

# **Amended Final Safety Assessment**

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## **Benzyl Alcohol, and Benzoic Acid and its Salts and Benzyl Ester**

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### **Cosmetic Ingredient Review**

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## **ABSTRACT**

*Benzyl alcohol, benzoic acid, and its salts and benzyl benzoate function mostly as fragrance ingredients/preservatives in cosmetic products. The Panel previously issued a final report with concentration limits for benzyl alcohol, benzoic acid, and sodium benzoate in cosmetics, and also determined that the available data were insufficient to support the safety of these ingredients during inhalation exposure. After reviewing the available data in the present report, it was concluded that benzyl alcohol, benzoic acid, and its salts and benzyl ester are safe in the present practices of use and concentration described in this safety assessment.*

## **INTRODUCTION**

This report assesses the safety of benzyl alcohol, and benzoic acid and its salts and benzyl ester as used in cosmetics. In cosmetics, these ingredients are reported to function as fragrance ingredients, pesticides, pH adjusters, preservatives, solvents, and/or viscosity decreasing agents in cosmetic products.

In 2001, the Cosmetic Ingredient Review (CIR) published a final report on the safety assessment of benzyl alcohol, benzoic acid, and sodium benzoate with the conclusion that benzyl alcohol, benzoic acid, and sodium benzoate are safe for use in cosmetic formulations at concentrations up to 5%.<sup>1</sup> Benzyl alcohol was safe for use in hair dyes at concentrations up to 10%. The available data were insufficient, however, to support the safety of these ingredients in cosmetic products in which a primary route of exposure is inhalation. The Expert Panel stated that inhalation toxicity data were needed.

The Personal Care Products Council provided a 4-week inhalation toxicity study (rats) of aerosolized benzyl alcohol and benzoic acid. After reviewing this study, the Expert Panel determined that the final report on the safety assessment of benzyl alcohol, benzoic acid, and sodium benzoate should be reopened to amend the original conclusion. Furthermore, the Panel determined that data available in the original safety assessment, with the addition of the new inhalation toxicity data, were sufficient to support the safety of other simple salts of benzoic acid (calcium, magnesium, and potassium benzoate) and the benzyl ester.

Additional new data reviewed included an Organization for Economic Cooperation and Development (OECD) Screening Information Data Sets (SIDS) for high volume chemicals initial assessment report on benzoic acid, sodium benzoate, potassium benzoate, and benzyl alcohol published by the United Nations Environmental Program Chemicals Branch (UNEP chemicals) in 2001,<sup>2</sup> the same year that the CIR Final Safety Assessment on benzyl alcohol, benzoic acid, and sodium benzoate was published. In addition to the SIDS data summaries included in this CIR safety assessment, the 2001 SIDS report may be consulted for further details. Further, the European Commissions' International Uniform Chemical Information Database (IUCLID) on benzyl alcohol, benzoic acid, sodium benzoate, and benzyl benzoate was created in the year 2000.<sup>3</sup> Data from the OECD SIDS assessment and IUCLID data sets are summarized in the Toxicology section. In the process of selecting new data for inclusion in the current report, efforts were made to eliminate redundancy between OECD SIDS, IUCLID, or other data found in the published literature and data included in the original final report.

## **CHEMISTRY**

### **Definition and Structure**

Definitions, other chemical names, and cosmetic ingredient functions for ingredients reviewed in this safety assessment are included in Table 1. Structural formulas are included in Figure 1.

### **Chemical and Physical Properties**

Chemical and physical properties of benzyl alcohol, benzoic acid and its salts, and the benzyl benzoate ester are included in Table 2.

The photodegradation of benzoic acid aqueous solution in the presence of UV light has been reported. Benzoic Acid absorbs UV light below 300 nm, and its concentration in solution was monitored using reverse-phase high-performance liquid chromatography (HPLC). The mercury lamp used emitted polychromatic radiation between 240 and 540 nm. The photon flow absorbed by the reaction system varied with the concentration of benzoic acid and with the presence of the

products of degradation. When the temperature was increased from 25°C to 50°C, photodegradation was increased by 20%. Photodegradation products were not mentioned.<sup>4</sup>

The conversion of benzyl alcohol to benzaldehyde and benzoic acid in aqueous solution has been demonstrated using liquid chromatography/spectrophotometry.<sup>5</sup> When benzyl benzoate (in ethanol) was exposed to sunlight for 11 days (at 25°C), 35 products were isolated from exposed samples and identified using gas chromatography/mass spectrometry. Benzyl alcohol and benzoic acid were among the products identified.<sup>6</sup>

### **Method of Manufacture**

#### **Benzyl Alcohol**

Large scale production of benzyl alcohol is achieved by the action of sodium or potassium carbonate on benzyl chloride.<sup>1</sup>

#### **Benzoic Acid**

Benzoic acid is produced via the decarboxylation of phthalic anhydride in the presence of catalysts. Another production method involves the chlorination of toluene to yield benzotrichloride, which is hydrolyzed to benzoic acid.<sup>7</sup>

#### **Sodium Benzoate**

Sodium benzoate is produced by the neutralization of benzoic acid with sodium bicarbonate, sodium carbonate, or sodium hydroxide.<sup>8</sup>

#### **Potassium Benzoate**

Potassium benzoate can be prepared by reacting methyl benzoate with potassium thioacetate.<sup>9</sup>

#### **Benzyl Benzoate**

Benzyl benzoate results from the condensation of benzoic acid and benzyl alcohol. It can also be generated from benzaldehyde via the Tishchenko reaction.<sup>10</sup>

### **UV Absorption**

The following UV absorption maxima have been reported for salts and an ester of benzoic acid: 228.6 nm (magnesium benzoate), 226.8 nm (potassium benzoate), and 256 nm (benzyl benzoate).<sup>11,12</sup>

### **Analytical Methods**

Benzoic acid has been analyzed using mass spectrometry,<sup>13</sup> and potassium benzoate has been analyzed using IR spectroscopy.<sup>14</sup> Benzyl benzoate has been analyzed using IR and NMR spectroscopy.<sup>15,16</sup>

### **Impurities**

#### **Benzoic Acid and Sodium Benzoate**

The effects of UVA exposure on benzene formation have been determined. Benzene formation was examined for samples contained in UV stabilized and non-UV stabilized packaging.<sup>17</sup> Some of the samples selected for UVA testing included model solutions prepared with 0.04% benzoate and 0.025% ascorbic acid in unbuffered water. The results of 24-h irradiation studies indicated that, under intense UV light, benzene levels increased by as much as 53% (to 315 ng/g benzene, compared to model solutions exposed to visible light [206 ng/g benzene]) in model solutions stored in non-UV stabilized bottles. However, the use of UV stabilized polyethylene terephthalate bottles reduced benzene formation by ~ 13%, relative to the non-UV stabilized bottles. Similar trends were observed following irradiation for 7 days. According to the Food and Drug Administration (FDA), the U.S. beverage industry voluntarily reformulated beverages that were found to contain benzene levels at or above the 5 ng/g maximum contaminant level (mcl) for drinking water established by the U.S. Environmental Protection Agency (EPA).<sup>17</sup> An mcl of 0.005 mg/L (5 ppb) is the EPA's official limit for benzene in drinking water.<sup>18</sup> Furthermore, in a 2008 survey conducted by FDA, all of the reformulated beverages tested were shown to contain ≤ 1.1 ng/g benzene. Most of the beverages were reformulated by removing or reducing either benzoate or ascorbic acid, or by adding EDTA.<sup>19</sup>

## **Benzyl Benzoate, Potassium Benzoate, and Sodium Benzoate**

According to the *United States Pharmacopoeia*, benzyl benzoate contains less than 99% and not more than 100.5% C<sub>14</sub>H<sub>12</sub>O<sub>2</sub> and potassium benzoate contains not less than 100.5% and not more than 99% C<sub>7</sub>H<sub>5</sub>KO<sub>2</sub>. Limits for water and heavy metals in potassium benzoate are stated as ≤ 1.5% and ≤ 0.001%, respectively.<sup>20</sup>

The Food Chemicals Codex specification for sodium benzoate includes the following limitations: heavy metals, as Pb (≤ 2 mg/kg), alkalinity, as NaOH (≤ 0.04%), and water (≤ 1.5%).<sup>21</sup>

### **USE**

#### **Cosmetic**

As stated in the *International Cosmetic Ingredient Dictionary and Handbook*, benzoic acid, benzyl alcohol, sodium benzoate, calcium benzoate, magnesium benzoate, potassium benzoate, and benzyl benzoate may function as fragrance ingredients, pesticides, pH adjusters, preservatives, solvents, viscosity decreasing agents, and pH adjusters in cosmetic products (See Table 1).<sup>22</sup>

According to information supplied to the Food and Drug Administration (FDA) by industry as part of the Voluntary Cosmetic Registration Program (VCRP) in 2010, the following ingredients were being used in personal care products: benzoic acid, benzyl alcohol, sodium benzoate, potassium benzoate, and benzyl benzoate.<sup>23</sup> Uses of calcium benzoate and magnesium benzoate were not reported to the VCRP. These data are summarized in Table 3.

Independent of the VCRP data, a survey of ingredient use concentrations was conducted by the Personal Care Products Council in 2010. The results of this survey indicated that the following ingredients were being used in cosmetic products: benzoic acid (0.000002 to 5%), benzyl alcohol (0.000006 to 10%), sodium benzoate (0.000001 to 1%), calcium benzoate (0.002 to 0.004%), potassium benzoate (0.002 to 0.003%), and benzyl benzoate (0.000005 to 4%).<sup>24</sup> No uses of magnesium benzoate were reported in the Council survey.

Personal care products containing the ingredients reported as being used may be applied to the skin, nails, or hair, or, incidentally, may come in contact with the eyes and mucous membranes. Products containing these ingredients may be applied as frequently as several times per day and may come in contact with the skin, nails, or hair for variable periods following application. Daily or occasional use may extend over many years.

In the European Union, the maximum authorized concentrations for benzoic acid and sodium benzoate as preservatives in cosmetic products are: rinse-off products, except oral care products (2.5%, as acid), oral care products (1.7%, as acid), and leave-on products (0.5%, as acid). For salts of benzoic acid other than sodium benzoate, the maximum authorized concentration is 0.5% (as acid).<sup>25</sup>

The Scientific Committee on Cosmetic Products and Non-Food Products Intended for Consumers (SCCNFP) was asked to review the data submitted to support the safety of benzoic acid and its salts and esters, when used at concentrations other than those laid down in Annex VI to Directive 76/768/EEC as preservatives, for other specific non-preservative purposes apparent from the presentation of the products and to answer the following question: Can benzoic acid and its salts and esters be used safely for non-preservative purposes in cosmetic rinse-off products at a maximum concentration of 2.5% and in cosmetic oral-care products at a maximum concentration of 1.7%. Data were provided to the SCCNFP in support of the use of benzoic acid as a non-preservative. The SCCNFP did not find the submission appropriate for a safety evaluation of benzoic acid and sodium benzoate for the applied “other uses” in cosmetic products.<sup>26</sup> Subsequently, a second data set was provided, and these data were considered adequate for arriving at a conclusion on the safety of benzoic acid and sodium benzoate, but not the other salts or esters. Based on these data, the Scientific Committee on Consumer Products (SCCP) issued the following opinion: The SCCP is of the opinion that benzoic acid and sodium benzoate are safe for use for preservative and non-preservative purposes in cosmetic rinse-off products at a maximum concentration of 2.5 % and in cosmetic oral-care products at a maximum concentration of 1.7%, and in leave-on products up to 0.5%.<sup>27</sup>

#### **Noncosmetic**

According to FDA’s Select Committee on Generally Recognized as Safe (GRAS) substances, benzoic acid (21 CFR [Code of Federal Regulations] 184.1021)<sup>28</sup> and sodium benzoate (21 CFR 184.1733)<sup>29</sup> are classified as GRAS food substances. Sodium benzoate has also been approved by FDA for use as an antimycotic when migrating from food-packaging material (21 CFR 181.23).<sup>30</sup> Other CFR citations relating to FDA-approved direct/indirect food additive

ingredient uses include: benzyl alcohol (21 CFR 172.515, 175.105, 175.300, and 177.1210)<sup>31,32,33,34</sup>; benzoic acid (21 CFR 150.141, 150.161, 166.110, 175.300, and 177.1390)<sup>35,36,37,33,38</sup>; sodium benzoate (21 CFR 150.141, 150.161, and 166.110)<sup>35,36,37</sup>; calcium benzoate (21 CFR 166.110 and 178.2010)<sup>37,39</sup>; potassium benzoate (21 CFR 166.110 and 177.1210)<sup>37,34</sup>; and benzyl benzoate (21 CFR 172.515 and 175.105).<sup>31,32</sup>

Benzyl alcohol is an FDA-approved diluent in color additive mixtures for external drug use.<sup>40</sup> In 2009, the FDA approved benzyl alcohol lotion, 5%, as a prescription medication for the treatment of head lice (*Pediculosis capitis*) infestation in patients 6 months of age and older.<sup>41</sup> The results of clinical trials supporting benzyl alcohol lotion 5% (Ulesfia™) as a safe and effective topical treatment for head lice are included in the section on Skin Irritation and Sensitization later in the report text.<sup>42</sup> Benzyl benzoate (10%) has been used in the treatment of scabies in developing countries.<sup>43</sup>

Benzyl alcohol has been classified by FDA regarding its use in the following types of over-the-counter (OTC) drug products (Category I: generally recognized as safe (S) and effective (E) for the claimed therapeutic indication; Category II: not generally recognized as safe and effective or unacceptable indications; Category III: insufficient data available to permit final classification):<sup>44</sup>

- anorectal (final monograph: 1 to 4% use concentration range established; Category I)
- external analgesic (both Category I and final monograph pending)
- external analgesic (final monograph: Category III)
- oral health care (Category I pending)
- oral discomfort care (Category II pending)
- oral discomfort care (Category IIISE pending)
- pediculicide (final monograph: Category III)
- skin protectant (final monograph: Category III)

Similarly, benzoic acid has been classified by FDA regarding its use in the following types of OTC drug products:<sup>44</sup>

- acne (final monograph: Category III)
- antifungal (final monograph: Category III)
- oral health care - (Category IIISE pending)
- skin protectant - (final monograph: Category III)

Sodium benzoate has been classified by FDA regarding its use in menstrual/diuretic OTC drug products (final monograph: Category III), and benzyl benzoate has been approved for use in OTC pediculicides (final monograph: Category II).<sup>44</sup>

Calcium benzoate, magnesium benzoate, and potassium benzoate are inert non-food ingredients that, under the Federal Insecticide, Fungicide, and Rodenticide Act, are permitted by the Environmental Protection Agency (EPA) for use in non-food use pesticide products.<sup>45</sup> Pesticide products containing benzoic acid as the active ingredient have been registered with EPA for use in the extermination of dust mites.<sup>46</sup>

Benzyl alcohol has been listed as a chemical in photographic developing systems,<sup>47</sup> and, at concentrations of 0.9 to 2%, is commonly used as an antibacterial agent in a variety of pharmaceutical formulations intended for intravenous administration.<sup>48</sup> Additionally, benzyl alcohol has been used in parenteral medications commonly administered to critically ill neonates.<sup>49</sup>

## **TOXICOKINETICS**

### **Oral Studies**

#### **Benzyl Alcohol, Benzoic Acid, Sodium Benzoate, and Potassium Benzoate**

Benzyl alcohol is metabolized to benzoic acid. Both benzoic acid and sodium benzoate are rapidly absorbed from the gastrointestinal tract of mammals and conjugated with glycine in the liver; the resulting hippuric acid is rapidly excreted in the urine.<sup>1</sup> Benzyl alcohol and benzoic acid and its sodium and potassium salt can be considered as a single category regarding human health, because they are all rapidly metabolized and excreted via a common pathway within 24 h.<sup>2</sup>

#### **Benzyl Benzoate**

Following oral administration (gavage, 0.5, 1, and 2 ml/kg) of benzyl benzoate to fasted female dogs, urine was collected for 5 to 6 days. All 3 doses resulted in creatinuria and urinary excretion of glucuronate conjugates.<sup>50</sup>

Following the administration of unspecified oral doses of benzyl benzoate to cats and dogs (number and breed not stated), small amounts of hippuric acid were detected in the 24 h urine. The metabolism of benzyl benzoate to benzoic acid *in vivo* has also been reported. Additional details for these 2 studies were not provided.<sup>3</sup>

In a human subject, urinary excretion of hippuric acid was measured after oral dosing (single dose, 2.02 g) with benzyl benzoate. Urine was collected for 6 h. Approximately 90% of the dose administered was excreted as hippuric acid in the urine. After oral dosing with 1 g, approximately 80% of the administered dose was excreted as hippuric acid. In a second subject, 71% of the administered dose was excreted as hippuric acid during the first 6 h and an additional 14% of the dose was excreted as such during the second 6 h after dosing.<sup>51</sup>

