are expected to be metabolized similarly, as shown by the metabolism simulator.
 - The structural differences between the target material and the read-across analog do not affect consideration of the toxicological endpoint.
 - Methyl benzoate (CAS # 93-58-3) was used as a read-across analog for the target material pentyl phenylacetate (CAS # 5137-52-0) for the skin sensitization endpoint.
 - The target substance and the read-across analog are structurally similar and belong to the structural class of aromatic esters.
 - The target substance and the read-across analog share an aromatic acid portion and straight chain saturated alcohol portion.
 - The key difference between the target substance and the read-across analog is that the target substance has a C5 alcohol portion while the read-across analog has a C1 (methyl alcohol) alcohol portion. In addition to the acid portion, the target substance has a carboxylic acid insulated by 1 carbon from the aromatic moiety while the carboxylic acid in the read-across analog has a carboxylic acid conjugated with the aromatic moiety. The carboxylic acid in the read-across analog is a stronger acid (lower pKa) compared to the acid portion in the target substance (higher pKa). This difference between the target substance and the read-across analog makes the read-across analog more reactive compared to the target substance.
 - Similarity between the target substance and the read-across analog is indicated by the Tanimoto score in the table above. Differences between the structures that affect the Tanimoto score do not affect consideration of the toxicological endpoint.
 - The physical–chemical properties of the target substance and the read-across analog are sufficiently similar to enable comparison of their toxicological properties.
 - According to the QSAR OECD Toolbox (v3.4), structural alerts for the toxicological endpoint are consistent between the target substance and the read-across analog.
 - The read-across analog has a protein binding alert by OASIS. This shows that the read-across analog is more reactive than the target substance. The data described in the skin sensitization section show that the read-across analog does not pose a concern for skin sensitization. Therefore, the alert will be superseded by the available data.
 - The target substance and the read-across analog are expected to be metabolized similarly, as shown by the metabolism simulator.
 - The structural differences between the target material and the read-across analog do not affect consideration of the toxicological endpoints.

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