## Dermal Studies

### Benzoic Acid

The penetration of benzoic acid through excised dorsal skin (full-thickness and with stratum corneum removed) from guinea pigs was examined. Skin preparations were mounted in a two-chamber diffusion cell. A suspension of excess amount of benzoic acid in saline was added to donor compartments. In full-thickness skin, permeation proceeded with a short lag time. Solubility ( $C_d$ , unit = mM) and permeability ( $K_p$ , unit =  $\times 10^{-2} \text{ cm} \cdot \text{h}^{-1}$ ) coefficients for benzoic acid were  $32.7 \pm 1.6$  and  $9.01 \pm 1.51$ , respectively. Removal of the stratum corneum by tape stripping and its delipidation using an organic solvent mixture enhanced the skin penetration of benzoic acid.<sup>52</sup>

### Benzyl Alcohol and Benzyl Benzoate

The percutaneous absorption of [ $7\text{-}^{14}\text{C}$ ]benzyl alcohol and [ $7\text{-}^{14}\text{C}$ ]benzyl benzoate *in vivo* was evaluated using 4 female Rhesus monkeys (10 to 19 years old). Each chemical was applied ( $4 \mu\text{g}/\text{cm}^2$ , without or with occlusion [glass chamber - G or plastic wrap - P]) to a clipped  $1 \text{ cm}^2$  area of abdominal skin; the vehicle was acetone ( $10$  to  $20 \mu\text{l}/\text{cm}^2$ ). The animals remained in metabolism chairs/metabolism cages for 5 days. The amount of absorbed compound in the urine was determined by liquid scintillation counting. Mean values ( $\pm$  SEM, 4 animals) for urinary excretion of the administered dose were:  $77.4 \pm 4.3\%$  (benzyl alcohol) and  $65.3 \pm 13.4\%$  (benzyl benzoate). Percutaneous absorption values (absorption, as % of applied dose) were as follows: benzyl alcohol ( $31.6 \pm 4.2\%$  [unoccluded];  $56.3 \pm 14.5\%$  [P];  $79.9 \pm 7.4\%$  [G]) and benzyl benzoate ( $57.0 \pm 10.4\%$  [unoccluded];  $71.2 \pm 4.4\%$  [P];  $64.7 \pm 10.2\%$  [G]). Over a 5-day period, the urinary excretion of radioactivity was  $65.3 \pm 13.4\%$  following dermal application of benzyl benzoate and  $56.5 \pm 7.7\%$  following dermal application of benzyl alcohol. When the acetone vehicle was replaced with a lotion ( $10 \text{ mg}/\text{cm}^2$ ), percutaneous absorption was increased. The increase was significant for benzyl benzoate ( $P < 0.05$ ), but not benzyl alcohol. There was no apparent correlation between the skin penetration of benzyl alcohol and benzyl benzoate and their octanol-water partition coefficients of 7.4 and 3.97, respectively.<sup>53</sup>

The percutaneous absorption of benzyl benzoate *in vitro* was evaluated using human epidermis (from abdominal skin) placed in a glass diffusion chamber. Saline (5 ml) was added to the chamber and was in contact with the bottom of the epidermis. Benzyl benzoate (0.2 ml) was applied to the top of the epidermis. Six experiments were performed and the mean value for skin penetration of benzyl benzoate was  $0.018 \pm 0.002\%$ .<sup>54</sup>

The penetration of benzyl alcohol through split-thickness cadaver skin was evaluated using nonoccluded Franz diffusion cells. Benzyl alcohol (spiked with  $^{14}\text{C}$  radiolabel) in ethanol was applied to the skin at doses ranging from 0.9 to  $10,600 \mu\text{g}/\text{cm}^2$ . After 24 h, the percentage of radioactivity that penetrated the skin increased gradually with dose, ranging from  $19.8 \pm 2.9\%$  (lowest dose) to  $29.2 \pm 3.0\%$  (highest dose). Also, less than 4% of the administered radioactivity was retained in the tissues at 24 h, and evaporation accounted for the remaining percentage. Data from this study were also analyzed in relation to a finite dose diffusion/evaporation model. The results of this analysis indicated that the increase in benzyl alcohol absorption with dose was consistent with a 3-fold increase in diffusivity in the stratum corneum, as the concentration of benzyl alcohol increased from tracer levels to saturation.<sup>55</sup>

## In Vitro Study

The rate of hydrolysis of benzyl benzoate (in acetonitrile) by 80% human plasma *in vitro* was studied. The  $t_{1/2}$  for the *in vitro* hydrolysis of benzyl benzoate to benzoic acid in human plasma was 19 minutes.<sup>56</sup>

## **TOXICOLOGICAL STUDIES**

An Organization for Economic Cooperation and Development (OECD) Screening Information Data Sets (SIDS) for high volume chemicals initial assessment report on benzoic acid, sodium benzoate, potassium benzoate, and benzyl alcohol was published by the United Nations Environmental Program, Chemicals Branch (UNEP chemicals) in 2001, with the following recommendation relating to human safety: "Taking into account the rapid metabolism and excretion, the non-bioaccumulation, the low toxicity after acute and repeated exposures, the non-reproductive toxicity, the non-genotoxicity and the non-carcinogenicity, the low irritating and non- to very low sensitizing properties of these substances, as well as the controlled (industrial settings) and /or regulated (pharma, cosmetics and /or food) uses, these substances will pose a minimal risk to humans (workers and consumers)." Because the database for this safety assessment is similar to that for the 2001 CIR Final Report on Benzyl Alcohol, Benzoic Acid, and Sodium Benzoate, the available data in the OECD SIDS assessment are summarized with minimal details in sections that follow.

### **Acute and Repeated Dose Inhalation Toxicity**

#### **Benzyl Alcohol and Benzoic Acid**

According to the OECD SIDS initial assessment report on Benzoates, 4 h of inhalation exposure to benzyl alcohol or benzoic acid at 4 and 12 mg/L aerosol/dust, respectively, did not cause death in rats. Thus, low acute toxicity was associated with these compounds.<sup>2</sup>

A 4-week inhalation toxicity study of aerosolized benzyl alcohol and benzoic acid was performed using groups of Crl:CD(SD) rats.<sup>57</sup> Four groups of rats (10/sex/group) were exposed (nose-only) to aerosolized benzyl alcohol 5 days per week (6 h/day) for 4 weeks. Each animal received a minimum of 20 exposures of benzyl alcohol, and target exposure concentrations were 30, 100, 300, and 1,000 mg/m<sup>3</sup> for the four groups. Two additional groups of rats were exposed to aerosolized benzoic acid (2.5 and 12.5 mg/m<sup>3</sup>) according to the same procedure. The group designated for exposure to benzyl alcohol (30 mg/m<sup>3</sup> target) was actually exposed to a mean atmosphere concentration that was 136% of the target concentration. For the remaining groups, all mean atmosphere concentrations were within 96.7% to 107.2% of the respective target concentrations. A concurrent control group was exposed to filtered air only.

There were no test-substance related deaths or effects on any of the following: body weight, food consumption, clinical pathology parameters, and organ weights. Additionally, there were no test substance-related macroscopic or microscopic findings. The (NOEL) and NOAEL were considered to be 1,072 mg/m<sup>3</sup> for benzyl alcohol and 12.6 mg/m<sup>3</sup> for benzoic acid.<sup>57</sup>

### **Acute Oral Toxicity**

Acute oral toxicity data are summarized in Table 4.

#### **Benzyl Alcohol, Benzoic Acid, Sodium Benzoate, and Potassium Benzoate**

According to the OECD SIDS initial assessment report on Benzoates, low acute oral toxicity is associated with benzoic acid and its sodium and potassium salts. The LD50 values were > 2000 mg/kg body weight, except for benzyl alcohol (LD50 = 1610 mg/kg [rats]), which needs to be considered harmful via the oral route.<sup>2</sup>

### **Acute Intraperitoneal Toxicity**

#### **Benzyl Benzoate**

Mice were dosed i.p. with benzyl benzoate (test procedure/number of mice not stated), and an LD50 of > 500 mg/kg body weight was reported.<sup>3</sup>

### **Acute Dermal Toxicity**

#### **Benzyl Alcohol, Benzoic Acid, Sodium Benzoate, and Potassium Benzoate**

According to the OECD SIDS initial assessment report on Benzoates, low acute dermal toxicity is associated with benzyl alcohol and benzoic acid and its sodium and potassium salts.<sup>2</sup>

## **Benzyl Alcohol and Benzyl Benzoate**

The acute dermal toxicity of benzyl benzoate was evaluated using rats (number not stated), and an LD50 of 4,000 mg/kg body weight was reported. Study details were not included. In a study involving rabbits, the acute dermal toxicity of undiluted benzyl benzoate was evaluated according to the Draize procedure (no GLP; number of animals not stated), and an LD50 of 4,448 mg/kg body weight was reported.<sup>3</sup> In another study involving rabbits, acute dermal LD50s of 2,000 mg/kg and 4,000 mg/kg have been reported for benzyl alcohol and benzyl benzoate, respectively.<sup>58</sup>

Single dermal applications of 25% benzyl benzoate (20 ml in isopropanol/water) to the backs of cats resulted in death. Additional study details were not provided.<sup>3</sup>

### **Acute Subcutaneous Toxicity**

#### **Benzyl Benzoate**

Groups of adult C57BL/6J mice (5 per group) were injected s.c. with 0.1 ml benzyl benzoate (~111 mg) daily for 4 weeks. An untreated group of sham injected animals served as the untreated concurrent control group. The adrenal glands were prepared for light and electron microscopic examination. The animals died within 3 h of initial injection. Additional study details were not provided.<sup>3</sup>

Guinea pigs (number not stated) received single s.c. injections of benzyl benzoate, and mortality rates at the administered doses were as follows: 0 (0.5 g/kg), 30% (1 g/kg), 80% (5 g/kg), 100% (10 g/kg), and 100% (20 g/kg).<sup>59</sup>

### **Acute Intramuscular Toxicity**

#### **Benzyl Benzoate**

Following benzyl benzoate intramuscular injection into cats (number not stated) i.m., doses of 2 and 10 g/kg were described as lethal.<sup>59</sup>

### **Repeated Dose Toxicity – Oral Studies**

#### **Benzyl Alcohol, Benzoic Acid, Sodium Benzoate, and Potassium Benzoate**

According to the OECD SIDS initial assessment report on Benzoates, repeated dose oral toxicity studies yielded an NOAEL of 800 mg/kg/day for benzoic acid and > 1000 mg/kg/day for the salts. At higher doses, increased mortality, decreased weight gain, and liver and kidney effects were observed. Systemic toxic effects of a similar nature (e.g. liver, kidney) were observed after dosing with benzyl alcohol, benzoic acid, sodium benzoate, and potassium benzoate. However, these effects were observed at higher doses of benzoic acid and its salts when compared to dosing with benzyl alcohol. Long-term studies on benzyl alcohol yielded an NOAEL of > 400 mg/kg body weight per day for rats and > 200 mg/kg body weight per day for mice. At higher doses, effects on body weight and lesions in the brain, thymus, skeletal muscle, and kidneys were observed. It was noted that one should take into account that oral administration was by gavage in these studies, whereby saturation of metabolic pathways is likely to occur. It was concluded that benzoic acid and its salts exhibited very low repeated dose toxicity and that benzyl alcohol exhibited low repeated dose toxicity.<sup>2</sup>

#### **Benzyl Benzoate**

No effects were observed in a 7-month toxicity study in which mice (no. and strain not stated) were dosed orally with benzyl benzoate (800 mg/kg).<sup>60</sup>

### **Repeated Dose Toxicity – Dermal Studies**

#### **Benzyl Benzoate**

The following effects were observed in groups of rats (3 males/3 females/group; strain not stated) that received dermal applications of 0.188, 0.488, 0.781, 1.25, or 2.0 g/kg for 30 days: hyperplasia of squamous epithelium, degeneration of hair follicles and sebaceous glands, subcutaneous fibrosis, and hyperplasia of the thyroid gland. Seven areas of the shaved back were dosed in rotation. Dosing with 2.0 g/kg also caused 4 deaths, decreased white blood cell numbers, and other blood effects. Body weight changes, decreased white blood cell numbers, and other blood effects were also observed following dosing with 1.25 g/kg. Additional study details were not provided.<sup>61</sup>



In a study involving rabbits, (non-GLP; number and strain not stated) dermal doses of benzyl benzoate were applied daily for 90 days.<sup>3</sup> The range of doses tested was not stated. Deaths occurred at doses of 2.2 g/kg body weight/day and greater. Testicular atrophy was noted at high doses, and, possibly, there was also an increased incidence of focal nephritis and encephalitis. Very mild skin irritation was also observed. Additional details relating to this study were not provided. Benzyl benzoate was also applied topically to rabbits at doses of 0.5 ml/kg/day daily for 90 days. Slight dermatitis and inanition were observed, and, at higher doses, slight to moderate testicular atrophy was apparent. Increased leucocyte counts were reported and results were also suggestive of kidney damage. Benzyl benzoate also caused death, but the number of mortalities and corresponding toxic doses were not reported. Repeated dermal applications of benzyl benzoate (doses not stated) did not cause toxic symptoms in the following species: dogs, horses, cows (females), sheep, or pigs. Additional study details were not provided.<sup>3</sup>

In another study, rabbits (no. and strain not stated) received dermal applications of benzyl benzoate at doses of 0.5, 1.0, 2.0, and 4.0 ml/kg daily for 90 days.<sup>62</sup> Lethality and effects on the following were reported after dosing: blood effects (at 2.0 ml/kg), kidneys, skin (slight dermatitis and other effects), reproduction, leukocyte (increased 2.5 x normal), testicular damage, and kidney damage. Animal deaths were not preceded by the usual pattern of systemic effects prior to death. Additional details were not included.

Five applications of benzyl benzoate (undiluted) were made to the backs of the following domestic animals: 1 sheep (250 ml dose), 1 pig (200 ml), 1 young cow (500 ml), and 1 horse (1000 ml). The animals were observed for 2 weeks. None of the animals died. In another study, 6 applications of undiluted benzyl benzoate were made to the backs (clipped skin, 4 " x 6" area) of 3 dogs as follows: 1 dog (200 ml volume) and 2 dogs (100 ml). The animals were observed for 2 weeks. No test substance-related effects were observed.<sup>63</sup>

Benzyl benzoate was rubbed onto cutaneous lesions on the shaved upper thighs of cats (no. not stated). Three applications per animal were made. Within 24 to 36 h, the test substance was lethal at doses of 8.7 to 12.8 g/kg.<sup>59</sup>

### **Repeated Dose Toxicity – Parenteral Study**

#### **Benzyl Benzoate**

Rabbits (1 or 2 animals) were injected s.c. once with benzyl benzoate in olive oil (doses of 1 to 2.5 ml/kg) or received 4 daily 0.25 mg/kg s.c. injections. Effects on leukocyte counts were monitored over an 11-day period and the animals were also examined for mortalities and clinical signs. The 2.5 ml/kg dose was lethal and clinical signs were observed. No effects were associated with the 0.25, 1, or 1.5 ml/kg doses.<sup>64</sup>

### **Ocular Irritation/Toxicity**

#### **Benzyl Alcohol**

The ocular toxicity of benzyl alcohol was evaluated using 9 New Zealand rabbits. The following test concentrations of benzyl alcohol (in stock solution of carboxymethylcellulose and polysorbate 80) were injected intravitreally, each into 3 eyes (15 eyes total injected): 0.0073%, 0.022%, 0.073%, 0.222%, and 0.733%. Control eyes (2) were injected intravitreally with 0.9% normal saline. Ocular toxicity was not observed at the lowest concentration; transient clinical changes (retinal hemorrhage and whitening), but no remarkable pathological changes, were noted in 1 of 3 eyes tested with 0.022% benzyl alcohol. Ocular toxicity was observed at the remaining concentrations; changes in the outer retina included loss and shortening of outer segments and receptors.<sup>65</sup>

Triamcinolone acetonide (TA) has been increasingly applied intravitreally in therapy for retinal diseases, and the vehicles of TA injections usually contain benzyl alcohol as a preservative. The ocular toxicity of 2 commercial vehicles (A and B) for drugs was evaluated using 3 groups of 12 New Zealand white rabbits (vehicle A, vehicle B, and balanced saline solution [BSS, negative control] groups). The 2 vehicles were similar in that each also contained benzyl alcohol (9.9 mg/ml), sodium carboxymethylcellulose (CBC, 7.5 mg/ml), and polysorbate 80 (0.4 mg/ml). Each group of 12 was divided into 2 groups, each receiving a different intravitreal dose (0.1 or 0.2 ml). Ocular examination (using ophthalmoscope) results were normal for group A and the negative control group; retinal damage was obvious in group B. However, at microscopic examination, retinal toxicity was observed in groups A and B. Vehicle B induced extensive retinal necrosis and atrophy, while vehicle A induced localized retinal changes.<sup>66</sup>

In a published comment on the preceding publication, the association of the benzyl alcohol preservative with retinal toxicity was acknowledged. However, it was noted that retinal toxicity is not observed at concentrations achieved following injection of 0.1 ml preserved TA in clinical practice.<sup>67</sup>

In a similar study, 4 groups of Chinchilla rabbits (pigmented) were injected intravitreally with the following 4 materials, respectively: 0.1 ml BSS, 0.1 ml TA (1.3 mg/ml), 0.1 ml vehicle (0.99% BA and other excipient) alone, and 0.1 ml TA + vehicle. Given the materials tested, it should be noted that commercially prepared TA was described as a suspension containing 40 mg TA plus vehicle containing 0.99% BA, 0.75% excipient sodium carboxymethylcellulose, and 0.04% polysorbate. At microscopic examination, morphologic changes in the ciliary body, lens, and retina were observed only in 2 groups (TA + vehicle and vehicle alone). Thus, it was concluded that benzyl alcohol was toxic to these tissues when injected intravitreally.<sup>68</sup>

### **Benzoic Acid, Benzyl Alcohol, Sodium Benzoate, and Potassium Benzoate**

According to the OECD SIDS initial assessment report on Benzoates, benzyl alcohol (4% aqueous) and benzoic acid (undiluted) were irritating to the eyes of rabbits and sodium benzoate (concentration not stated) was only slightly irritating. Data on potassium benzoate were not available; however, it was expected that this chemical would induce slight ocular irritation.<sup>2</sup>

### **Benzyl Benzoate**

Undiluted benzyl benzoate was instilled (0.1 ml) into the eyes of 6 rabbits (3 males, 3 females), and reactions were scored up to 72 h post-instillation. Ocular irritation was observed in 3 rabbits (only reactions reported) at 1 h post-instillation.<sup>69</sup>

In an ocular irritation study involving human subjects (test procedure/number of subjects not stated), undiluted benzyl benzoate was classified as an ocular irritant.<sup>3</sup>

### **Skin Irritation and Sensitization**

Skin irritation and sensitization data (human and non-human) are included in Table 5.

### **Benzyl Alcohol, Benzoic Acid, Sodium Benzoate, and Potassium Benzoate**

According to the OECD SIDS initial assessment report on Benzoates, benzoic acid and benzyl alcohol were slightly irritating to the skin, whereas, sodium benzoate was non-irritating. Skin irritation data on potassium benzoate were not available; however, it is not expected that this compound would be irritating to the skin.<sup>2</sup> In the same report, benzoic acid did not induce sensitization in animal studies; however, a low incidence of positive reactions was observed in dermatologic patients patch tested. Similar results were reported for sodium benzoate, and it has been suggested that the positive reactions observed were actually non-immunologic contact urticaria. Both positive and negative results were reported for benzyl alcohol in animal skin sensitization studies. A maximum skin sensitization incidence of 1% was reported for benzyl alcohol in human patch tests. Occupational exposure to benzyl alcohol, benzoic acid, or sodium benzoate has not resulted in skin sensitization over a period of decades.<sup>2</sup>

### **Case Reports**

Case reports involving patch, intradermal, or prick testing of benzyl alcohol, benzoic acid, sodium benzoate, or benzyl benzoate are included in Table 6.

### **Cytotoxicity**

#### **Sodium Benzoate**

The potential for sodium benzoate and other preservatives to induce lipid peroxidation in normal and  $\alpha$ -linolenic acid -loaded cultured hepatocytes from male Wistar rats was evaluated. Cell injury was measured by the release of lactic acid dehydrogenase into the culture medium after 10 h of incubation. At concentrations up to 10 mM, sodium benzoate was not cytotoxic to either group of hepatocytes. Also, lipid peroxidation was not detected in either group of hepatocytes incubated with sodium benzoate.<sup>70</sup>

#### **Benzyl Benzoate**

In an *in vitro* assay, benzyl benzoate was toxic to ascites sarcoma BP8 cell cultures. Additional details were not provided.<sup>3</sup>

## Phototoxicity

### Benzyl Benzoate

The phototoxicity of benzyl benzoate was evaluated using hairless mice (number and strain not stated). Details relating to the test protocol were not included. Following test substance application, test sites were irradiated for a total of 3 or 4, 24 h exposures. No reactions were observed after 3 exposures; however, a slightly positive reaction was observed after 4 exposures.<sup>3</sup>

The phototoxicity of benzyl benzoate was evaluated, using 5 female Hartley guinea pigs, at concentrations of 10%, 30%, and 50% in acetone. At 4 h after depilation, the test substance was applied to a circular area (diameter = 1.5 cm) on both sides of the back. Two applications (left and right side) of each concentration were made to the back of each animal. One side of each animal was covered with aluminum foil and the other was irradiated with 5 Toshiba model FL-40 BLB lamps (320 to 400 nm) at a distance of 10 cm for 70 min. Photo-irritation was not observed at any of the 3 test concentrations.<sup>71</sup>

## Photohemolysis

### Benzyl Alcohol, Sodium Benzoate, and Benzyl Benzoate

The photohemolytic activity of benzyl alcohol and sodium benzoate *in vitro* was evaluated in a test involving human erythrocyte suspensions, incubated with either chemical ( $10^{-3}$  mol/l) for 1 h and exposed to UVA or UVB radiation. The UVA rich light source was a UVASUN 5000 lamp (320 to 460 nm; 42 mW/cm<sup>2</sup>) and the UVB rich light source was a lamp with TL 20 W/12 light bulbs (between 275 and 365 nm; 1 mW/cm<sup>2</sup> [UVB] and 0.4 mW/cm<sup>2</sup> [UVA]) Cultures without either chemical served as negative controls. Neither benzyl alcohol nor sodium benzoate induced significant hemolysis.<sup>72</sup>

A similar *in vitro* photohemolysis test (human erythrocyte suspensions) was used to evaluate the phototoxicity of benzyl alcohol and benzyl benzoate. The radiation sources described in the preceding study were used. Moderate hemolysis (~6%) was induced by benzyl alcohol (at  $10^{-4}$  mol/l) only in the presence of UVA, classifying it as a weakly phototoxic compound. Benzyl benzoate (test concentration not stated) was not phototoxic in the presence of UVA or UVB light.<sup>73</sup>

The photohemolytic activity of benzyl benzoate (0.1%) *in vitro* was evaluated using human red blood cells. Cultures were exposed to UVA or UVB radiation for up to 3 h and then centrifuged. Release of hemoglobin in the supernatant was determined as cyanomethaemoglobin. No photohemolytic activity was noted for 0.1% benzyl benzoate.<sup>74</sup>

## REPRODUCTIVE AND DEVELOPMENTAL TOXICITY

### Benzoic Acid, Sodium Benzoate, Potassium Benzoate, and Benzyl Alcohol

According to the OECD SIDS initial assessment report on benzyl alcohol, benzoic acid, and its sodium and potassium salts, benzoic acid did not induce reproductive effects in a four-generation reproductive toxicity study (NOAEL > 750 mg/kg).<sup>2</sup> Groups of rats (20 rats/sex/group) received benzoic acid doses of 375 or 750 mg/kg/day in the diet continuously. Animals of the third generation were killed after 16 weeks. Also, test substance-related effects on reproductive organs (based on gross and microscopic examination) were not observed in subchronic studies (rats and mice) on benzyl alcohol and sodium benzoate.

In groups of rats fed sodium benzoate (doses up to 5600 mg/kg/day) during each day of gestation, developmental effects were observed only in the presence of marked maternal toxicity (reduced food intake and decreased body weight; NOAEL = 1400 mg/kg/day). All developmental effects were observed at doses  $\geq$  2800 mg/kg/day. On days 6 through 10, 15, or 18 of gestation, dosing with sodium benzoate by gavage in hamsters (NOEL = 300 mg/kg body weight), rabbits (NOEL = 250 mg/kg), and CD-1 mice (NOEL = 175 mg/kg), did not result in maternal toxicity; doses > NOEL were not tested. In mice of an unspecified strain dosed with benzyl alcohol by gavage, an NOAEL of 550 mg/kg (only dose) body weight for developmental toxicity was reported. An LOAEL of 750 mg/kg/day (only dose) for developmental toxicity was reported for CD-1 mice dosed orally (gavage) with benzyl alcohol. In this study, maternal toxicity (increased mortality, decreased body weight, and clinical toxicity) was observed.<sup>2</sup>

## Sodium Benzoate and Benzyl Benzoate

The effect of sodium benzoate on biochemical aspects of pregnant female albino rats and survival of their offspring was evaluated using groups of 10 Sprague-Dawley rats.<sup>75</sup> Two groups of weanling female rats were fed (*ad libitum*) sodium benzoate at doses of 0.01 g and 0.0125 g in the diet, respectively, daily for 12 weeks. Standard diet was fed to the control group. After mating, the animals were fed *ad libitum* during pregnancy. When compared to the control group, both doses of sodium benzoate induced a decrease in serum bilirubin and an increase in serum urea.

Both doses of sodium benzoate also induced an increase in serum uric acid ( $p < 0.01$  and  $p < 0.05$ , respectively). Serum alanine aminotransferase (ALT) activity was significantly greater ( $p < 0.01$ ) in the high dose group, but not in the lower dose group. The high dose did not induce a significant increase in serum creatinine. Statistically significant ( $p < 0.01$ ) decreases in food intake, hemoglobin, and hematocrit were also reported. Pregnant rats that received high doses of sodium benzoate sustained a 13.6% decrease (statistically significant,  $p < 0.05$ ) in mean weight of their pups, when compared to mean pup weight in the corresponding control group.<sup>75</sup>

The teratogenicity of benzyl benzoate was evaluated using 21 rats (strains not stated) per dose; doses administered in the diet were defined as 0.04 or 1.0% ( $\approx 24$  or 595 mg/kg body weight/day).<sup>3</sup> The test substance was administered daily from day 0 of gestation to day 21 post-parturition. The results of examinations for external, skeletal, or visceral anomalies indicated that benzyl benzoate did not induce harmful effects in fetuses.

In a reproductive and developmental toxicity study, pregnant rats (no. and strain not stated) were fed diets supplemented with 0.04 or 1.0% benzyl benzoate from day 0 of gestation to day 21 post-parturition.<sup>76</sup>

Another group of rats was fed a control diet. There were no effects on reproductive or developmental toxicity parameters; no external, skeletal, or visceral anomalies were observed in fetuses from either treatment group. Minor variations were observed. A significantly decreased number of fetuses with incomplete sternebrae was noted in the 1% benzyl benzoate treatment group.

Testicular atrophy was observed in 2 repeated dose dermal toxicity studies (90 days) on benzyl benzoate involving rabbits.<sup>3</sup> The exact doses at which this finding occurred were not stated. However, testicular atrophy was observed at daily doses  $> 0.5$  g/kg/day. These studies are summarized in the Repeated Dose Toxicity – Dermal Studies section earlier in the report text.

Pregnant NMRI mice (34 animals) were injected s.c. with an unspecified dose of benzyl benzoate in castor oil on days 1 and 11 of gestation.<sup>77</sup> The animals were killed on gestation day 17. Untreated mice served as controls. Fetal observations relating to the following were made: counts, sex, weight, and malformations. There was no evidence of test substance-related effects on fetuses. Additional details were not included.

A developmental toxicity study was performed using *Drosophila melanogaster*.<sup>78</sup> One group of males and females was raised on food medium with 4% benzyl benzoate in ethanol and the other group was raised on control feed medium. Virgin Canton-S males and females were collected daily for 4 days and then mated. The percentage of eggs hatched was 6.5 times greater in flies on feed containing benzyl benzoate, compared to those on control feed.

The safety of benzyl benzoate lotion (25% benzyl benzoate) as a topical treatment for scabies during pregnancy was assessed using a population of 444 pregnant women and their matched controls (1,776 pregnant women).<sup>79</sup> The study population consisted of refugee and migrant women attending antenatal clinics on the Thai-Burmese border between August of 1993 and April of 2006. Most first treatments took place during the second and third trimesters, and the overall median gestation exposure was 24.5 weeks. Treated women (444) received 559 applications of butyl benzoate lotion (79.5%, 15.5%, 4.5%, and 0.15% receiving 1, 2, 3, and 4 treatments, respectively).

Conditional Poisson regression was used to estimate risk ratios for outcomes of pregnancy (proportion of abortions, congenital abnormalities, neonatal deaths, stillbirths, and premature babies), mean birth weight, and estimated median gestational age for scabies and scabies-free women. Regarding pregnancy outcomes, there were no statistically significant differences between women treated with the lotion and their matched controls. Thus, there was no evidence of adverse effects on pregnancy outcome due to topical application of benzyl benzoate (25%) lotion.<sup>79</sup>

## Benzoic Acid, Benzyl benzoate - Estrogenic effects

The estrogenic activity of benzoic acid was evaluated in the recombinant yeast human estrogen ( $ER_{\alpha}$ ) receptor assay *in vitro* and in uterotrophic assays involving immature female Alpk:AP<sub>f</sub>SD rats (21 to 22 days old) and immature female Alpk:AP<sub>f</sub>CD-1 mice.<sup>80</sup> Immature mouse uterotrophic assays involved 3 daily s.c. injections of benzoic acid in corn oil (100

and 1,000 µg/kg doses; 5 ml/kg = dose volume), ending on day 4. The same protocol was used for the rat assays, with the exception that benzoic acid in arachis oil was injected (doses of 10, 100, and 1,000 µg/kg).

Benzoic acid ( $10^{-7}$  to  $10^{-3}$  M) was negative in the recombinant yeast human estrogen ( $ER_{\alpha}$ ) receptor assay *in vitro*. Overall (rat and mouse assays), benzoic acid produced one statistically significant and 3 statistically nonsignificant increases in average uterine weight and one significant and 6 nonsignificant decreases in average uterine weight. Therefore, in all uterotrophic assays, benzoic acid did not produce any reproducible evidence of estrogenic activity. Results for vehicle controls were negative and the positive control, estradiol, was uterotrophic.<sup>80</sup>

The estrogenic activity of benzyl benzoate (concentration range:  $10^{-9}$  to  $10^{-4}$  M) *in vitro* was evaluated in the E-screen test using MCF-7 breast cancer cells. Untreated cultures served as controls. In this test, MCF-7 cells proliferate in the presence of estrogen. Cell numbers were assessed by measurement of the total protein content, using the sulforhodamine B assay. Compared to control cultures, benzyl benzoate did not increase the proliferation of MCF-7 cells ( $p > 0.05$ ) over the range of concentrations tested.<sup>81</sup>

The estrogenic activity of benzyl benzoate, benzyl salicylate, and butylphenylmethylpropional (Lilial) in the estrogen-responsive MCF7 human breast cancer cell line was evaluated using the following assays: competitive binding assay to estrogen receptor (ER) of MCF7 cytosol, competitive binding assay to recombinant  $ER_{\alpha}$  and  $ER_{\beta}$ , and the assay of stably transfected ERE-CAT reporter gene in MCF7 cells.<sup>82</sup> In the latter assay, The ERE-CAT vector consisted of the estrogen response element (ERE) of the vitellogenin A2 gene from -331 to -295 bp cloned into the pBLCAT2 vector upstream of the thymidine kinase (tk) promoter. Cell proliferation experiments were also performed.

The following results indicate that all 3 chemicals produced estrogenic responses in cultured human breast cancer cells *in vitro*.<sup>82</sup> At 3,000,000-fold molar excess, each chemical was able to partially displace [<sup>3</sup>H]estradiol from recombinant human estrogen receptors  $ER_{\alpha}$  and  $ER_{\beta}$ , and from cytosolic ER of MCF7 cells.

At concentrations in the  $5 \times 10^{-5}$  to  $5 \times 10^{-4}$  M range, benzyl benzoate and the other 2 chemicals were able to increase the expression of a stably integrated estrogen-responsive reporter gene (ERE-CAT) and of the endogenous estrogen-responsive pS2 gene in MCF7 cells; however, these effects were at a lesser extent when compared to  $10^{-8}$  M 17 $\beta$ -estradiol (5,000 to 50,000 molar excess of benzyl benzoate).

In cell proliferation experiments, each chemical increased the proliferation of estrogen-dependent cells over a 7-day period. Cell proliferation was inhibited by fulvestrant (antiestrogen), suggesting an ER-mediated mechanism. However, over a 35-day period, the extent of proliferation in the presence of  $10^{-4}$  M benzyl benzoate, benzyl salicylate, or butylphenylmethylpropional increased to the same magnitude as that observed in the presence of  $10^{-8}$  M 17 $\beta$ -estradiol over a 14-day period (10,000 molar excess of benzyl benzoate).<sup>82</sup>

## GENOTOXICITY

### **Benzoic Acid, Sodium Benzoate, Potassium Benzoate, and Benzyl Alcohol**

According to the OECD SIDS initial assessment report on benzyl alcohol, benzoic acid, and its sodium and potassium salt, each chemical was not mutagenic in *in vitro* Ames tests. Various results (negative and positive [chromosomal/chromatid responses]) for sodium benzoate, potassium benzoate, and benzyl alcohol were obtained in other *in vitro* genotoxicity assays. However, while some mixed and/or equivocal *in vitro* chromosomal/chromatid responses have been observed, sodium benzoate and benzyl alcohol were not genotoxic in the *in vivo* cytogenetic assay, the micronucleus test, or in other *in vivo* assays. The weight of evidence of the *in vitro* and *in vivo* genotoxicity data indicates that these chemicals are not mutagenic or clastogenic.<sup>2</sup>

In the *in vitro* comet assay, the DNA-damaging potential of benzoic acid and benzyl alcohol was evaluated using human blood cells (lymphocytes).<sup>83</sup> This assay is able to detect single and double strand breaks in DNA. Benzoic acid was evaluated at concentrations ranging from 0.05 to 5 mM and, benzyl alcohol, at concentrations ranging from 1 to 50 mM. Benzoic acid was tested at lower concentrations ( $\leq 5$  mM) because this chemical has apoptotic effects at higher concentrations. The tail moment and % tail DNA in chemicals evaluated were compared to results for the solvent control (distilled water). Significantly increased tail moment and % tail DNA were noted only with 5 mM benzoic acid and with 25 and 50 M benzyl alcohol, indicating genotoxicity. A dose response was not observed.

Benzoic acid (in DMSO) was evaluated in the micronucleus test at concentrations ranging from 250 to 1,000 µg/ml (with and without metabolic activation) using L5178Y TK+/- clone 3.7.2C mouse lymphoma cells. Cultures without benzoic

acid served as negative controls and mitomycin C served as the positive control. Benzoic acid was non-genotoxic over the range of concentrations tested, both with and without metabolic activation. The positive control was genotoxic.<sup>84</sup>

The genotoxicity of sodium benzoate was evaluated in the sister chromatid exchanges assay (SCE assay) using human blood lymphocyte cultures incubated for 72 h. Preparations were scored blindly for cells in their first mitosis, second mitosis, and third and subsequent divisions. Forty SCEs second division cells from each culture were scored for SCEs. Compared to negative control cultures, sodium benzoate (0.02, 0.2, 2, 4, and 8 mM) caused statistically significant delays in cell division ( $p < 0.01$ ), indicative of weak cytostatic activity, but did not induce cytotoxicity. Also, when compared to negative control cultures, 8 mM sodium benzoate induced a statistically significant increase ( $p < 0.01$ ) in SCEs/cell. Sodium benzoate (2 mM) induced a statistically significant decrease ( $p < 0.05$ ) in SCEs/cell when compared to the 8 mM sodium benzoate culture.<sup>85</sup>

The DNA-damaging activity of sodium benzoate and potassium benzoate was evaluated in the spore rec-assay using *Bacillus subtilis* M45 (rec<sup>-</sup>) and H17 (rec<sup>+</sup>) strains with and without metabolic activation. Sodium benzoate (in water) was tested at doses of 16 and 20 mg/disk and potassium benzoate (in mixed solution of water and ethanol [1:1]) was tested at doses of 15 and 20 mg/disk. Results for both chemicals were judged positive in this assay.<sup>86</sup>

### **Benzyl Benzoate**

In the Ames test, benzyl benzoate was not mutagenic to *Salmonella typhimurium* strains TA98 or TA 100 at doses up to 5,000 µg/plate (plate incorporation assay) or 5,000 mg/plate (pre-incubation assay). Benzyl Benzoate also was not mutagenic to the following strains at doses up to 3 µmol/plate (pre-incubation assay) with or without metabolic activation: TA98, TA100, TA1525, and TA 1537. In the recombination assay, benzyl benzoate was not mutagenic to *Bacillus subtilis* strains H17 or M45 at a concentration of 10 mg/disk.<sup>3</sup>

In another Ames test, the mutagenicity of benzyl benzoate (in ethanol) was evaluated using *Salmonella typhimurium* strains TA98, TA100, TA1535, and TA1537. Benzyl benzoate was tested at a concentration of 3 µmol/plate with and without metabolic activation and was not found to be mutagenic.<sup>87</sup>

## **Photogenotoxicity**

### **Benzoic Acid and Sodium Benzoate**

The photogenotoxicity of benzoic acid (0.5%) and sodium benzoate (0.5%) and other food additives was evaluated using *Escherichia coli* cell suspensions.<sup>88</sup> Initially, the cytotoxic and genotoxic potential of each in the absence of sunlight (200 to 3,000 nm; mean intensity =  $5.8 \times 10^2 \text{ Wm}^{-2}$ ) was evaluated by allowing the cells to remain in contact with the additive for 80 min in the dark. Neither benzoic acid nor sodium benzoate affected cell viability (i.e., not cytotoxic) or the number of spontaneous mutations in the absence of sunlight. The number of spontaneous mutations was 2.09 mutations per  $10^8$  cells. Also, these chemicals did not induce mutations in the absence of sunlight.

Plates containing the cellular suspension (5 ml) and a food additive (0.5% concentration) were exposed to sunlight for periods ranging from 0 min to 60 min. Exposure to sunlight resulted in cell death in the presence or absence of benzoic acid or sodium benzoate. When cells were exposed to direct sunlight in the absence of benzoic acid or sodium benzoate, the number of induced mutations increased with duration of exposure ( $5.1 \times 10^2$  mutations per  $10^8$  cells after 60 min). In the presence of benzoic acid or sodium benzoate, the number of induced mutations increased with the duration of exposure to sunlight. This increase was almost additive over that observed with sunlight exposure alone. After 60 min, there were approximately  $1 \times 10^3$  mutations per  $10^8$  cells.<sup>88</sup>

## **Effect on DNA Synthesis**

### **Sodium Benzoate**

Aqueous sodium benzoate, at a concentration of 7.20 mg/ml, was added to *Tetrahymena pyriformis* in an experimental series consisting of six 100 ml cultures. Another experimental series, without sodium benzoate, served as the control. The quantitative analysis of DNA content of the protozoan nuclei was performed using an image analysis system. Sodium benzoate caused a statistically significant increase ( $p < 0.01$ ) in DNA content, suggestive of stimulation of mitosis. The authors speculated that this observation might represent a direct cytotoxic effect due to increased DNA content of nuclei.<sup>89</sup>

## CARCINOGENICITY

The following carcinogenicity study summaries are included in the CIR final report on benzyl alcohol, benzoic acid, and sodium benzoate that was published in 2001.<sup>1</sup>

### **Benzyl Alcohol and Benzoic Acid**

Groups of 100 F344/N rats (50 each sex) were dosed orally with 200 or 400 mg/kg benzyl alcohol in corn oil, 5 days per week for 103 weeks. Groups of 100 B6C3F<sub>1</sub> mice were dosed with 100 or 200 mg/kg benzyl alcohol following the same schedule. During week 80, mice were mistakenly dosed for four days with 375 (low-dose group) and 750 mg/kg (high-dose group) of  $\alpha$ -methylbenzyl alcohol. Dose-related negative trends were noted in the incidences of anterior pituitary gland neoplasms in female rats (vehicle control, 29/50; low dose, 17/47; high dose, 9/49) and of Harderian gland adenomas in male mice (8/50; 3/50; 2/50). Epithelial hyperplasia of the nonglandular stomach was noted in 4 of 50 high-dose male rats; it was not found in controls or low-dose male rats. An increased incidence of adenomas of the adrenal cortex noted in high-dose male mice (0/48; 0/44; 3/48) was within historical range and not considered compound-related. The NTP investigators considered the study negative for benzyl alcohol-induced carcinogenicity. However, reviewing the study, the EPA considered the 3 of 48 incidence of adrenal cortex adenoma to be “equivocal evidence of carcinogenic activity rather than negative.”

From the Final Report on benzyl alcohol, benzoic acid, and sodium benzoate<sup>1</sup>

A skin painting study was performed using groups of 120 Eppley Swiss mice (60 per sex). A non-oxidative hair dye containing 2.0% benzyl alcohol and 0.016% benzoic acid was painted onto the skin at a dose of 0.05 ml/application, three times weekly for 20 months. Sites were shaved of hair 24 h before each application and a new bottle of dye was used each week. Two groups of control animals were shaved but not treated. Nine months into the study, 10 mice/sex/group were killed. Body weights and survival differed little between treatment and control groups. Varying degrees of chronic dermal inflammation were noted in all groups, including the controls. A significant ( $p < .01$ ) increase in malignant lymphomas was noted in treated females (23/60). However, the researchers noted that one concurrent control group had a very low incidence (7/60 or 12%) of that tumor type. The rate was 22% for the other control group and had averaged 33% for three control groups in previous studies. Thus, the findings were not considered treatment-related. The incidence of pulmonary adenomas and hepatic hemangiomas, which are common in this mouse strain, were similar between treated and control groups. No unusual neoplasms were observed.

From the Final Report on benzyl alcohol, benzoic acid, and sodium benzoate<sup>1</sup>

### **Sodium Benzoate**

For 18 to 24 months, groups of Fischer 344 rats (50 males and 52 females per group) received feed containing 2% or 1% sodium benzoate. The doses corresponded to the maximum tolerated dose (MTD) and  $\frac{1}{2}$ MTD as determined in 6-week toxicity studies. A control group of 25 male and 43 female rats received untreated feed. Average daily sodium benzoate intake was 280 and 202 mg, respectively, for male and female rats of the 2% group, and 141 and 102 mg, respectively, for male and female rats of the 1% group. No clinical signs of toxicity or differences in average body weight or mortality rates were noted in treated rats when compared with controls. Neoplasms that were present in treated rats were similar in type and number to those in controls. No evidence of sodium benzoate-related carcinogenicity was observed.

From the Final Report on benzyl alcohol, benzoic acid, and sodium benzoate<sup>1</sup>

In a life-time drinking water study, 100 Albino Swiss mice (50 of each sex) were supplied with water containing 2% sodium benzoate. A control group of 200 mice was supplied with untreated water. Average daily intake of sodium benzoate was 124.0 and 119.2 mg for males and females, respectively. Sodium benzoate treatment did not affect survival. No carcinogenic effect attributable to treatment was noted at necropsy.

From the Final Report on benzyl alcohol, benzoic acid, and sodium benzoate<sup>1</sup>

## SUMMARY

The following cosmetic ingredients are reviewed in this safety assessment: Benzyl alcohol, benzoic acid, sodium benzoate, calcium benzoate, magnesium benzoate, potassium benzoate, and benzyl benzoate. Most of these ingredients function as fragrance ingredients/preservatives in cosmetic products. Together, data reported to the Food and Drug Administration's Voluntary Cosmetic Registration Program in 2010 and the results of a 2010 Personal Care Products industry survey indicated use of the following ingredients in cosmetics: benzyl alcohol, benzoic acid, sodium benzoate, calcium

benzoate, potassium benzoate, and benzyl benzoate. According to this industry survey, ingredient use concentrations have ranged from 0.000001% (sodium benzoate) to 10% (benzyl alcohol).

Among the methods of manufacture identified are the production of benzyl alcohol via the action of sodium or potassium carbonate on benzyl chloride and the production of benzoic acid via the decarboxylation of phthalic anhydride. Hydroxyl radical generated by the metal-catalyzed reduction of  $O_2$  and  $H_2O_2$  by ascorbic acid can attack benzoic acid to produce benzene under conditions that are prevalent in beverages. Additionally, exposure to UV light and elevated temperature over the shelf life of beverages may result in benzene formation in products containing benzoic and ascorbic acids. The U.S. beverage industry voluntarily reformulated beverages that were found to contain benzene levels at or above the maximum contaminant level for drinking water established by the U.S. Environmental Protection Agency. Limits for heavy metal impurities in food-grade sodium benzoate (as Pb,  $\leq 2$  mg/kg) and pharmaceutical-grade potassium benzoate (heavy metals,  $\leq 0.001\%$ ) have been established.

Benzyl alcohol is metabolized to benzoic acid via simple oxidation. Benzoic acid and sodium benzoate are rapidly absorbed from the gastrointestinal tract of mammals, conjugated with glycine in the liver, and then excreted as hippuric acid. In *in vitro* studies, the percutaneous absorption of benzoic acid through guinea pig skin and the percutaneous absorption of benzyl alcohol and benzyl benzoate through human skin was reported. The percutaneous absorption and urinary excretion of benzyl alcohol and benzyl benzoate was observed following dermal application to rhesus monkeys.

Benzoic acid (up to 1,000  $\mu\text{g}/\text{kg}$ , s.c. injection) did not induce any reproducible evidence of estrogenic activity in uterotrophic assays involving immature female Alpk:AP<sub>r</sub>SD rats and female Alpk:AP<sub>r</sub>CD-1 mice. However, benzyl benzoate induced estrogenic responses in cultured human breast cancer cells *in vitro*. At 3,000,000-fold molar excess, benzyl benzoate was able to partially displace [<sup>3</sup>H]estradiol from recombinant human estrogen receptors ER $\alpha$  and ER $\beta$ , and from cytosolic ER of MCF7 cells. The antiparasitic activity of benzyl benzoate has been demonstrated *in vitro*.

Neither acute (up to 12 mg/L aerosol/dust) nor repeated inhalation (up to 1,000 mg/m<sup>3</sup>) exposures to benzyl alcohol or benzoic acid caused death in rats. Results of the repeated dose inhalation toxicity study also indicated no test substance-related macroscopic or microscopic findings for either test substance. Overall, the results of acute oral and dermal toxicity studies indicated a low level of toxicity for benzyl alcohol, benzoic acid, and benzyl benzoate. The same was true for repeated dose oral toxicity studies on benzoic acid and its salts and benzyl alcohol. Systemic/dermal effects induced by benzyl benzoate were observed in rats receiving repeated dermal doses up to 2.0 g/kg and in rabbits receiving repeated dermal doses up to 4.0 ml/kg. However, test substance-related toxic signs were not observed in other species that received repeated doses of benzyl benzoate.

Benzyl alcohol (4% aqueous), benzoic acid (undiluted), and its sodium salt (concentration not stated) were irritating to the eyes of rabbits, and it was expected that potassium benzoate would be irritating as well. Benzyl benzoate (undiluted) was irritating to the eyes of rabbits and humans. Undiluted benzyl alcohol and benzoic acid were slightly irritating and benzyl benzoate was non-irritating to the skin at concentrations up to 50% in animal studies. Undiluted benzyl benzoate was a non-irritant in human subjects. In animal studies, benzyl benzoate induced sensitization reactions ranging from none to moderate. Moderate sensitization was observed at a concentration of 10% benzyl benzoate, but not at lower concentrations, and there was one report of mild sensitization at a concentration of 40%. Benzyl benzoate was classified as non-sensitizer when tested at a concentration of 30% in healthy human subjects. Most of the skin sensitization studies on benzyl benzoate involved groups of patients, and mixed results regarding sensitization potential were reported.

A slightly positive reaction was observed in a phototoxicity study involving hairless mice tested with an unspecified concentration of benzyl benzoate (4, 24 h exposures). Photo-irritation was not observed in guinea pigs tested with benzyl benzoate at concentrations up to 30%. Moderate photohemolytic activity in human erythrocytes was observed in the presence of benzyl benzoate, but significant photohemolytic activity was not associated with sodium benzoate or benzyl alcohol in these cells.

Developmental effects only were observed in the presence of marked maternal toxicity in rats fed sodium benzoate in the diet ( $\geq 2800$  mg/kg/day; NOAEL = 1400 mg/kg/day). Testicular atrophy was observed in rabbits that received repeated dermal doses of benzyl benzoate ( $> 0.5$  g/kg/day). However, overall, benzyl alcohol, benzoic acid, and sodium benzoate were not classified as reproductive/developmental toxicants in oral and dermal animal studies. There was no evidence of adverse effects on pregnancy outcome due to topical application of benzyl benzoate (25%) lotion in pregnant women.

Mixed/equivocal results were apparent in *in vitro* assays, but negative results were reported in *in vivo* genotoxicity assays on benzyl alcohol, benzyl benzoate, benzoic acid, and its sodium and potassium salt. The photogenotoxicity of



benzoic acid (0.5%) and sodium benzoate (0.5%) has been demonstrated in vitro using bacterial cell suspensions. Benzyl alcohol was negative for carcinogenicity when dermally tested on mice at 2.00% in a nonoxidative hair dye. NTP considered it negative for carcinogenicity following 2 years of oral dosing in rats (up to 400 mg/kg) and mice (up to 200 mg/kg), but the EPA considered the 3 of 48 incidence of adrenal cortex adenoma (male mice) to be equivocal evidence of carcinogenic activity rather than negative. Following dermal application to mice, a nonoxidative hair dye containing 2.0% benzyl alcohol and 0.016% benzoic acid was negative for carcinogenicity. Sodium benzoate was negative for carcinogenicity when administered orally at up to 2% to rats (in feed for up to 2 years) or mice (in a life-time drinking water study).

## **DISCUSSION**

The CIR Expert Panel noted gaps in the available safety data for some of the ingredients in this safety assessment. The available data on many of the ingredients are sufficient, however, structure activity relationships and metabolism suggest that the available data can be extrapolated to support the safety of the entire group.

The negative inhalation toxicity data provided satisfied the Panel's request for data and supported the safe use of these ingredients in cosmetic products in which inhalation is a primary route of exposure.

The Panel did note the formation of adenomas in the adrenal cortex of male B6C3F<sub>1</sub> mice receiving a high dose of benzyl alcohol in an oral carcinogenicity study. The low rate of occurrence was within historical control values for this benign tumor. The Panel acknowledged that EPA reviewed the results of this study and concluded that the results suggest equivocal evidence of carcinogenicity, but it was the Panel's view that there was no evidence of carcinogenicity in the animals.

Data from model studies, pertaining to the beverage industry, reviewed by the Panel raised the possibility that benzene could be formed from benzoic acid and benzoates in cosmetics. The yields of benzene, however, that could be generated by photodegradation are sufficiently low that they do not constitute a relevant hazard.

In its previous safety assessment of benzyl alcohol, benzoic acid, and sodium benzoate, the Panel established a 5% safe concentration limit for benzyl alcohol, benzoic acid, and sodium benzoate and a 10% safe concentration limit for benzyl alcohol in hair dyes. On review of the current practices of use and concentration, it was clear that industry has adhered to those limits. Accordingly, reference to present practices of use and concentration is sufficient to assure safety.

## **CONCLUSION**

The CIR Expert Panel concluded that the following cosmetic ingredients are safe in the present practices of use and concentration described in this safety assessment:

- benzyl alcohol
- benzoic acid
- sodium benzoate
- calcium benzoate
- magnesium benzoate\*
- potassium benzoate
- benzyl benzoate

\*Were ingredients in this group not in current use to be used in the future, the expectation is that they would be used in product categories and at concentrations comparable to others in the group.

**Table 1.** Benzyl Alcohol and Benzoic Acid and its Salts and Ester<sup>22</sup>

Chemical Names/CAS Nos.	Definitions*	Functions in Cosmetics
<b>Benzyl Alcohol</b> Alcohol benzylicus; benzenemethanol; benzylic alcohol; phenylcarbinol; phenylmethanol; phenylmethyl alcohol; and $\alpha$ -toluenol; CAS No. 100- 51-6	An aromatic alcohol	External analgesics; fragrance ingredients; oral health care drugs; preservatives; solvents; and viscosity decreasing agents
<b>Benzoic Acid</b> Acidum benzoicum; benzenecarboxylic acid; benzeneformic acid; benzenemethanoic acid; carboxybenzene; dracyclic acid; phenylcarboxylic acid; and phenylformic acid; CAS No. 65-85-0	An aromatic acid	Fragrance ingredients; preservatives; and pH adjusters
<b>Sodium Benzoate</b> Benzoic acid, sodium salt and natrii benzoas (EP); CAS No. 532-32-1	The sodium salt of benzoic acid	Corrosion inhibitors; fragrance ingredients; and preservatives
<b>Calcium Benzoate</b> Benzoic Acid, Calcium Salt and Calcium Dibenzoate; CAS No. 2090- 05-3	The calcium salt of benzoic acid	Preservatives
<b>Magnesium Benzoate</b> Benzoic Acid, magnesium salt and magnesium dibenzoate; CAS No. 553- 70-8	The magnesium salt of benzoic acid	Preservatives
<b>Potassium Benzoate</b> Benzoic Acid, Potassium Salt; CAS No. 582-25-2	The potassium salt of benzoic acid	Preservatives
<b>Benzyl Benzoate</b> Benylate; benzoic acid, benzyl ester; benzoic acid, phenylmethyl ester; benzylis benzoas (EP); and phenylmethyl benzoate; CAS No. 120- 51-4	The ester of benzyl alcohol and benzoic	Fragrance ingredients; pesticides; and solvents

\*See Figure 1 for structures, completing the definition of each chemical.

**Table 2.** Properties of Benzyl Alcohol and Benzoic Acid and its Salts and Ester<sup>90</sup>

Properties	Benzyl Alcohol	Benzoic Acid	Sodium Benzoate	Calcium Benzoate	Magnesium Benzoate	Potassium Benzoate	Benzyl Benzoate
Form	Liquid	Monoclinic plates	White granules		White powder	White solid <sup>2</sup>	Leaflets/oily liquid
Molecular weight	108.14	122.12	144.1	121.12 <sup>91</sup>	266.53	160.21	212.24
Density	1.045	1.321					1.118
Melting point	-15.19°C	122.4°C	330.6 <sup>0</sup> C <sup>2</sup>		200 <sup>0</sup>	330.6 <sup>0</sup> C <sup>2</sup>	21°C
Boiling point	204.7°C	249.2 <sup>0</sup> C	464.9 <sup>0</sup> C <sup>2</sup>			464.9 <sup>0</sup> C <sup>2</sup>	323-324 <sup>0</sup> C
Water Solubility	dissolves (1g) in ~ 25 ml water	2.9 g/l	1 g dissolves in 1.8 ml water		Soluble in 20 parts water	556 g/l <sup>2</sup>	Practically insoluble in water
logP	1.1 <sup>2</sup>	1.8 <sup>2</sup>	-2.269 <sup>2</sup>		1.53 <sup>92</sup>	-2.269 <sup>2</sup>	3.68 <sup>93</sup>
UV absorption max.					228.6 nm <sup>91</sup>	226.8 nm <sup>91</sup>	256 nm <sup>91</sup>

**Table 3.** Current Frequency and Concentration of Use According to Duration and Type of Exposure Provided in 2011<sup>23,24</sup>

	<b>Benzoic Acid</b>		<b>Benzyl Alcohol</b>		<b>Sodium Benzoate</b>		<b>Calcium Benzoate</b>	
	# of Uses	Conc. (%)	# of Uses	Conc. (%)	# of Uses	Conc. (%)	# of Uses	Conc. (%)
<b>Exposure Type</b>								
<i>Eye Area</i>	23	0.00005 to 0.2	109	0.000008 to 0.8	58	0.00001 to 0.4	NR	NR
<i>Incidental Ingestion</i>	64	0.003 to 2	23	0.000006 to 0.9	33	0.001 to 0.6	NR	NR
<i>Incidental Inhalation-sprays</i>	16	0.000003 to 0.5	496	0.0002 to 3	43	0.001 to 0.5	NR	0.004
<i>Incidental Inhalation-powders</i>	4	0.000002 to 0.001	28	0.0001 to 0.3	4	0.3	NR	0.002 to 0.004
<i>Dermal Contact</i>	503	0.000002 to 5	1673	0.000008 to 3	1075	0.000001 to 1	NR	NR
<i>Deodorant (underarm)</i>	1	0.03 to 0.5	38	0.03 to 0.5	2	0.5		
<i>Hair - Non-Coloring</i>	97	0.00002 to 1	361	0.0002 to 3	431	0.000005 to 1	NR	NR
<i>Hair-Coloring</i>	22	0.004 to 0.03	56	0.00007 to 10	18	0.008 to 0.5	NR	NR
<i>Nail</i>	7	0.0005 to 0.2	3	0.00002 to 0.5	7	0.00001 to 0.001	NR	NR
<i>Mucous Membrane</i>	201	0.000003 to 2	269	0.000006 to 1	407	0.0004 to 1	NR	NR
<i>Baby Products</i>	3	0.000002 to 0.08	25	0.004 to 0.9	26	0.3 to 1	NR	NR
<b>Duration of Use</b>								
<i>Leave-On</i>	378	0.000002 to 5	1491	0.000008 to 3	619	0.000001 to 1	NR	0.002 to 0.004
<i>Rinse off</i>	295	0.000003 to 5	651	0.000006 to 10	936	0.000001 to 1	NR	NR
<i>Diluted for (bath) use</i>	23	0.000003 to 0.08	20	0.001 to 0.9	15	0.004 to 1	NR	NR
<b>Totals/Conc. Range</b>	696	0.000002 to 5	2162	0.000006 to 10	1570	0.000001 to 1	NR	0.002 to 0.004
	<b>Potassium Benzoate</b>		<b>Benzyl Benzoate</b>		<b>Magnesium Benzoate</b>			
	# of Uses	Conc. (%)	# of Uses	Conc. (%)	# of Uses	Conc. (%)		
<b>Exposure Type</b>								
<i>Eye Area</i>	NR	NR	5	0.0006 to 0.002	NR	NR		
<i>Incidental Ingestion</i>	NR	NR	82	0.000005 to 0.2	NR	NR		
<i>Incidental Inhalation-sprays</i>	NR	NR	583	0.002 to 4	NR	NR		
<i>Incidental Inhalation-powders</i>	NR	NR	30	0.001 to 0.3	NR	NR		
<i>Dermal Contact</i>	NR	0.002	1321	0.0001 to 4	NR	NR		
<i>Deodorant (underarm)</i>	NR	NR	47	NR	NR	NR		
<i>Hair - Non-Coloring</i>	1	NR	259	0.006 to 4	NR	NR		
<i>Hair-Coloring</i>	NR	NR	21	0.07 to 2	NR	NR		
<i>Nail</i>	1	0.0003	26	0.2 to 0.8	NR	NR		
<i>Mucous Membrane</i>	NR	0.0002	227	0.000005 to 2	NR	NR		
<i>Baby products</i>	NR	NR	3	NR				
<b>Duration of Use</b>								
<i>Leave-On</i>	2	0.0003	1396	0.000005 to 4	NR	NR		
<i>Rinse off</i>	NR	0.0002	282	0.0001 to 2	NR	NR		
<i>Diluted for (bath) use</i>			31	0.0005 to 2	NR	NR		
<b>Totals/Conc. Range</b>	2	0.0002 to 0.0003	1709	0.000005 to 4	NR	NR		

NR = Not Reported; Totals = Rinse-off + Leave-on Product Uses.

Note: Because each ingredient may be used in cosmetics with multiple exposure types, the sum of all exposure type uses may not equal the sum total uses.

**Table 4.** Acute Oral Toxicity Studies on Potassium and Benzyl Benzoate

Material	Animals	Test Procedure	Results
Potassium Benzoate	Rats, mice, and guinea pigs	Not stated	LD50 > 10,000 mg/kg <sup>2</sup>
Benzyl Benzoate (undiluted)	10 rats/dose	Oral dosing followed by 6-day observation period; no GLP*	LD50 = 1891 mg/kg <sup>3</sup>
Benzyl Benzoate	20 rats	Single dose by gavage; 14-day observation period; no GLP	LD50 = 2800 mg/kg <sup>3</sup>
Benzyl Benzoate	Rats (no. not stated)	Not stated	LD50 = 500 mg/kg <sup>3</sup>
Benzyl Benzoate (in arachis oil)	Rats (no. not stated)	OECD guideline 401; 100 to 2000 mg/kg oral doses; GLP study	LD50 > 2000 mg/kg <sup>3</sup>
Benzyl Benzoate (in arachis oil)	10 Sprague-Dawley albino rats	Single 5.0 ml/kg dose; 14-day observation period	LD50 > 2000 mg/kg <sup>94</sup>
Benzyl Benzoate	Rats (no. not stated)	Not stated	LD50 = 1700 mg/kg <sup>95</sup>
Benzyl Benzoate (1% in gum tragacanth suspension)	5 groups of 10 Sprague-Dawley rats	Doses up to 4000 mg/kg total (i.e., four 250 mg/kg doses at 40 ml/kg dose volume/dose); 14-day observation period	LD50 = 1.16 g/kg <sup>96</sup>
Benzyl Benzoate (in Tween 80 + gum tragacanth vehicle)	6 groups of 10 Sprague-Dawley rats 5 males, 5 females/group)	6 different dose levels (dose volume = 40 ml/kg)	LD50 = 1160 mg/kg <sup>97</sup>
Benzyl Benzoate (in tween 80 solution + methylcellulose vehicle)	Rats (25 males, 25 females)	Not stated	LD50 = 1550 mg/kg <sup>98</sup>
Benzyl Benzoate (in tween 80 solution + gum tragacanth vehicle)	Sprague-Dawley rats (5 males, 5 females)	10 mg/kg total dose (i.e., four 2500 mg/kg doses); 14-day observation period	LD50 > 10 g/kg <sup>99</sup>
Benzyl Benzoate	10 rats/dose	Dosing followed by 6-day observation period	LD50 = 1.7 ml/kg <sup>100</sup>
Benzyl Benzoate	20 rats	Dosing followed by 2-week observation period	LD50 = 2.8 g/kg <sup>63</sup>
Benzyl Benzoate (undiluted)	10 mice/dose	Dosing followed by 6-day observation period; no GLP	LD50 = 1557 mg/kg <sup>3</sup>
Benzyl Benzoate	10 mice/dose	Dosing followed by 6-day observation period	LD50 = 1.4 ml/kg <sup>100</sup>
Benzyl Benzoate	Mice (no. not stated)	Not stated	LD50 = 1400 mg/kg <sup>3</sup>
Benzyl Benzoate (undiluted)	10 rabbits/dose	Dosing followed by 6-day observation period; no GLP	LD50 = 2002 mg/kg <sup>3</sup>
Benzyl Benzoate	10 rabbits/dose	Dosing followed by 6-day observation period	LD50 = 1.8 ml/kg <sup>100</sup>
Benzyl Benzoate	9 rabbits	Single dose by gavage; 14-day observation period; no GLP	LD50 = 1680 mg/kg <sup>3</sup>
Benzyl Benzoate	12 rabbits	Dosing followed by 2-week observation period	LD50 = 1.68 g/kg <sup>63</sup>
Benzyl Benzoate	11 cats	Single dose by gavage; 14-day observation period; no GLP	LD50 = 2240 mg/kg <sup>3</sup>
Benzyl Benzoate	4 dogs	Single dose by gavage; 14-day observation period; no GLP	LD50 > 2244 mg/kg <sup>3</sup>
Benzyl Benzoate (undiluted)	10 guinea pigs/dose	Oral dosing followed by 6-day observation period ; no GLP	LD50 = 1112 mg/kg <sup>3</sup>
Benzyl Benzoate	Guinea pigs (no. not stated)	Not stated	LD50 = 1000 mg/kg <sup>3</sup>
Benzyl Benzoate	10 guinea pigs/dose	Dosing followed by 6-day observation period	LD50 = 1 mg/kg <sup>100</sup>

\*GLP = good laboratory practice

**Table 5. Skin Irritation and Sensitization Studies**

Test Substance	Animals/Subjects	Doses/Concentrations Tested	Procedure	Results
<b>Animal Studies</b>				
Benzyl alcohol (purity not stated)	Rabbits (no. not stated)	undiluted benzyl alcohol (10 mg)	dose applied for 24 h	Slightly irritating <sup>2</sup>
Benzoic Acid (purity not stated)	3 albino rabbits	undiluted benzoic acid (0.5 g/kg)	dose applied to flank under semi-occlusive dressing for 4 h	Slightly irritating <sup>2</sup>
Sodium Benzoate	Rabbits (no. not stated)	undiluted benzyl alcohol (10 mg)	dose applied for 24 h	Slightly irritating <sup>2</sup>
Sodium Benzoate	Rabbits (no. not stated)	Concentration not stated	OECD Guide - line 404 protocol	Non-irritating <sup>2</sup>
Benzyl Benzoate	Guinea pigs (no. not stated)	undiluted benzyl benzoate (15 g/kg)	Dose applied to intact abdominal skin (duration not stated)	Non-irritating <sup>59</sup>
Benzyl Benzoate (50%)	Guinea pigs (no. not stated)	up to 50%	Not stated	Non-irritant <sup>71</sup>
Benzyl Benzoate (40%)	10 female Hartley albino guinea pigs	10% induction concentration; challenge concentrations up to 40%	10% w/v in complete adjuvant (intradermal induction); 10% w/v in Freund's complete adjuvant + saline (topical induction); topical challenge concentrations up to 40% benzyl benzoate in propylene glycol + acetone	No reactions at challenge concentrations of 5% and 10% benzyl benzoate; mild reactions at 40% challenge concentration (4 of 10 animals)
Benzyl Benzoate (25% in acetone/PEG vehicle)	4 male guinea pigs	up to 25%	24-h occlusive patch test	Non-irritant <sup>101</sup>
Benzyl Benzoate (10%)	Guinea pigs (no. not stated)	10%	Maximization test	Moderate sensitization <sup>102</sup>
Benzyl Benzoate (10% in ethanol)	4 female guinea pigs	up to 10%	24-h occlusive patch test; up to 10% during induction and challenge with 2.5%	Non-irritating at 5 and 10% concentrations. Barely perceptible erythema (at 24 and 48 h) at 2.5% concentration <sup>103</sup>
Benzyl Benzoate (10% in acetone)	10 guinea pigs	10%	Modified Freund's complete adjuvant test	Weak sensitization <sup>104</sup>
Benzyl Benzoate (1 and 10%)	Guinea pigs (no. not stated)	1% and 10% (15 or 30 mg)	Modified Freund's complete adjuvant test: Challenge with 1% and 10% benzyl benzoate (in acetone)	Weak sensitization at both concentrations <sup>105</sup>
Benzyl Benzoate	Guinea pigs (no. not stated)	Not stated	Maximization test (no GLP)	Non-sensitizer <sup>3</sup>
Benzyl Benzoate	Guinea pigs (6 to 8 per group)	Not stated	Open epicutaneous test (24 h application)	Non-sensitizer <sup>3</sup>
Benzyl Benzoate	4 female New Zealand albino rabbits	Undiluted benzyl benzoate (0.5 ml/2.5 cm <sup>2</sup> )	4 h semiocclusive patch test	Non-irritant <sup>106</sup>
Benzyl Benzoate	Rabbits (no. not stated)	Not stated	3-week subacute skin toxicity test	Very mild gross skin irritation <sup>100</sup>
<b>Predictive Tests</b>				
Benzyl Alcohol	31 healthy subjects	0.2 ml benzyl alcohol	Hill Top chamber with Webril pad applied to upper arm for up to 4 h	Skin irritation in 1 subject <sup>107</sup>
Benzyl Alcohol lotion 5% (Ulesfia TM)	485 subjects	Lotion tested as supplied	Topical treatment of head lice program	Low frequency of application site irritation (2.3% incidence) <sup>42</sup>

**Table 5. Skin Irritation and Sensitization Studies**

Test Substance	Animals/Subjects	Doses/Concentrations Tested	Procedure	Results
Benzoic Acid	86 healthy subjects (mean age = 37.3 years)	1 M in petrolatum	Finn chambers applied to volar arm for 20 min	Caused contact urticaria <sup>108</sup>
Benzoic Acid	58 healthy subjects (mean age = 39 years)	1 M in petrolatum (50 µl volume)	Finn chambers applied to volar arm for 20 min	Caused contact urticaria <sup>109</sup>
Benzyl Benzoate	18 subjects (18+ years)	5 to 10 drops of undiluted benzyl benzoate	Closed patch containing benzyl benzoate applied to healthy and locally inflamed skin for up to 48 h	No skin irritation <sup>59</sup>
Benzyl Benzoate	Male and female subjects (no. not stated; 18+ years)	undiluted benzyl benzoate (1cm-diameter area)	24-h closed patch test	No skin irritation <sup>110</sup>
Benzyl Benzoate (50% in diethyl phthalate/ethanol)	21 subjects (24 to 70 years old)	50% (0.3 ml)	24-h occlusive patch test	Minimal erythema in 3 subjects; negligible irritation potential <sup>111</sup>
Benzyl Benzoate (50% in ethanol/diethyl phthalate)	129 subjects (46 males, 83 females)	50% (0.3 ml)	RIPT (24-h applications)	No effects <sup>112</sup>
Benzyl Benzoate (30% in petrolatum)	Patients (no. not stated)	30%	Maximization test	No positive reaction <sup>3</sup>
Benzyl Benzoate (30% in petrolatum)	20 healthy males	30%	Occlusive patch test; 5 48 h applications to evaluate sensitization potential	No positive reaction <sup>113</sup>
Benzyl Benzoate (20% in vaselinum aldum or unguentum hydrophilicum vehicle)	34 healthy male and female subjects	20%	48-h closed patch test	No skin irritation <sup>114</sup>
Benzyl Benzoate (4% in petrolatum)	25 healthy male and female subjects	4%	48-h closed patch test	No skin irritation <sup>115</sup>
Benzyl Benzoate (2% in unguentum simplex or unguentum hydrophilicum vehicle)	30 healthy male and female subjects (18 + years)	2%	24- to 72-h closed patch test	No skin irritation <sup>114</sup>
<b>Provocative Tests/Treatment</b>				
Benzyl Alcohol (1% in petrolatum)	11,373 dermatitis patients (males and females)	1%	24 h or 48 h patch test (Finn chambers)	Questionable/irritant reactions (62 patients); sensitization reactions (46 patients) <sup>116</sup>
Benzyl Alcohol (5% in petrolatum)	102 patients	5%	24 h patch tests (Finn chambers)	8 subjects with positive reactions <sup>117</sup>
Benzyl Alcohol (1%)	2,166 dermatitis patients	1%	24 or 48 h patch tests	0.3% with positive reactions <sup>118</sup>
Benzoic Acid (5% in physiological saline)	7 patients (males and females) with birch pollen allergy	5%	45 min epicutaneous application (AL-tests on Scanpor)	Erythema at application sites of all patients (contact urticaria) <sup>119</sup>
Benzoic Acid (5% in petrolatum)	102 patients	5%	24 h patch tests (Finn chambers)	20 subjects with positive reactions <sup>117</sup>
Benzoic Acid	1,252 patients with oral mucosal disease	Concentration not stated	Contact urticaria test; Delayed hypersensitivity test	366 patients positive (urticaria test); 124 patients positive (hypersensitivity test) <sup>120</sup>

**Table 5.** Skin Irritation and Sensitization Studies

Test Substance	Animals/Subjects	Doses/Concentrations Tested	Procedure	Results
Benzoic Acid	417 patients	Concentration not stated	48 h occlusive patch tests (Finn chambers; 10 mm diameter)	5% of patients with questionable/irritant reactions; 4.3% with allergic reactions <sup>121</sup>
Benzoic Acid	465 patients	Concentration not stated	Not stated	Positive reactions (allergenicity) in 2.1% of patients <sup>122</sup>
Sodium Benzoate	465 patients	Concentration not stated	Not stated	Positive reactions (allergenicity) in 1.9% of patients <sup>122</sup>
Benzyl Benzoate (20% emulsion)	1,000 scabies patients	20%	Treatment for scabies	No dermatitis after treatment <sup>123</sup>
Benzyl Benzoate (0.1, 2, and 20% in vaseline ointment or cream base)	175 patients with and without dermatoses	0.1, 2, and 20%	Closed patch test (no GLP) to evaluate sensitization potential	No positive reactions <sup>3</sup>
Benzyl Benzoate (10% in water)	73 dermatitis patients	10%	Patch test (no GLP) to evaluate sensitization potential	No positive reactions <sup>3</sup>
Benzyl Benzoate (5%)	11 male and female patients (18+ years)	5%	24 h application	Positive reaction in 3 of 8 patients <sup>104</sup>
Benzyl Benzoate (5% in petrolatum)	20 male and female patients (18+ years)	5%	48-h patch test (aluminum-backed strips)	Positive reaction in 1 of 20 patients <sup>124</sup>
Benzyl Benzoate (5% in petrolatum)	45 dermatitis patients (18+ years)	5%	48-h closed patch test	No reaction <sup>125</sup>
Benzyl Benzoate (5% in petrolatum)	73 eczematous dermatitis patients	5%	Patch test (no GLP)	1 positive reaction <sup>3</sup>
Benzyl Benzoate (5% in petrolatum)	Eczema patients (no. not stated)	5%	Patch test (no GLP)	No positive reactions <sup>3</sup>
Benzyl Benzoate (5% in petrolatum)	8 dermatitis patients	5%	Patch test (no GLP)	1 positive reaction <sup>3</sup>
Benzyl Benzoate (5% in petrolatum)	19 dermatitis patients	5%	Patch test (no GLP)	No positive reactions <sup>3</sup>
Benzyl Benzoate (5% in yellow soft paraffin)	241 patients	5%	Patch test (no GLP)	No positive reactions <sup>3</sup>
Benzyl Benzoate (5% in vaseline)	103 patients	5%	Patch test (no GLP)	12 positive reactions <sup>3</sup>
Benzyl Benzoate (5% in vaseline)	115 patients	5%	Patch test (no GLP)	14 positive reactions <sup>3</sup>
Benzyl Benzoate (5% in vaseline)	465 dermatitis patients	5%	Patch test (no GLP)	7 positive reactions <sup>3</sup>
Benzyl Benzoate	443 patients (383 tested and 60 controls)	5%	Patch test (no GLP)	2 positive reactions (1 cosmetic dermatitis patient; 1 non-cosmetic dermatitis patient) <sup>3</sup>
Benzyl Benzoate (5%)	271 patients (225 tested and 46 controls)	5%	Patch test (no GLP)	2 positive reactions (1 cosmetic dermatitis patient; 1 non-cosmetic dermatitis patient) <sup>3</sup>
Benzyl Benzoate (5% in petrolatum)	73 patients (18+ years)	5%	Not stated	Positive reaction in 1 patient <sup>3</sup>

**Table 5.** Skin Irritation and Sensitization Studies

Test Substance	Animals/Subjects	Doses/Concentrations Tested	Procedure	Results
Benzyl Benzoate (5% in petrolatum)	5 male and female urticaria patients (18+ years)	5%	30 min closed patch test	No reaction <sup>126</sup>
Benzyl Benzoate (5% in petrolatum)	12 patients (18+ years)	5%	Patch test to evaluate sensitization potential	No positive reactions <sup>127</sup>
Benzyl Benzoate (5% in petrolatum)	3 female allergic dermatitis patients	5%	Patch tests (Finn chambers) to evaluate sensitization potential	No positive reactions <sup>128</sup>
Benzyl Benzoate (5%; vehicle not reported)	50 dermatitis patients	5%	Patch tests to evaluate sensitization potential	Positive reaction in 1 patient <sup>123</sup>
Benzyl Benzoate (5%)	70 dermatitis patients; 19 eyelid dermatitis patients	5%	48 h patch tests (Finn chambers) to evaluate sensitization potential	2.9% with positive reactions (70 patients); no positive reactions (19 patients) <sup>129</sup>
Benzyl Benzoate (5% in petrolatum, vaseline, or alcohol)	115 patients with positive reactions to Balsam of Peru	5%	48 h closed patch tests to evaluate sensitization potential	Positive reactions in 14 patients <sup>130</sup>
Benzyl Benzoate (5%; vehicle not stated)	225 patients (18+ years)	5%	Patch test to evaluate sensitization potential	Positive reactions in 2 patients <sup>131</sup>
Benzyl Benzoate (5%; vehicle not stated)	658 male and female patients	5%	48 h patch tests (Finn chambers) to evaluate sensitization potential	Positive reaction in 1 patient <sup>132</sup>
Benzyl Benzoate (5% in white petrolatum)	102 contact dermatitis patients (18+ years)	5%	24 h patch tests (Finn chambers) to evaluate sensitization potential	Positive reactions in 4 patients <sup>117</sup>
Benzyl Benzoate (5% in petrolatum)	317 eczema patients (18+ years)	5%	Patch test to evaluate sensitization potential	No positive reactions <sup>133</sup>
Benzyl Benzoate (5% in petrolatum)	73 patients with eczematous dermatitis	5%	Patch test to evaluate sensitization potential	1 positive reaction (allergic contact dermatitis) <sup>123</sup>
Benzyl Benzoate (5% in petrolatum)	102 patients	5%	24 h patch tests (Finn chambers)	4 subjects with positive reactions <sup>117</sup>
Benzyl Benzoate (2 or 5% in vaseline)	144 patients (14 controls included) per test concentration	2 or 5%	Closed patch test (no GLP) to evaluate sensitization potential	1 positive reaction in cosmetic dermatitis patient <sup>3</sup>
Benzyl Benzoate (2 or 5% in petrolatum)	64 dermatitis patients (18+ years)	2 or 5%	Closed patch test to evaluate sensitization potential	Positive reaction in 1 patient (5% concentration); no positive reactions (2%) <sup>134</sup>
Benzyl Benzoate (2%)	Patients (no. not stated)	2%	Patch test (no GLP)	Negative reaction in contact dermatitis patient <sup>3</sup>
Benzyl Benzoate (2%)	198 patients	2%	Patch test (no GLP)	No positive reactions <sup>3</sup>
Benzyl Benzoate (2% in petrolatum)	335 patients	2%	Patch test (no GLP)	No positive reactions <sup>3</sup>



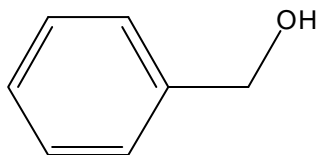
**Table 5.** Skin Irritation and Sensitization Studies

Test Substance	Animals/Subjects	Doses/Concentrations Tested	Procedure	Results
Benzyl Benzoate (2% in yellow paraffin)	539 patients (18+ years; 50 with photosensitivity dermatitis, 32 with polymorphic light eruption (PLE) and 457 with contact dermatitis)	2%	48 h closed patch test to evaluate sensitization potential	Positive reaction in 1 PLE patient <sup>74</sup>
Benzyl Benzoate (2% in petrolatum)	284 patients (18+ years)	2%	Patch test to evaluate sensitization potential	1% with positive reactions <sup>135</sup>
Benzyl Benzoate (2% in petrolatum)	335 patients (18+ years)	2%	Patch test to evaluate sensitization potential	No positive reactions <sup>135</sup>
Benzyl Benzoate (2% in petrolatum)	198 patients (18+ years)	2%	Patch test to evaluate sensitization potential	No positive reactions <sup>136</sup>
Benzyl Benzoate (2% in yellow soft paraffin)	241 patients (18+ years)	2%	Finn chamber technique to evaluate sensitization potential	Positive reaction in 1 male patient <sup>137</sup>
Benzyl Benzoate (2% in white petrolatum)	284 contact dermatitis patients (30 to 67 years)	2%	Patch test to evaluate sensitization potential	No positive reactions <sup>138</sup>
Benzyl Benzoate (1% in yellow petrolatum)	28 patients allergic to perfumes (18+ years)	1%	Sensitization test (procedure not stated)	No positive reactions <sup>139</sup>
Benzyl Benzoate (1%)	2,003 dermatitis patients	1%	24 or 48 h patch tests to evaluate sensitization potential	No positive reactions <sup>118</sup>
Benzyl Benzoate (0.1% in ethanol or non-irritative cream base)	111 male and female dermatitis patients (18+ years)	0.1%	24-h to 48-h closed patch test to evaluate sensitization potential	No positive reactions <sup>114</sup>
Benzyl Benzoate	Cosmetic dermatitis patients (no. not stated)	Concentration not stated	Patch test (no GLP) to evaluate sensitization potential	1 positive reaction <sup>3</sup>
Benzyl Benzoate	15 patients	Concentration not stated	Patch test (no GLP)	delayed in reaction in 1 patient sensitized to Peru balsam <sup>3</sup>
Benzyl Benzoate	111 patients	Concentration not stated	Patch test (no GLP)	No positive reactions <sup>3</sup>
Benzyl Benzoate	142 patients	Concentration not stated	Patch test (no GLP) to evaluate sensitization potential	10 positive reactions in patients sensitive to balsam of Peru <sup>3</sup>
Benzyl Benzoate	155 scabies patients	Concentration not stated	Treatment with benzyl benzoate (concentration not stated)	Dermatitis in 17 patients <sup>3</sup>
Benzyl Benzoate	460 patients	Concentration not stated	Patch test (no GLP) to evaluate sensitization potential	3 positive reactions in patients sensitive to cosmetics <sup>3</sup>
Benzyl Benzoate	25 patients	Concentration not stated	Maximization test (no GLP)	No positive reactions <sup>3</sup>
Benzyl Benzoate	465 patients	Concentration not stated	Not stated	Positive reactions (allergenicity) in 1.5% of patients <sup>122</sup>

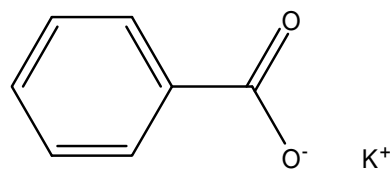
**Table 6.** Case Reports

Test Substance	Subjects	Procedure/Results
Benzyl Alcohol	30-year-old facial dermatitis patient	Patch testing (no test concentration) produced macular erythema <sup>140</sup>
Benzyl Alcohol	38-year old eczema patient	1% aqueous benzyl alcohol: negative prick test results and positive (++) intradermal injection test results. Negative injection test results in 10 healthy controls <sup>141</sup>
Benzyl Alcohol	39-year-old female with pruritic erythema of foot	5% benzyl alcohol in petrolatum: weak (+) reaction in patch test and strong positive reaction in repeated open application test <sup>142</sup>
Benzyl Alcohol	67-year-old male with leg dermatitis	1% benzyl alcohol in petrolatum: 2+ occlusive patch test reaction. 0.9% benzyl alcohol in saline: negative prick test reaction at 0.5 h reading, but marked induration and proximal spread over arm at days 3 to 8 <sup>143</sup>
Benzyl Alcohol	53-year-old with stasis dermatitis	1% benzyl alcohol in petrolatum: redness and swelling at 1 h after patch application, wheal 1 day later, and mild urticaria at day 5 <sup>144</sup>
Benzyl Alcohol	16-year-old female with possible anaphylactic reactions after i.m. injection with B12 preparation containing 0.9% benzyl alcohol	Benzyl alcohol preparation: negative in prick tests, but positive in intradermal tests <sup>145</sup>
Benzyl Alcohol	57-year-old female with pruritic dermatitis	Allergic contact dermatitis (+, ++, or +++ reaction) after patch testing with benzyl alcohol <sup>146</sup>
Benzyl Alcohol	40-year-old female with dermatitis	Positive patch test reaction (+++, allergic contact dermatitis) to 9.5% benzyl alcohol in petrolatum <sup>147</sup>
Benzyl Alcohol	65-year-old female with eyelid dermatitis	Macular erythema after patch testing with benzyl alcohol (concentration not stated) <sup>148</sup>
Benzyl Alcohol	30-year-old female with eyelid dermatitis	Positive (1+) patch test reaction to benzyl alcohol (concentration not stated) <sup>140</sup>
Benzoic Acid	30-year-old facial dermatitis patient	Patch testing (no test concentration) produced 1+ reaction <sup>140</sup>
Benzoic Acid	46-year-old female with history of erythema and itching	5% benzoic acid in petrolatum: positive reaction <sup>149</sup>
Sodium Benzoate	64-year-old female with erythema and edema of finger	Positive patch test reaction to 5% sodium benzoate on days 2 and 4; negative patch test results at interdigital area at 2 months <sup>150</sup>
Sodium Benzoate	52-year-old female with hand dermatitis	Negative patch test reaction to sodium benzoate (concentration not stated) <sup>151</sup>
Sodium Benzoate	75-year-old female with history of pruritus	Relapse of pruritus after 100 mg oral dose of sodium benzoate <sup>152</sup>
Benzyl Benzoate	25-year-old male with acute bullous eruption after application of 30% benzyl benzoate preparation	Negative patch test reaction to 1% benzyl benzoate in petrolatum <sup>153</sup>
Benzyl Benzoate (unspecified concentration or 25% emulsion in aqueous solution of triethanolamine stearate)	4 scabies patients (ages not stated)	Severe skin irritation; pruritic dermatitis within 12 to 20 h after treatment (unspecified concentration or 25% emulsion) <sup>154</sup>
Benzyl Benzoate	Scabies patients (no. not included)	Treatment with benzyl benzoate solution (concentration not stated) did not cause dermatitis <sup>3</sup>
Benzyl Benzoate	4 scabies patients	Treatment with benzyl benzoate solution caused dermatitis <sup>3</sup>
Benzyl Benzoate (5% in petrolatum)	1 patient (18 + years)	Reaction read at 48 h and 96 h. No sensitization <sup>155</sup>
Benzyl Benzoate 2% in petrolatum)	23-year old male patient	Negative patch test results; no sensitization <sup>156</sup>

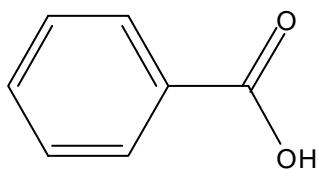
**Figure 1. Chemical Structures**



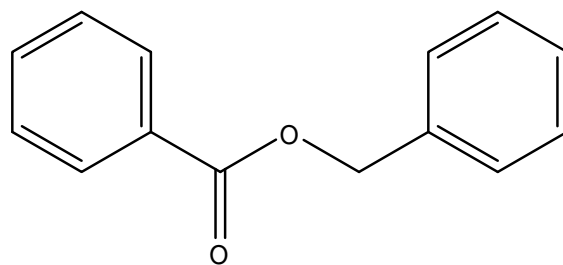
**benzyl alcohol**



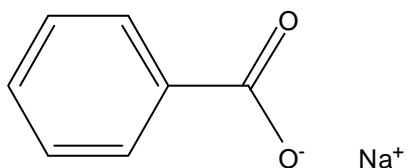
**potassium benzoate**



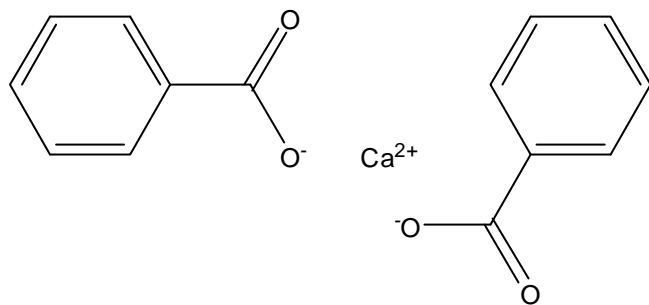
**benzoic acid**



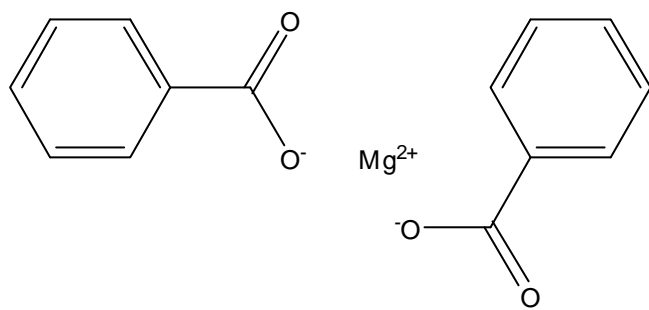
**benzyl benzoate**



**sodium benzoate**



**calcium benzoate**



**magnesium benzoate**

## References

1. Nair, B. Bergfeld W. F. Belsito D. V. Snyder P. W. Klaassen C. D. Schroeter A. L. Shank R. C. and Slaga T. J. and Andersen F. A. Final report on the safety assessment of benzyl alcohol, benzoic acid, and sodium benzoate. *International Journal of Toxicology*. 2001;20:23-50.
2. OECD. SIDS initial assessment report for 13th SIDS initial assessment meeting. Benzoates: benzoic acid, sodium benzoate, potassium benzoate, and benzyl alcohol. <http://webnet3.oecd.org/eChemPortal>. 2001. Date Accessed 7-12-2010.
3. European Commission - European Chemicals Bureau. IUCLID dataset on benzyl benzoate. <http://tcsweb3.jrc.it/e\sis/>. 2000. Date Accessed 11-18-2010.
4. Oussi, D. Mokriani A. Chamarro E. and Esplugas S. Photodegradation of benzoic acid in aqueous solutions. *Environmental Technology*. 1998;19:(9):955-960.
5. Sudareva, N. N. and Chubarova E. V. Time-dependent conversion of benzyl alcohol to benzaldehyde and benzoic acid in aqueous solutions. *J.Pharm.Biomed.Anal.* 2006;41:(4):1380-1385.
6. Shibamoto, T. and Umamo K. Photochemical products of benzyl benzoate: Possible formation of skin allergens. *J.Toxicol Cutaneous Ocul.Toxicol.* 1985;4:(2):97-104.
7. Sax, N. I. and Lewis R. J. Condensed chemical dictionary. 11 ed. New York: Van Nostrand Reinhold Company, 1987.
8. Food and Drug Administration (FDA). Code of Federal Regulations. Title 21 part 184.1733. Washington, D.C.: Government Printing Office, 2010.
9. Bhattacharya, A. Patel N. Vasques T. Tichkule R. Darmar G. and Wu J. Surfactant-mediated solvent-free dealkylative cleavage of ethers and esters and trans-alkylation under neutral conditions. *Tetradron Letters*. 2006;47:565-567.
10. Kamm, O. and Kamm W. F. Benzyl benzoate. *Org.Synth.* 1941;1:104.
11. Lewandowski, W. and Baranska H. *J.Mol.Struct.* 1988;174:417-422.
12. Khan, K. M. Maharvi G. M. Hayat S. Zia U. Choudhary M. I. and Atta-ur-Rahman T. *Tetrahedron*. 2003;59:5549-5554.
13. Sun, L., Li, G., Wan, S., and An, T. Mechanistic study and mutagenicity assessment of intermediates in photocatalytic degradation of gaseous toluene. *Chemosphere*. 2010;78:(3):313-318.
14. Tsipis, C. A. Hadjikostas C. C. and Sigalas M. P. *Spectrochim.Acta Part A*. 1982;38:41-50.
15. Morris. *J.Assoc.Off.Anal.Chem.* 1973;56:1037-1043.
16. Butler, L. G. Cheng C. P. and Brown T. L. *J.Phys.Chem.* 1981;85:2738-2740.
17. Nyman, P. J. Wamer W. G. Begley T. H. Diachenko G. W. and Perfetti G. A. Evaluation of accelerated UV and thermal testing for benzene formation in beverages containing benzoate and ascorbic acid. *Journal of Food Science*. 2010;75:C263-C267.

18. Environmental Protection Agency (EPA). Basic information about benzene in drinking water. <http://water.epa.gov/drink/contaminants/basicinformation/benzene.cfm>. 2011. Date Accessed 8-7-2011.
19. Nyman, P. J., Diachenko, G. W., Perfetti, G. A., Mcneal, T. P., Hiatt, M. H., and Morehouse, K. M. Survey results of benzene in soft drinks and other beverages by headspace gas chromatography/mass spectrometry. *J.Agric.Food Chem.* 2008;56:(2):571-576.
20. United States Pharmacopoeial Convention, Inc. The United States Pharmacopoeia. Monograph on benzyl benzoate. 32 ed. Rockville: United States Pharmacopoeial Convention, Inc., 2009.
21. The United States Pharmacopoeial Convention. Food Chemicals Codex. 6th ed. Rockville: The United States Pharmacopoeial Convention, 2009.
22. Gottschalck, T. E. and Bailey J. E. International Cosmetic Ingredient Dictionary and Handbook. 13th ed. Washington, D.C.: Personal Care Products Council, 2010.
23. Food and Drug Administration (FDA). Information supplied to FDA by industry as part of the VCRP FDA database. 2011. Washington, D.C.: FDA.
24. Personal Care Products Council. Use concentration data from industry survey. Unpublished data submitted by the Personal Care Products Council, May 19, 2010. 2010. pp.1-7.
25. European Union. Consolidated version of Cosmetic Directive 76/768/EEC, as amended, Annexes I through IX. 2008. <http://ec.europa.eu/enterprise/sectors/cosmetics/documents/directive/#h2-technical-adaptations-to-be-incorporated-in-the-consolidated-textDate> Accessed 10-15-2010
26. Scientific Committee on Cosmetic Products and Non-Food Products Intended for Consumers. Opinion of the Scientific Committee on Consumer Products and Non-Food Products Intended for Consumers concerning benzoic acid and sodium benzoate. 2002. pp.1-20.
27. Scientific Committee on Consumer Products (SCCP). Opinion on benzoic acid and sodium benzoate. pp.1-30.
28. Food and Drug Administration (FDA). Direct food substances affirmed as generally recognized as safe. Benzoic acid. 21 CFR 184.1021. 2011.
29. Food and Drug Administration (FDA). Direct food substances affirmed as generally recognized as safe. Sodium Benzoate. 21CFR 184.1733. 2011.
30. Food and Drug Administration (FDA). Antimycotics. Sodium benzoate. 21 CFR 181.23. 2011.
31. Food and Drug Administration (FDA). Food additives permitted for direct addition to food for human consumption. Benzyl alcohol and benzyl benzoate. 21 CFR 172.515. 2011.
32. Food and Drug Administration (FDA). Substances for use only as components of coatings. Benzyl alcohol and benzyl benzoate. 21 CFR 175.105. 2011.
33. Food and Drug Administration (FDA). Substances for use as components of coatings. Benzyl alcohol and benzoic acid. 21 CFR 175.300. 2011.

34. Food and Drug Administration (FDA). Substances for use as basic components of single and repeated use food contact surfaces. Benzyl alcohol and potassium benzoate. 21 CFR 177.1210. 2011.
35. Food and Drug Administration (FDA). Artificially sweetened fruit jelly - optional ingredients. Benzoic acid and sodium benzoate. 21 CFR 150.141. 2011.
36. Food and Drug Administration (FDA). Artificially sweetened fruit preservatives and jams - optional ingredients. Benzoic acid and sodium benzoate. 21 CFR 150.161. 2011.
37. Food and Drug Administration (FDA). Requirements for specific standardized margarine - preservatives. Benzoic acid, sodium benzoate, potassium benzoate, and calcium benzoate. 21 CFR 166.110. 2011.
38. Food and Drug Administration (FDA). Substances for use as basic components of single and repeated use food contact surfaces. Benzoic acid. 21 CFR 177.1390. 2011.
39. Food and Drug Administration (FDA). Antioxidants and/or stabilizers for polymers. Calcium benzoate. 21 CFR 178.2010. 2011.
40. Food and Drug Administration (FDA). Diluents in color additive mixtures for drug use exempt from certification. Benzyl alcohol. 21 CFR 73.1001. 2011.
41. FDA. FDA News Release. FDA approves benzyl alcohol lotion for the treatment of head lice. <http://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm149562.htm>. 2009. Date Accessed 1-25-2011.
42. Meinking, T. L., Villar, M. E., Vicaria, M., Eyerdam, D. H., Paquet, D., Mertz-Rivera, K., Rivera, H. F., Hiriart, J., and Reyna, S. The clinical trials supporting benzyl alcohol lotion 5% (Ulesfia): a safe and effective topical treatment for head lice (pediculosis humanus capitis). *Pediatr.Dermatol.* 2010., Jan.1. 27(1):19-24.:(1:19-24):ediatric.
43. Brooks, P. A. and Grace R. F. Ivermectin is better than benzyl benzoate for childhood scabies in developing countries. *J.Paediatr.Child Health*. 2002;38:(4):401-404.
44. FDA. OTC active drug ingredients. <http://www.fda.gov/downloads/AboutFDA/CentersOffices/CDER/UCM135688.pdf>. 2010. Date Accessed 12-12-2010.
45. Environmental Protection Agency. Inert ingredients permitted for use in nonfood use pesticide products. <http://www.epa.gov/opprd001/inerts/lists.html>. 2011.
46. Environmental Protection Agency. Suspension of registrations under FIFRA section 3(c)(2)(B). <http://www.epa.gov/pesticides/regulating/registering/suspensions.htm>. 2010. Date Accessed 10-14-2010.
47. Scheman, A. J. and Katta R. Photographic allergens: an update. *Contact Dermatitis*. 1997;37:(3):130-130.
48. Smith, A. L. Haider K. Schachtner J. M. Mathur S. Vanorden R. and Gentile T. C. Fatal hemolysis after high-dose etoposide: Is benzyl alcohol to blame? *Pharmacotherapy*. 2001;21:(6):764-766.

49. Shehab, N., Lewis, C. L., Streetman, D. D., and Donn, S. M. Exposure to the pharmaceutical excipients benzyl alcohol and propylene glycol among critically ill neonates. *Pediatr.Crit Care Med.* 2009, Mar. 10(2):256-9.:(2:256-9):ediatric.
50. Pack, G. T. and Underhill F. P. The influence of benzyl benzoate upon nitrogen metabolism. *Journal Metabolic Research.* 1922;2:73-105.
51. Snapper, J Grunbaum A. and Sturkop s. About the fission and oxidation of benzyl alcohol and benzyl esters in the human organism. *Biochemische Zeitschrift.* 1925;155:163-173.
52. Kitagawa, S. and Li H. Effects of removal of stratum corneum, delipidization and addition of enhancers, ethanol and *l*-menthol, on skin permeation of benzoic acid and its 4-*n*-alkyl substituents in excised guinea pig dorsal skin. *Chem.Pharm.Bull.* 1999;47:(1):44-47.
53. Bronaugh, R. L. Wester R. C. Bucks D. Maibach H. I. and Sarason R. In vivo percutaneous absorption of fragrance ingredients in Rhesus monkeys and humans. *Fd.Chem.Toxic.* 1990;28:(5):369-373.
54. Jimbo, Y. Penetration of fragrance compounds through human epidermis. *Journal of Dermatology.* 1983;10:(3):229-239.
55. Miller, M. A. Bhatt V. and Kasting G. B. Dose and airflow dependence of benzyl alcohol disposition on skin. *J.Pharm.Sci.* 2006;95:(2):281-291.
56. Nielsen, N. M. and Bundgaard H. Prodrugs as delivery systems. 68. Chemical and plasma-catalyzed hydrolysis of various esters of benzoic acid: A reference system for designing prodrug esters of carboxylic acid agents. *International Journal of Pharmacology.* 1987;39:75-85.
57. WIL Research Laboratories, LLC. A 4-week inhalation toxicity study of aerosolized benzyl alcohol and benzoic acid in Sprague-Dawley rats. WIL-703002. Unpublished data submitted by the Personal Care Products Council on 2-25-2010. 2010. pp.1-990.
58. Nishimura, H. Saito S. Kishida F. and Matsuo M. Analysis of acute toxicity (LD50 value) of organic chemicals to mammals by solubility parameter ( $\delta$ ). 3. Acute dermal toxicity to rabbits. *Jpn.J.Ind.Health.* 1994;36:(6):428-434.
59. Postel, E. Benzyl benzoate, a component of Peru balsam. *Klinische Wochenschrift.* 1943;22:362-364.
60. Ornellas, M. R. Chemotherapeutic activity of benzyl benzoate. *Bulletin Cancer.* 1965;52:(4):371-378.
61. Moreno, O. M. Ratcliffe H. L and Singh H. 30-day dermal toxicity study of benzylbenzoate in rats. Unpublished report 4000. Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1980.
62. Lehman, A. J. Insect repellants. *Association Food Drug Officials United States.* 1955;19:(2):87-99.
63. Graham, B. E. and Kuizenga M. H. Toxicity studies on benzyl benzoate and related benzyl compounds. *The Journal of Pharmacology and Experimental Therapeutics.* 1945;84:(4):358-362.
64. Heller, A. E. and Steinfield E. Nonleucotoxic properties of benzyl benzoate. *New York Medical Journal.* 1920;112:160-161.

65. Morrison, V. L. Koh H. J. Cheng L. Bessho K. Davidson M. C. and Freeman W. R. Intravitreal toxicity of the kenalog vehicle (benzyl alcohol) in rabbits. *Retina*. 2006;26:(3):339-344.
66. Li, Q., Wang, J., Yang, L., Mo, B., Zeng, H., Wang, N., and Liu, W. A morphologic study of retinal toxicity induced by triamcinolone acetonide vehicles in rabbit eyes. *Retina*. 2008, Mar. 28(3):504-10. (3:504-10):Retina.
67. Carvounis, P. E. and Holz E. R. Comment on the publication "Vehicle of triamcinolone acetonide is associated with retinal toxicity and transient increase in lens density" by Kai et al. *Graefes Arch.Clin.Exp.Ophthalmol*. 2007;245:(1):179-180.
68. Zhengyu, S., Fang, W., and Ying, F. Vehicle used for triamcinolone acetonide is toxic to ocular tissues of the pigmented rabbit. *Curr.Eye Res*. 2009;34:(9):769-776.
69. International Flavors and Fragrances. Acute eye irritation test in rabbits with benzyl benzoate. Unpublished data available from the Research Institute for Fragrance materials (RIFM). 1982.
70. Sugihara, N. Shimomichi K. and Furuno K. Cytotoxicity of food preservatives in cultured rat hepatocytes loaded with linoleic acid. *Toxicology*. 1997;120:(1):29-36.
71. Takasago International Corporation. Toxicology studies of benzyl alcohol in the guinea pig. Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1981. November 18A.
72. Placzek, M. Krosta I. Gaube S. Eberlein-KÖNIG B. and Przybilla B. Evaluation of phototoxic properties of antimicrobials used in topical preparations by a photohaemolysis test. *Acta Derm.Venereol*. 2005;85:(1):13-16.
73. Placzek, M. Frömel W. Eberlein B. Gilbertz K. P. and Przybilla B. Evaluation of phototoxic properties of fragrances. *Acta.Derm.Venereol*. 2007;87:(4):312-316.
74. Addo, H. A. Ferguson J. Johnson B. E. and Frain-Bell W. The relationship between exposure to fragrance materials and persistent light reaction in the photosensitivity dermatitis with actinic reticuloid syndrome. *British Journal of Dermatology*. 1982;107:(3):261-274.
75. Mowafy, A. R. Darwish A. M. El-Kholy S. A. and Abdel-Moshen S. H. Effect of food preservatives on mother rats and survival of their offspring. *J.Egypt Public Health Assoc*. 2001;76:(3-4):281-295.
76. Morita, S. Yamada A. Ahgaki S. Noda T. and Taniguchi S. Safety evaluation of chemicals for use household products. I. Teratological studies on benzyl benzoate and 2-(morpholinothio) benzothiazole in rats. *Annual Report Osaka City Inst.Public Health Environment Sci*. 1980;43:90-97.
77. Eibs, H. Speilmann H. Haegele M. and Klose I. Effects of Steroid Sex Hormones on the Development of Early Mouse Embryos In Vitro and In Vivo. Chapter: 6. In: *Advances in the Study of Birth Defects, 2 (Teratological Testing)*. 1979:113-137.
78. Gonzalez, F. W. Effect of benzyl benzoate on egg-hatching in *Drosophila melanogaster*. *Newsletter Environmental Mutagen Society*. 1971;5:36.
79. Mytton, O. T. McGready R. Lee S. J. Roberts C. H. Ashley E. A. Carrara V. I. Thwai K. L. Jay M. P. Wiangambun T. Singhasivanon P. and Nosten F. Safety of benzyl benzoate lotion and permethrin in pregnancy: a retrospective matched cohort study. *BJOG*. 2007;114:(5):582-587.



80. Ashby, J. Lefevre P. A. Odum J. Tinwell H. Kennedy S. J. Beresford N. and Sumpter J. P. Failure to confirm estrogenic activity for benzoic acid and clofibrate: Implications for lists of endocrine-disrupting agents. *Regul.Toxicol.Pharmacol.* 1997;26:96-101.
81. Hashimoto, Y. Kawaguchi M. Miyazaki K. and Nakamura M. Estrogenic activity of tissue cojnditioners in vitro. *Dent.Mater.* 2003;19:(4):341-346.
82. Charles, A. K. and Darbre P. D. Oestrogenic activity of benzyl salicylate, benzyl benzoate and butylphenylmethylpropional (Lilial) in MCF7 human breast cancer cells *in vitro*. *J.Appl.Toxicol.* 2009;422-434.
83. Demir, E., lu, S., and Kaya, B. Assessment of genotoxic effects of benzyl derivatives by the comet assay. *Food Chem.Toxicol.%2010., May.* 48(5):1239-42.:(5:1239-42):Food.
84. Nesslany, F. and Marzin D. A micromethod for the *in vitro* micronucleus assay. *Mutagenesis.* 1999;14:(4):403-410.
85. Mpountoukas, P., Vantarakis, A., Sivridis, E., and Lialiaris, T. Cytogenetic study in cultured human lymphocytes treated with three commonly used preservatives. *Food Chem.Toxicol.%2008., Jul.* 46(7):2390-3.:(7:2390-3):Food.
86. Ishizaki, M. and Ueno S. The DNA-damaging activity of natural and synthetic food additives (V). *J.Food Hyg.Soc.Jpn.* 1989;30:(5):447-451.
87. Florin, I. Rutberg L. Curvall M. and Enzell C. R. Screening of tobacco smoke constituents for mutagenicity using the Ames test. *Toxicology.* 1980;18:(3):219-232.
88. Salih, F. M. Risk assessment of combined photogenotoxic effects of sunlight and food additives. *Sci.total Environ.* 2006;362:(1-3):68-73.
89. Stefanidou, M., Alevizopoulos, G., and Spiliopoulou, C. DNA content of Tetrahymena pyriformis as a biomarker for different toxic agents. *Chemosphere.%2008., Dec.* 74(1):178-80.:(1:178-80):Chemosphere.
90. O'Neil, M. J. The Merck Index Online. Whitehouse Station, NJ: Merck & Co., Inc., 2010.
91. Scientific and Technical Information Network. Beilstein record on calcium benzoate. <https://stnweb.cas.org>. 2010. Date Accessed 12-7-2010.
92. National Library of Medicine. ChemIDplus record on magnesium benzoate. <http://www.nlm.nih.gov>. 2010. Date Accessed 12-7-2010.
93. Barratt, M. D. Quantitative structure-activity relationships for skin irritation and corrosivity of neutral and electrophilic organic chemicals. *Toxicology In Vitro.* 1996;10:(3):247-256.
94. Haarmann & Reimer GmbH. OECD acute toxicity test: An assessment of the acute oral toxicity of benzyl benzoate in the rat. Unpublished (July 03a). Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1985.
95. Nishimura, F. T. Saito S. Kishida F. and Matsuo M. Analysis of acute toxicity (LD50-value) of organic chemicals to mammals by solubility parameter (delta). 1. Acute oral toxicity to rats. *Japanese Journal Industrial Health.* 1994;36:(5):314-323.

96. International Flavors and Fragrances. Acute oral toxicity study in rats with benzyl benzoate (14 days). Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1982.
97. International Flavors and Fragrances. Acute oral toxicity study in rats with benzyl benzoate (14 days). Unpublished (June 9a, June 9b). Unpublished data available from the Research institute for Fragrance Materials (RIFM). 1982.
98. International Flavors and Fragrances. Acute oral toxicity study in rats with benzyl benzoate (14 days). Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1982.
99. International Flavors and Fragrances. Acute oral toxicity study in rats with benzyl benzoate (14 days). Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1982.
100. Draize, J. H. Alvarez E. Whitesell M. F. Woodard G. Hagan E. C. and Nelson A. A. Toxicological investigations of compounds proposed for use as insect repellents. *The Journal of Pharmacology and Experimental Therapeutics*. 1948;93:26-39.
101. Quest International. Guinea pig skin irritation test with benzyl benzoate. Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1981.
102. Ishihara, M. Itoh M. Nishimura M. Konoshita M. Kantoh H. Nogami T. and Yamada K. Closed epicutaneous test. *Skin Research*. 1986;28:(2):230-240.
103. Quest International. Guinea pig skin irritation test with benzyl benzoate. Unpublished data available from the Research Institute for Fragrance Materials (RIFM). 1977.
104. Hausen, B. M. Evers P. Stuwe H. T. Konig W. A. and Wollenweber E. Propolis allergy (IV). Studies with further sensitizers from propolis and constituents common to propolis, poplar buds, and balsam of Peru. *Contact Dermatitis*. 1992;26:(1):34-44.
105. Hausen, B. M. Simatupang T. Bruhn G. Evers P. and Koenig W. A. Identification of new allergenic constituents and proof of evidence of coniferyl benzoate in Balsam of Peru. *Am J. Contact Dermat*. 1995;6:(4):199-208.
106. Research Institute for Fragrance Materials, inc. Primary irritation test in rabbits. Report to the Research Institute for Fragrance Materials (RIFM, report 3099). Unpublished data available from RIFM. 1985.
107. Basketter, D. A. Chamberlain M. Griffiths H. A. Rowson M. Whittle E. and York M. The classification of skin irritants by human patch test. *Food and Chemical Toxicology*. 1997;35:(8):845-852.
108. Coverly, J. Peters L. Whittle E. and Basketter D. A. Susceptibility of skin stinging, non-immunologic contact urticaria and acute skin irritation; is there a relationship? *Contact Dermatitis*. 1998;38:(2):90-95.
109. Marriott, M. Holmes J. Peters L. Cooper K. Rowson M. and Basketter D. A. The complex problem of sensitive skin. *Contact Dermatitis*. 2005;53:(2):93-99.
110. Katz, A. E. Dermal irritating properties of essential oils and aromatic chemicals. *The Spice Mill*. 1946;69:46-51.

111. Research Institute for Fragrance Materials (RIFM). Modified primary dermal irritation test(MPDI) with benzyl benzoate (report 44230). Unpublished data available from RIFM. 2003.
112. Research Institute for Fragrance Materials (RIFM). Repeated insult patch test with benzyl benzyl benzoate (report 47159). Unpublished data available from RIFM. 2004. July 7b.
113. Research Institute for Fragrance Materials (RIFM). The contact sensitizing potential of fragrance materials in humans (report 1760). Unpublished data available from RIFM. 1970.
114. Fujii, T. Furukawa S. and Suzuki S. Studies on compounded perfumes for toilet goods. On the non-irritative compounded perfumes for soaps. *Yukugaku*. 1972;21:(12):904-908.
115. Research Institute for Fragrance Materials (RIFM). Report on human maximization studies (report 1787). Unpublished data available from RIFM. 1978.
116. Schnuch, A. Geier J. Uter W. and Frosch P. J. Patch testing with preservatives, antimicrobials and industrial biocides. Results from a multicentre study. *British Journal of Dermatology*. 1998;138:(3):467-476.
117. Hausen, B. M. Contact allergy to Balsam of Peru. II. Patch test results in 102 patients. *Am J.Contact Dermat*. 2001;12:(2):93-102.
118. Schnuch, A. Uter W. Geier J. Lessman H. and Frosch P. J. Sensitization to 26 fragrances to be labelled according to current European regulation. *Contact Dermatitis*. 2007;57:(1):1-10.
119. Wallengren, J. and Larsson B. Nitric oxide participates in prick test and irritant patch test reactions in human skin. *Arch.Dermatol.Res*. 2001;293:(3):121-125.
120. Wray, D. Rees S. R. Gibson J. and Forsyth A. The role of allergy in oral mucosal diseases. *Q J Med*. 2000;93:(8):507-511.
121. Kanerva, L. Rantanen T. Aalto-Korte K. et al. A multicenter study of patch test reactions with dental screening series. *American Journal of Contact Dermatitis*. 2001;12:(2):83-87.
122. Meynadier, J.-M. Meynadier J. Colmas A. Castelain P. Y. Ducombs G. Chabeau G. Lacroix M. Martin P. and Ngangu Z. Allergy to preservatives. *Ann.Dermatol.Venereol*. 1982;109:(12):1017-1023.
123. Meneghini, C. L. Vena G. A. and Angelini G. Contact dermatitis to scabicides. *Contact Dermatitis*. 1982;8:(4):285-286.
124. Larsen, W. G. Perfume dermatitis: A study of 20 patients. *Archives of Dermatology*. 1977;113:623-626.
125. Itoh, M. Sensitization potency of some phenolic compounds - with special emphasis on the relationship between chemical structure and allergenicity. *Journal of Dermatology*. 1982;9:(3):223-233.
126. Forsbeck, M. and Skog E. Immediate reactions to patch tests with balsam of Peru. *Contact Dermatitis*. 1977;3:201-205.
127. Rudzki, E. and Grzywa Z. Dermatitis from propolis. *Contact Dermatitis*. 1983;9:40.
128. Ljunggren, B. Contact dermatitis to estradiol benzoate. *Contact Dermatitis*. 1981;7:(3):141-144.

129. Nethercott, J. R. Nield G. and Holness D. L. A review of 79 cases of eyelid dermatitis. *Journal of the American Academy of Dermatology*. 1989;21:223-230.
130. Hjorth, N. Other aromatic substances related to balsams. Chapter: 10. In: *Eczematous allergy to balsams*. 1961:94-111.
131. Nishimura, M. Ishihara M. Itoh M. Hosono K. and Kantoh H. Results of patch tests on cosmetic ingredients conducted between 1979 and 1982. *Skin Research*. 1982;26:(4):945-954.
132. Heydorn, S. Johansen J. D. Andersen K. E. Bruze M. Svedman C. White I. R. Basketter D. A. and Menne T. Fragrance allergy in patients with hand eczema - a clinical study. *Contact Dermatitis*. 2003;48:(6):317-323.
133. Heydorn, S. Johansen J. D. Andersen K. E. Basketter D. Bruze M. Karlberg A. T. White I. and Menne T. Identification of fragrances relevant to hand eczema. *Contact Dermatitis*. 2002;46:(4):21-22.
134. Ishihara, M. Itoh S. Hayashi S. and Satake T. Methods of diagnosis in cases of cosmetic dermatitis and facial melanosis in females. *Nishinohon J.Dermatology*. 1979;41:(3):426-439.
135. Mitchell, J. C. Adams R. M. Glendenning W. E. Fisher A. Kanof N. Larsen W. Maibach H. I. Rudner E. J. Schnorr W. Storrs F. and Taylor J. S. Results of standard patch tests with substances abandoned. *Contact Dermatitis*. 1982;8:(5):336-337.
136. Rudner, E. J. North American group results. *Contact Dermatitis*. 1977;3:(3):208-209.
137. Ferguson, J. and Sharma S. Cinnamic aldehyde test concentrations. [Letter to the Editor]. *Contact Dermatitis*. 1984;10:191-192.
138. Angelini, G. Vena G. a. Giglio G. Fiordalisi F. and Meneghini C. L. Contact dermatitis due to cosmetics. *Journal of Applied Cosmetology*. 1985;3:(3):223-233.
139. Meynadier, J. M. Meynadier J. Peyron J. L. and Peyron L. Clinical forms of skin manifestatins in allergy to perfume. *Annales Dermatologie Venereologie*. 1986;113:(1):31-39.
140. Jacob, S. E. and Stechschulte, S. Eyelid dermatitis associated with balsam of Peru constituents: benzoic acid and benzyl alcohol. *Contact Dermatitis*. 2008; Feb. 58(2):111-2.:(2:111-2):Contact.
141. Verecken, P. Birringer C. Knitelius A. C. Herbaut D. and Germaux M. A. Sensitization to benzyl alcohol: a possible cause of "corticosteroid allergy". *Contact Dermatitis*. 1998;38:(2):106-106.
142. Podda, M. Grundmann-Kollmann M. Kaufmann R. and Boehncke W. H. Allergic contact dermatitis from benzyl alcohol during topical antimycotic treatment. *Contact Dermatitis*. 1999;41:(5):302-303.
143. Shaw, D. W. Allergic contact dermatitis to benzyl alcohol in a hearing aid impression material. *Am J.Contact Dermat*. 1999;10:(4):228-232.
144. Guin, J. D. and Goodman J. Contact urticaria from benzyl alcohol presenting as intolerance to saline soaks. *Contact Dermatitis*. 2001;45:(3):182-183.
145. Turvey, S. E. Cronin B. Arnold A. D. Twarog F. J. and Dioun A. F. Adverse reactions to vitamin B12 injections dud to benzyl alcohol sensitivity. *Allergy*. 2004;59:(9):1023-1024.

146. Curry, E. J. and Warshaw E. M. Benzyl alcohol allergy: Importance of patch testing with personal products. *Dermatitis*. 2005;16:(4):203-208.
147. Carrascosa, J.-M. Domingo H. Soria X. and Ferrándiz C. Allergic contact dermatitis due to benzyl alcohol in a hair dye. *Contact Dermatitis*. 2006;55:(2):124-125.
148. Jacob, S. E. Ciclosporin ophthalmic emulsion - a novel therapy for benzyl alcohol-associated eyelid dermatitis. *Contact Dermatitis*.%2008., Mar. 58(3):169-70.:(3:169-70):Contact.
149. Sestini, S. Mori M. and Francalanci S. Allergic contact dermatitis from benzyl alcohol in multiple medicaments. *Contact Dermatitis*. 2004;50:(5):316-317.
150. Vilaplana, J. and Romaguera C. Fixed drug eruption from sodium benzoate. *Contact Dermatitis*. 2003;49:(6):290-291.
151. Sutton, T. and Nixon R. Allergic contact dermatitis to sodium benzoate chloroacetamide in a sorbolene lotion. *Australas.J.Dermatol*. 2006;47:(3):209-210.
152. Asero, R. Sodium benzoate-induced pruritus. *Allergy*. 2006;61:(10):1240-1241.
153. Stransky, L. Vasileva S. and Mateev G. Contact bullous pemphigoid? *Contact Dermatitis*. 1996;35:(3):182-182.
154. Daughtry, D. C. Complications following a rapid treatment of scabies. *JAMA*. 1945;127:(1):88-89.
155. Calnan, C. D. Cinnamon dermatitis from an ointment. *Contact Dermatitis*. 1976;2:167-170.
156. McIli, M. C. Giorgini S. and Sertoli A. Occupational dermatitis in a bee-keeper. *Contact Dermatitis*. 1983;9:(5):427-428.